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# **LAMMPS Documentation**

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**The LAMMPS Developers**

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# LAMMPS DOCUMENTATION

<b>I</b>	<b>User Guide</b>	<b>3</b>
<b>1</b>	<b>Introduction</b>	<b>5</b>
1.1	Overview of LAMMPS	5
1.2	What does a LAMMPS version mean	6
1.3	LAMMPS features	6
1.3.1	General features	7
1.3.2	Particle and model types	7
1.3.3	Interatomic potentials (force fields)	8
1.3.4	Atom creation	9
1.3.5	Ensembles, constraints, and boundary conditions	9
1.3.6	Integrators	9
1.3.7	Diagnostics	10
1.3.8	Output	10
1.3.9	Multi-replica models	10
1.3.10	Pre- and post-processing	10
1.3.11	Specialized features	11
1.4	LAMMPS non-features	11
1.5	LAMMPS open-source license	12
1.5.1	GPL version of LAMMPS	12
1.5.2	LGPL version of LAMMPS	13
1.6	Authors of LAMMPS	13
1.7	Citing LAMMPS	14
1.7.1	Core Algorithms	14
1.7.2	DOI for the LAMMPS source code	15
1.7.3	Home page	15
1.7.4	Citing contributions	15
1.8	Additional website links	15
<b>2</b>	<b>Install LAMMPS</b>	<b>17</b>
2.1	Download an executable for Linux	17
2.1.1	Pre-built Ubuntu Linux executables	18
2.1.2	Pre-built Fedora Linux executables	18
2.1.3	Pre-built EPEL Linux executable	19
2.1.4	Pre-built OpenSuse Linux executable	19
2.1.5	Gentoo Linux executable	20
2.1.6	Archlinux build-script	20
2.2	Download an executable for Mac	20
2.3	Download an executable for Windows	21
2.4	Download an executable for Linux or Mac via Conda	22
2.5	Download source and documentation as a tarball	22

2.6	Download the LAMMPS source with git	23
<b>3</b>	<b>Build LAMMPS</b>	<b>27</b>
3.1	Build LAMMPS with CMake	27
3.1.1	Advantages of using CMake	27
3.1.2	Getting started	28
3.1.3	Configuration and build options	29
3.1.4	Multi-configuration build systems	29
3.1.5	Installing CMake	29
3.2	Build LAMMPS with make	30
3.2.1	Requirements	30
3.2.2	Getting started	30
3.2.3	Customized builds and alternate makefiles	31
3.3	Link LAMMPS as a library to another code	31
3.3.1	Link with LAMMPS as a static library	32
3.3.2	Link with LAMMPS as a shared library	33
3.4	Basic build options	35
3.4.1	Serial vs parallel build	35
3.4.2	Choice of compiler and compile/link options	37
3.4.3	Build the LAMMPS executable and library	39
3.4.4	Including or removing debug support	41
3.4.5	Build LAMMPS tools	41
3.4.6	Install LAMMPS after a build	42
3.5	Optional build settings	42
3.5.1	C++11 standard compliance	43
3.5.2	FFT library	43
3.5.3	Size of LAMMPS integer types and size limits	45
3.5.4	Output of JPG, PNG, and movie files	46
3.5.5	Read or write compressed files	47
3.5.6	Memory allocation alignment	48
3.5.7	Workaround for long long integers	48
3.5.8	Exception handling when using LAMMPS as a library	49
3.5.9	Trigger selected floating-point exceptions	49
3.6	Include packages in build	50
3.6.1	Information for both build systems	51
3.6.2	CMake presets for installing many packages	52
3.6.3	Make shortcuts for installing many packages	53
3.7	Packages with extra build options	54
3.7.1	COMPRESS package	54
3.7.2	GPU package	55
3.7.3	KIM package	58
3.7.4	KOKKOS package	60
3.7.5	LATTE package	64
3.7.6	ML-IAP package	65
3.7.7	MSCG package	65
3.7.8	OPT package	66
3.7.9	POEMS package	66
3.7.10	PYTHON package	67
3.7.11	VORONOI package	68
3.7.12	ADIOS package	68
3.7.13	ATC package	69
3.7.14	AWPMD package	70
3.7.15	COLVARS package	71
3.7.16	ELECTRODE package	72

3.7.17	ML-PACE package	72
3.7.18	PLUMED package	73
3.7.19	H5MD package	74
3.7.20	ML-HDNNP package	75
3.7.21	INTEL package	76
3.7.22	MDI package	77
3.7.23	MESONT package	77
3.7.24	MOLFILE package	78
3.7.25	NETCDF package	78
3.7.26	OPENMP package	79
3.7.27	QMMM package	79
3.7.28	ML-QUIP package	80
3.7.29	SCAFACOS package	81
3.7.30	MACHDYN package	82
3.7.31	VTK package	82
3.8	Build the LAMMPS documentation	83
3.8.1	Build using GNU make	83
3.8.2	Build using CMake	84
3.8.3	Prerequisites for HTML	84
3.8.4	Prerequisites for PDF	85
3.8.5	Prerequisites for ePUB and MOBI	85
3.8.6	Instructions for Developers	85
3.9	Notes for building LAMMPS on Windows	86
3.9.1	General remarks	87
3.9.2	Running Linux on Windows	87
3.9.3	Using a GNU GCC ported to Windows	87
3.9.4	Using Microsoft Visual Studio	87
3.9.5	Using Intel oneAPI Compilers and Libraries	88
3.9.6	Using a cross-compiler	89
3.10	Notes for saving disk space when building LAMMPS from source	89
3.11	Development build options	90
3.11.1	Monitor compilation flags (CMake only)	90
3.11.2	Enable static code analysis with clang-tidy (CMake only)	90
3.11.3	Report missing and unneeded '#include' statements (CMake only)	90
3.11.4	Address, Undefined Behavior, and Thread Sanitizer Support (CMake only)	91
3.11.5	Code Coverage and Unit Testing (CMake only)	91
3.11.6	Coding style utilities	96
3.11.7	Clang-format support	96
<b>4</b>	<b>Run LAMMPS</b>	<b>97</b>
4.1	Basics of running LAMMPS	97
4.2	Command-line options	98
4.3	Screen and logfile output	106
4.4	Running LAMMPS on Windows	108
<b>5</b>	<b>Commands</b>	<b>111</b>
5.1	LAMMPS input scripts	111
5.2	Parsing rules for input scripts	112
5.3	Input script structure	114
5.3.1	Initialization	115
5.3.2	System definition	115
5.3.3	Simulation settings	115
5.3.4	Run a simulation	115
5.4	Commands by category	115

5.4.1	Initialization:	116
5.4.2	Setup simulation box:	116
5.4.3	Setup atoms:	116
5.4.4	Force fields:	116
5.4.5	Settings:	116
5.4.6	Operations within timestepping (fixes) and diagnostics (computes):	116
5.4.7	Output:	117
5.4.8	Actions:	117
5.4.9	Input script control:	117
5.5	General commands	117
5.6	Fix commands	118
5.7	Compute commands	119
5.8	Pair_style potentials	120
5.9	Bond_style potentials	122
5.10	Angle_style potentials	122
5.11	Dihedral_style potentials	122
5.12	Improper_style potentials	123
5.13	KSpace solvers	123
5.14	Dump commands	123
5.15	Removed commands and packages	124
5.15.1	Fix ave/spatial and fix ave/spatial/sphere	124
5.15.2	Reset_ids command	124
5.15.3	MEAM package	124
5.15.4	REAX package	124
5.15.5	USER-CUDA package	124
5.15.6	restart2data tool	124
<b>6</b>	<b>Optional packages</b>	<b>125</b>
6.1	Available Packages	125
6.2	Package details	127
6.2.1	ADIOS package	128
6.2.2	AMOEBA package	129
6.2.3	ASPHERE package	129
6.2.4	ATC package	130
6.2.5	AWPMD package	130
6.2.6	BOCS package	130
6.2.7	BODY package	131
6.2.8	BPM package	131
6.2.9	BROWNIAN package	132
6.2.10	CG-DNA package	132
6.2.11	CG-SPICA package	133
6.2.12	CLASS2 package	133
6.2.13	COLLOID package	133
6.2.14	COLVARS package	134
6.2.15	COMPRESS package	134
6.2.16	CORESHELL package	135
6.2.17	DIELECTRIC package	135
6.2.18	DIFFRACTION package	136
6.2.19	DIPOLE package	136
6.2.20	DPD-BASIC package	137
6.2.21	DPD-MESO package	137
6.2.22	DPD-REACT package	138
6.2.23	DPD-SMOOTH package	138
6.2.24	DRUDE package	139

6.2.25	EFF package	139
6.2.26	ELECTRODE package	140
6.2.27	EXTRA-COMPUTE package	140
6.2.28	EXTRA-DUMP package	141
6.2.29	EXTRA-FIX package	141
6.2.30	EXTRA-MOLECULE package	141
6.2.31	EXTRA-PAIR package	141
6.2.32	FEP package	142
6.2.33	GPU package	142
6.2.34	GRANULAR package	143
6.2.35	H5MD package	143
6.2.36	INTEL package	144
6.2.37	INTERLAYER package	145
6.2.38	KIM package	145
6.2.39	KOKKOS package	146
6.2.40	KSPACE package	147
6.2.41	LATBOLTZ package	147
6.2.42	LATTE package	148
6.2.43	MACHDYN package	148
6.2.44	MANIFOLD package	149
6.2.45	MANYBODY package	149
6.2.46	MC package	150
6.2.47	MDI package	150
6.2.48	MEAM package	151
6.2.49	MESONT package	151
6.2.50	MGPT package	152
6.2.51	MISC package	152
6.2.52	ML-HDNNP package	153
6.2.53	ML-IAP package	153
6.2.54	ML-PACE package	154
6.2.55	ML-QUIP package	154
6.2.56	ML-RANN package	155
6.2.57	ML-SNAP package	155
6.2.58	MOFFF package	156
6.2.59	MOLECULE package	156
6.2.60	MOLFILE package	157
6.2.61	MPIIO package	157
6.2.62	MSCG package	158
6.2.63	NETCDF package	158
6.2.64	OPENMP package	159
6.2.65	OPT package	159
6.2.66	ORIENT package	160
6.2.67	PERI package	160
6.2.68	PHONON package	161
6.2.69	PLUGIN package	161
6.2.70	PLUMED package	162
6.2.71	POEMS package	162
6.2.72	PTM package	163
6.2.73	PYTHON package	163
6.2.74	QEQ package	163
6.2.75	QMMM package	164
6.2.76	QTB package	164
6.2.77	REACTION package	165
6.2.78	REAXFF package	165

6.2.79	REPLICA package	166
6.2.80	RIGID package	166
6.2.81	SCAFACOS package	167
6.2.82	SHOCK package	167
6.2.83	SMTBQ package	168
6.2.84	SPH package	168
6.2.85	SPIN package	169
6.2.86	SRD package	169
6.2.87	TALLY package	170
6.2.88	UEF package	170
6.2.89	VORONOI package	171
6.2.90	VTK package	171
6.2.91	YAFF package	172
<b>7</b>	<b>Accelerate performance</b>	<b>173</b>
7.1	Benchmarks	173
7.2	Measuring performance	174
7.3	General tips	175
7.4	Accelerator packages	176
7.4.1	GPU package	177
7.4.2	INTEL package	180
7.4.3	KOKKOS package	188
7.4.4	OPENMP package	194
7.4.5	OPT package	197
7.5	Comparison of various accelerator packages	200
<b>8</b>	<b>Howto discussions</b>	<b>203</b>
8.1	General howto	203
8.1.1	Restart a simulation	203
8.1.2	Visualize LAMMPS snapshots	204
8.1.3	Run multiple simulations from one input script	205
8.1.4	Multi-replica simulations	206
8.1.5	Library interface to LAMMPS	207
8.1.6	Coupling LAMMPS to other codes	207
8.1.7	Using LAMMPS with the MDI library for code coupling	208
8.1.8	Bonded particle models	210
8.1.9	Broken Bonds	211
8.2	Settings howto	212
8.2.1	2d simulations	212
8.2.2	Type labels	213
8.2.3	Triclinic (non-orthogonal) simulation boxes	214
8.2.4	Thermostats	217
8.2.5	Barostats	219
8.2.6	Walls	220
8.2.7	NEMD simulations	221
8.2.8	Long-range dispersion settings	221
8.3	Analysis howto	222
8.3.1	Output from LAMMPS (thermo, dumps, computes, fixes, variables)	222
8.3.2	Use chunks to calculate system properties	227
8.3.3	Calculate temperature	230
8.3.4	Calculate elastic constants	231
8.3.5	Calculate thermal conductivity	232
8.3.6	Calculate viscosity	232
8.3.7	Calculate diffusion coefficients	235



8.3.8	Output structured data from LAMMPS . . . . .	235
8.4	Force fields howto . . . . .	241
8.4.1	CHARMM, AMBER, COMPASS, and DREIDING force fields . . . . .	241
8.4.2	AMOEBA and HIPPO force fields . . . . .	243
8.4.3	TIP3P water model . . . . .	248
8.4.4	TIP4P water model . . . . .	249
8.4.5	SPC water model . . . . .	251
8.5	Packages howto . . . . .	252
8.5.1	Finite-size spherical and aspherical particles . . . . .	252
8.5.2	Granular models . . . . .	255
8.5.3	Body particles . . . . .	256
8.5.4	Polarizable models . . . . .	262
8.5.5	Adiabatic core/shell model . . . . .	263
8.5.6	Drude induced dipoles . . . . .	266
8.5.7	Tutorial for Thermalized Drude oscillators in LAMMPS . . . . .	267
8.5.8	Manifolds (surfaces) . . . . .	274
8.5.9	Magnetic spins . . . . .	275
8.6	Tutorials howto . . . . .	275
8.6.1	Using CMake with LAMMPS tutorial . . . . .	275
8.6.2	LAMMPS GitHub tutorial . . . . .	283
8.6.3	PyLammps Tutorial . . . . .	295
8.6.4	Using LAMMPS on Windows 10 with WSL . . . . .	308
<b>9</b>	<b>Example scripts . . . . .</b>	<b>319</b>
9.1	Lowercase directories . . . . .	319
9.2	Uppercase directories . . . . .	321
<b>10</b>	<b>Auxiliary tools . . . . .</b>	<b>323</b>
10.1	Pre-processing tools . . . . .	323
10.2	Post-processing tools . . . . .	323
10.3	Miscellaneous tools . . . . .	324
10.4	Tool descriptions . . . . .	324
10.4.1	amber2lmp tool . . . . .	324
10.4.2	binary2txt tool . . . . .	324
10.4.3	ch2lmp tool . . . . .	324
10.4.4	chain tool . . . . .	325
10.4.5	LAMMPS coding standard . . . . .	325
10.4.6	colvars tools . . . . .	325
10.4.7	createatoms tool . . . . .	326
10.4.8	drude tool . . . . .	326
10.4.9	eam database tool . . . . .	326
10.4.10	eam generate tool . . . . .	326
10.4.11	eff tool . . . . .	327
10.4.12	emacs tool . . . . .	327
10.4.13	fep tool . . . . .	327
10.4.14	i-pi tool . . . . .	327
10.4.15	ipp tool . . . . .	327
10.4.16	kate tool . . . . .	328
10.4.17	LAMMPS shell . . . . .	328
10.4.18	lmp2arc tool . . . . .	331
10.4.19	lmp2cfg tool . . . . .	331
10.4.20	Magic patterns for the “file” command . . . . .	331
10.4.21	matlab tool . . . . .	332
10.4.22	micelle2d tool . . . . .	332

10.4.23	moltemplate tool	332
10.4.24	msi2lmp tool	332
10.4.25	Scripts for building LAMMPS when offline	333
10.4.26	phonon tool	334
10.4.27	polybond tool	334
10.4.28	pymol_asphere tool	335
10.4.29	python tool	335
10.4.30	replica tool	335
10.4.31	smd tool	335
10.4.32	spin tool	336
10.4.33	singularity/apptainer tool	336
10.4.34	stl_bin2txt tool	336
10.4.35	SWIG interface	336
10.4.36	vim tool	338
10.4.37	xmgrace tool	338
<b>11</b>	<b>Errors</b>	<b>339</b>
11.1	Common problems	339
11.2	Error and warning details	340
11.2.1	Unknown identifier in data file	341
11.3	Reporting bugs	341
11.4	Debugging crashes	341
11.4.1	Using the GDB debugger to get a stack trace	343
11.4.2	Using valgrind to get a stack trace	344
11.5	Error messages	345
11.6	Warning messages	469
<b>II</b>	<b>Programmer Guide</b>	<b>483</b>
<b>12</b>	<b>LAMMPS Library Interfaces</b>	<b>485</b>
12.1	LAMMPS C Library API	485
12.1.1	Creating or deleting a LAMMPS object	486
12.1.2	Executing commands	491
12.1.3	System properties	493
12.1.4	Per-atom properties	503
12.1.5	Compute, fixes, variables	504
12.1.6	Scatter/gather operations	509
12.1.7	Neighbor list access	512
12.1.8	Configuration information	514
12.1.9	Utility functions	521
12.1.10	Extending the C API	528
12.2	LAMMPS Python APIs	529
12.3	LAMMPS Fortran API	529
12.3.1	The LIBLAMMPS Fortran Module	529
12.3.2	Creating or deleting a LAMMPS object	530
12.3.3	The LIBLAMMPS module API	532
12.4	LAMMPS C++ API	533
12.4.1	Using the C++ API directly	533
12.4.2	Creating or deleting a LAMMPS object	533
12.4.3	Executing LAMMPS commands	534
<b>13</b>	<b>Use Python with LAMMPS</b>	<b>535</b>
13.1	Overview	535

13.2	Installation	537
13.2.1	Installing the LAMMPS Python Module and Shared Library	537
13.2.2	Extending Python to run in parallel	543
13.3	Run LAMMPS from Python	544
13.3.1	Running LAMMPS and Python in serial	545
13.3.2	Running LAMMPS and Python in parallel with MPI	545
13.3.3	Running Python scripts	546
13.3.4	Creating or deleting a LAMMPS object	546
13.3.5	Executing commands	548
13.3.6	System properties	550
13.3.7	Per-atom properties	552
13.3.8	Compute, fixes, variables	553
13.3.9	Scatter/gather operations	555
13.3.10	Neighbor list access	556
13.3.11	Configuration information	557
13.4	The <code>lammops</code> Python module	558
13.4.1	The <code>lammops</code> class API	558
13.4.2	The <code>PyLammops</code> class API	577
13.4.3	The <code>IPyLammops</code> class API	582
13.4.4	Additional components of the <code>lammops</code> module	583
13.5	Extending the Python interface	585
13.6	Calling Python from LAMMPS	586
13.7	Output Readers	586
13.8	Example Python scripts	587
13.9	Handling LAMMPS errors	588
13.10	Troubleshooting	589
13.10.1	Testing if Python can launch LAMMPS	589
<b>14</b>	<b>Modifying &amp; extending LAMMPS</b>	<b>591</b>
14.1	Overview	591
14.2	Submitting new features for inclusion in LAMMPS	592
14.2.1	Communication with the LAMMPS developers	593
14.2.2	Packages versus individual files	593
14.2.3	Time and effort required	593
14.2.4	Submission procedure	593
14.2.5	Licensing	594
14.2.6	External contributions	594
14.3	LAMMPS programming style and requirements for contributions	594
14.3.1	Motivation	594
14.3.2	Licensing requirements (strict)	595
14.3.3	Using Pull Requests on GitHub (preferred)	595
14.3.4	Integration Testing (strict)	595
14.3.5	Documentation (strict)	596
14.3.6	Examples (preferred)	596
14.3.7	Howto document (optional)	597
14.3.8	Programming Style Requirements (varied)	597
14.3.9	Contributing a package	601
14.3.10	Build system (strongly preferred)	601
14.3.11	Citation reminder (suggested)	601
14.3.12	Testing (optional)	601
14.4	Atom styles	602
14.5	Pair styles	604
14.6	Bond, angle, dihedral, improper styles	604
14.7	Compute styles	605

14.8	Fix styles	606
14.9	Input script command style	607
14.10	Dump styles	608
14.11	Kspace styles	608
14.12	Minimization styles	608
14.13	Region styles	609
14.14	Body styles	609
14.15	Thermodynamic output options	609
14.16	Variable options	610
<b>15</b>	<b>Information for Developers</b>	<b>611</b>
15.1	Source files	611
15.2	Class topology	612
15.3	Code design	615
15.3.1	Object oriented code	615
15.3.2	I/O and output formatting	619
15.3.3	Memory management	620
15.4	Parallel algorithms	621
15.4.1	Partitioning	621
15.4.2	Communication	622
15.4.3	Neighbor lists	624
15.4.4	Long-range interactions	626
15.4.5	OpenMP Parallelism	628
15.5	Communication patterns	630
15.5.1	Owned and ghost atoms	630
15.5.2	Higher level communication	631
15.6	How a timestep works	633
15.7	Writing new styles	636
15.7.1	Writing a new fix style	636
15.8	Notes for developers and code maintainers	640
15.8.1	Reading and parsing of text and text files	640
15.8.2	Requesting and accessing neighbor lists	641
15.8.3	Fix contributions to instantaneous energy, virial, and cumulative energy	643
15.8.4	KSpace PPPM FFT grids	644
15.9	Notes for updating code written for older LAMMPS versions	646
15.9.1	Setting flags in the constructor	646
15.9.2	Rename of pack/unpack_comm() to pack/unpack_forward_comm()	646
15.9.3	Use ev_init() to initialize variables derived from eflag and vflag	647
15.9.4	Use utils::numeric() functions instead of force->numeric()	648
15.9.5	Use utils::open_potential() function to open potential files	648
15.9.6	Simplify customized error messages	648
15.9.7	Use of “override” instead of “virtual”	649
15.9.8	Simplified function names for forward and reverse communication	649
15.9.9	Simplified and more compact neighbor list requests	650
15.9.10	Split of fix STORE into fix STORE/GLOBAL and fix STORE/PERATOM	650
15.9.11	Use Output::get_dump_by_id() instead of Output::find_dump()	651
15.10	Writing plugins	652
15.10.1	Members of lammpsplugin_t	653
15.10.2	Pair style example	653
15.10.3	Fix style example	654
15.10.4	Command style example	654
15.10.5	Additional Details	655
15.10.6	Compiling plugins	656
15.11	Adding tests for unit testing	656

15.11.1	Tests for utility functions	657
15.11.2	Tests for individual LAMMPS commands	658
15.11.3	Tests for the C-style library interface	659
15.11.4	Tests for the Python module and package	660
15.11.5	Tests for the Fortran interface	660
15.11.6	Tests for the C++-style library interface	660
15.11.7	Tests for reading and writing file formats	660
15.11.8	Tests for styles computing or modifying forces	660
15.11.9	Tests for programs in the tools folder	663
15.12	C++ base classes	663
15.12.1	LAMMPS Class	664
15.12.2	LAMMPS Atom and AtomVec Base Classes	665
15.12.3	LAMMPS Input Base Class	668
15.13	Platform abstraction functions	669
15.13.1	Time functions	669
15.13.2	Platform information functions	670
15.13.3	File and path functions and global constants	671
15.13.4	Standard I/O function wrappers	674
15.13.5	Environment variable functions	675
15.13.6	Dynamically loaded object or library functions	676
15.13.7	Compressed file I/O functions	676
15.14	Utility functions	677
15.14.1	I/O with status check and similar functions	677
15.14.2	String to number conversions with validity check	679
15.14.3	String processing	682
15.14.4	Potential file functions	687
15.14.5	Argument processing	688
15.14.6	Convenience functions	690
15.14.7	Customized standard functions	692
15.15	Special Math functions	692
15.16	Tokenizer classes	694
15.17	Argument parsing classes	699
15.18	File reader classes	702
15.19	Memory pool classes	706
15.20	Eigensolver functions	710
15.21	Communication buffer coding with <i>ubuf</i>	711

### III Command Reference

713

#### 16 Commands

715

16.1	angle_coeff command	715
16.1.1	Syntax	715
16.1.2	Examples	715
16.1.3	Description	715
16.1.4	Restrictions	716
16.1.5	Related commands	716
16.1.6	Default	716
16.2	angle_style command	716
16.2.1	Syntax	716
16.2.2	Examples	717
16.2.3	Description	717
16.2.4	Restrictions	718
16.2.5	Related commands	718

16.2.6	Default	718
16.3	atom_modify command	718
16.3.1	Syntax	718
16.3.2	Examples	719
16.3.3	Description	719
16.3.4	Restrictions	720
16.3.5	Related commands	720
16.3.6	Default	721
16.4	atom_style command	721
16.4.1	Syntax	721
16.4.2	Examples	721
16.4.3	Description	722
16.4.4	Restrictions	725
16.4.5	Related commands	726
16.4.6	Default	726
16.5	balance command	726
16.5.1	Syntax	726
16.5.2	Examples	727
16.5.3	Description	727
16.5.4	Restrictions	733
16.5.5	Related commands	733
16.5.6	Default	733
16.6	bond_coeff command	733
16.6.1	Syntax	733
16.6.2	Examples	733
16.6.3	Description	734
16.6.4	Restrictions	734
16.6.5	Related commands	734
16.6.6	Default	735
16.7	bond_style command	735
16.7.1	Syntax	735
16.7.2	Examples	735
16.7.3	Description	735
16.7.4	Restrictions	737
16.7.5	Related commands	737
16.7.6	Default	737
16.8	bond_write command	737
16.8.1	Syntax	737
16.8.2	Examples	737
16.8.3	Description	737
16.8.4	Restrictions	738
16.8.5	Related commands	738
16.8.6	Default	738
16.9	boundary command	738
16.9.1	Syntax	738
16.9.2	Examples	738
16.9.3	Description	739
16.9.4	Restrictions	740
16.9.5	Related commands	740
16.9.6	Default	740
16.10	box command	740
16.10.1	Syntax	740
16.10.2	Examples	740
16.10.3	Description	740

16.10.4	Restrictions	741
16.10.5	Related commands	741
16.10.6	Default	741
16.11	change_box command	741
16.11.1	Syntax	741
16.11.2	Examples	742
16.11.3	Description	742
16.11.4	Restrictions	745
16.11.5	Related commands	745
16.11.6	Default	746
16.12	clear command	746
16.12.1	Syntax	746
16.12.2	Examples	746
16.12.3	Description	746
16.12.4	Restrictions	746
16.12.5	Related commands	746
16.12.6	Default	746
16.13	comm_modify command	747
16.13.1	Syntax	747
16.13.2	Examples	747
16.13.3	Description	747
16.13.4	Restrictions	749
16.13.5	Related commands	749
16.13.6	Default	749
16.14	comm_style command	749
16.14.1	Syntax	749
16.14.2	Examples	750
16.14.3	Description	750
16.14.4	Restrictions	750
16.14.5	Related commands	750
16.14.6	Default	750
16.15	compute command	750
16.15.1	Syntax	750
16.15.2	Examples	751
16.15.3	Description	751
16.15.4	Restrictions	757
16.15.5	Related commands	757
16.15.6	Default	757
16.16	compute_modify command	757
16.16.1	Syntax	757
16.16.2	Examples	758
16.16.3	Description	758
16.16.4	Restrictions	758
16.16.5	Related commands	758
16.16.6	Default	758
16.17	create_atoms command	758
16.17.1	Syntax	758
16.17.2	Examples	760
16.17.3	Description	760
16.17.4	Restrictions	765
16.17.5	Related commands	765
16.17.6	Default	765
16.18	create_bonds command	765
16.18.1	Syntax	765

16.18.2	Examples	766
16.18.3	Description	766
16.18.4	Restrictions	768
16.18.5	Related commands	768
16.18.6	Default	768
16.19	create_box command	768
16.19.1	Syntax	768
16.19.2	Examples	769
16.19.3	Description	769
16.19.4	Restrictions	770
16.19.5	Related commands	770
16.19.6	Default	770
16.20	delete_atoms command	770
16.20.1	Syntax	770
16.20.2	Examples	771
16.20.3	Description	771
16.20.4	Restrictions	773
16.20.5	Related commands	773
16.20.6	Default	773
16.21	delete_bonds command	773
16.21.1	Syntax	773
16.21.2	Examples	774
16.21.3	Description	774
16.21.4	Restrictions	775
16.21.5	Related commands	775
16.21.6	Default	775
16.22	dielectric command	775
16.22.1	Syntax	775
16.22.2	Examples	776
16.22.3	Description	776
16.22.4	Restrictions	776
16.22.5	Related commands	776
16.22.6	Default	776
16.23	dihedral_coeff command	776
16.23.1	Syntax	776
16.23.2	Examples	776
16.23.3	Description	777
16.23.4	Restrictions	778
16.23.5	Related commands	778
16.23.6	Default	778
16.24	dihedral_style command	778
16.24.1	Syntax	778
16.24.2	Examples	778
16.24.3	Description	778
16.24.4	Restrictions	780
16.24.5	Related commands	780
16.24.6	Default	780
16.25	dimension command	780
16.25.1	Syntax	780
16.25.2	Examples	780
16.25.3	Description	780
16.25.4	Restrictions	781
16.25.5	Related commands	781
16.25.6	Default	781



16.26	displace_atoms command	781
16.26.1	Syntax	781
16.26.2	Examples	782
16.26.3	Description	782
16.26.4	Restrictions	783
16.26.5	Related commands	783
16.26.6	Default	783
16.27	dump command	783
16.28	dump vtk command	783
16.29	dump h5md command	783
16.30	dump molfile command	783
16.31	dump netcdf command	783
16.32	dump image command	783
16.33	dump movie command	783
16.34	dump atom/adios command	783
16.35	dump custom/adios command	783
16.36	dump cfg/uef command	783
16.36.1	Syntax	783
16.36.2	Examples	786
16.36.3	Description	786
16.36.4	Restrictions	793
16.36.5	Related commands	794
16.36.6	Default	794
16.37	dump atom/adios command	794
16.38	dump custom/adios command	794
16.38.1	Syntax	794
16.38.2	Examples	794
16.38.3	Description	794
16.38.4	Restrictions	795
16.38.5	Related commands	795
16.39	dump cfg/uef command	795
16.39.1	Syntax	795
16.39.2	Examples	795
16.39.3	Description	796
16.39.4	Restrictions	796
16.39.5	Related commands	796
16.39.6	Default	796
16.40	dump h5md command	796
16.40.1	Syntax	796
16.40.2	Examples	797
16.40.3	Description	797
16.40.4	Restrictions	798
16.40.5	Related commands	798
16.41	dump image command	798
16.42	dump movie command	798
16.42.1	Syntax	798
16.43	dump_modify options for dump image/movie	800
16.43.1	Syntax	800
16.43.2	Examples	801
16.43.3	Description	801
16.43.4	Dump_modify keywords for dump image and dump movie	807
16.43.5	Restrictions	810
16.43.6	Related commands	811
16.43.7	Default	811

16.44	dump_modify command	814
16.45	dump_modify command for image/movie options	814
16.45.1	Syntax	814
16.45.2	Examples	816
16.45.3	Description	816
16.45.4	Restrictions	825
16.45.5	Related commands	825
16.45.6	Default	825
16.46	dump molfile command	826
16.46.1	Syntax	826
16.46.2	Examples	826
16.46.3	Description	826
16.46.4	Restrictions	827
16.46.5	Related commands	827
16.46.6	Default	827
16.47	dump netcdf command	828
16.48	dump netcdf/mpiio command	828
16.48.1	Syntax	828
16.48.2	Examples	828
16.48.3	Description	828
16.48.4	Restrictions	829
16.48.5	Related commands	829
16.49	dump vtk command	829
16.49.1	Syntax	829
16.49.2	Examples	829
16.49.3	Description	829
16.49.4	Restrictions	831
16.49.5	Related commands	831
16.49.6	Default	831
16.50	dynamical_matrix command	831
16.50.1	Syntax	831
16.50.2	Examples	832
16.50.3	Description	832
16.50.4	Restrictions	832
16.50.5	Related commands	833
16.50.6	Default	833
16.51	echo command	833
16.51.1	Syntax	833
16.51.2	Examples	833
16.51.3	Description	833
16.51.4	Restrictions	833
16.51.5	Related commands	833
16.51.6	Default	834
16.52	fix command	834
16.52.1	Syntax	834
16.52.2	Examples	834
16.52.3	Description	834
16.52.4	Restrictions	843
16.52.5	Related commands	843
16.52.6	Default	843
16.53	fix_modify command	843
16.53.1	Syntax	843
16.53.2	Examples	844
16.53.3	Description	844

16.53.4	Restrictions	846
16.53.5	Related commands	846
16.53.6	Default	846
16.54	group command	846
16.54.1	Syntax	846
16.54.2	Examples	847
16.54.3	Description	847
16.54.4	Restrictions	850
16.54.5	Related commands	850
16.54.6	Default	850
16.55	group2ndx command	850
16.56	ndx2group command	850
16.56.1	Syntax	850
16.56.2	Examples	851
16.56.3	Description	851
16.56.4	Restrictions	851
16.56.5	Related commands	851
16.56.6	Default	851
16.57	hyper command	851
16.57.1	Syntax	851
16.57.2	Examples	852
16.57.3	Description	852
16.57.4	Restrictions	854
16.57.5	Related commands	854
16.57.6	Default	854
16.58	if command	854
16.58.1	Syntax	854
16.58.2	Examples	854
16.58.3	Description	855
16.58.4	Restrictions	857
16.58.5	Related commands	857
16.58.6	Default	857
16.59	improper_coeff command	857
16.59.1	Syntax	857
16.59.2	Examples	857
16.59.3	Description	858
16.59.4	Restrictions	858
16.59.5	Related commands	858
16.59.6	Default	859
16.60	improper_style command	859
16.60.1	Syntax	859
16.60.2	Examples	859
16.60.3	Description	859
16.60.4	Restrictions	860
16.60.5	Related commands	860
16.60.6	Default	860
16.61	include command	860
16.61.1	Syntax	860
16.61.2	Examples	861
16.61.3	Description	861
16.61.4	Restrictions	861
16.61.5	Related commands	861
16.61.6	Default	861
16.62	info command	861

16.62.1	Syntax	861
16.62.2	Examples	862
16.62.3	Description	862
16.62.4	Restrictions	863
16.62.5	Related commands	863
16.62.6	Default	863
16.63	jump command	863
16.63.1	Syntax	863
16.63.2	Examples	864
16.63.3	Description	864
16.63.4	Restrictions	865
16.63.5	Related commands	865
16.63.6	Default	865
16.64	kim command	866
16.64.1	Syntax	866
16.64.2	Examples	866
16.64.3	Description	866
16.64.4	Using OpenKIM IMs with LAMMPS ( <i>kim init, kim interactions</i> )	867
16.64.5	Using OpenKIM Web Queries in LAMMPS ( <i>kim query</i> )	871
16.64.6	Accessing KIM Model Parameters from LAMMPS ( <i>kim param</i> )	874
16.64.7	Writing material properties in standard KIM Property Instance format ( <i>kim property</i> )	878
16.64.8	Citation of OpenKIM IMs	884
16.64.9	Restrictions	884
16.64.10	Related commands	884
16.65	kspace_modify command	884
16.65.1	Syntax	884
16.65.2	Examples	885
16.65.3	Description	886
16.65.4	Restrictions	890
16.65.5	Related commands	890
16.65.6	Default	890
16.66	kspace_style command	891
16.66.1	Syntax	891
16.66.2	Examples	892
16.66.3	Description	893
16.66.4	Restrictions	897
16.66.5	Related commands	897
16.66.6	Default	897
16.67	label command	898
16.67.1	Syntax	898
16.67.2	Examples	898
16.67.3	Description	898
16.67.4	Restrictions	898
16.67.5	Related commands	898
16.67.6	Default	899
16.68	labelmap command	899
16.68.1	Syntax	899
16.68.2	Examples	899
16.68.3	Description	899
16.68.4	Restrictions	900
16.68.5	Related commands	900
16.68.6	Default	900
16.69	lattice command	900
16.69.1	Syntax	900

16.69.2	Examples	901
16.69.3	Description	901
16.69.4	Restrictions	904
16.69.5	Related commands	904
16.69.6	Default	904
16.70	log command	904
16.70.1	Syntax	904
16.70.2	Examples	904
16.70.3	Description	904
16.70.4	Restrictions	905
16.70.5	Related commands	905
16.70.6	Default	905
16.71	mass command	905
16.71.1	Syntax	905
16.71.2	Examples	905
16.71.3	Description	905
16.71.4	Restrictions	906
16.71.5	Related commands	906
16.71.6	Default	906
16.72	mdi command	906
16.72.1	Syntax	906
16.72.2	Examples	907
16.72.3	Description	907
16.72.4	Restrictions	911
16.72.5	Related commands	911
16.72.6	Default	911
16.73	min_modify command	912
16.73.1	Syntax	912
16.73.2	Examples	912
16.73.3	Description	912
16.73.4	Restrictions	914
16.73.5	Related commands	914
16.73.6	Default	914
16.74	min_style spin command	914
16.75	min_style spin/cg command	914
16.76	min_style spin/lbfgs command	914
16.76.1	Syntax	914
16.76.2	Examples	914
16.76.3	Description	914
16.76.4	Restrictions	915
16.76.5	Related commands	915
16.76.6	Default	916
16.77	min_style cg command	916
16.78	min_style hftn command	916
16.79	min_style sd command	916
16.80	min_style quickmin command	916
16.81	min_style fire command	916
16.82	min_style fire/old command	916
16.83	min_style spin command	916
16.84	min_style spin/cg command	916
16.85	min_style spin/lbfgs command	916
16.85.1	Syntax	916
16.85.2	Examples	916
16.85.3	Description	917

16.85.4	Restrictions	918
16.85.5	Related commands	918
16.85.6	Default	918
16.86	minimize command	918
16.87	minimize/kk command	918
16.87.1	Syntax	918
16.87.2	Examples	919
16.87.3	Description	919
16.87.4	Restrictions	923
16.87.5	Related commands	923
16.87.6	Default	923
16.88	molecule command	923
16.88.1	Syntax	923
16.88.2	Examples	924
16.88.3	Description	924
16.88.4	Restrictions	930
16.88.5	Related commands	931
16.88.6	Default	931
16.89	neb command	931
16.89.1	Syntax	931
16.89.2	Examples	931
16.89.3	Description	932
16.89.4	Restrictions	936
16.89.5	Related commands	936
16.89.6	Default	936
16.90	neb/spin command	936
16.90.1	Syntax	936
16.90.2	Examples	937
16.90.3	Description	937
16.90.4	Restrictions	941
16.90.5	Related commands	941
16.90.6	Default	941
16.91	neigh_modify command	941
16.91.1	Syntax	941
16.91.2	Examples	942
16.91.3	Description	942
16.91.4	Restrictions	944
16.91.5	Related commands	945
16.91.6	Default	945
16.92	neighbor command	945
16.92.1	Syntax	945
16.92.2	Examples	945
16.92.3	Description	945
16.92.4	Restrictions	946
16.92.5	Related commands	946
16.92.6	Default	946
16.93	newton command	947
16.93.1	Syntax	947
16.93.2	Examples	947
16.93.3	Description	947
16.93.4	Restrictions	947
16.93.5	Related commands	947
16.93.6	Default	948
16.94	next command	948

16.94.1	Syntax . . . . .	948
16.94.2	Examples . . . . .	948
16.94.3	Description . . . . .	948
16.94.4	Restrictions . . . . .	950
16.94.5	Related commands . . . . .	950
16.94.6	Default . . . . .	950
16.95	package command . . . . .	950
16.95.1	Syntax . . . . .	950
16.95.2	Examples . . . . .	952
16.95.3	Description . . . . .	952
16.95.4	Restrictions . . . . .	958
16.95.5	Related commands . . . . .	959
16.95.6	Default . . . . .	959
16.96	pair_coeff command . . . . .	959
16.96.1	Syntax . . . . .	959
16.96.2	Examples . . . . .	959
16.96.3	Description . . . . .	960
16.96.4	Restrictions . . . . .	961
16.96.5	Related commands . . . . .	961
16.96.6	Default . . . . .	961
16.97	pair_modify command . . . . .	962
16.97.1	Syntax . . . . .	962
16.97.2	Examples . . . . .	962
16.97.3	Description . . . . .	963
16.97.4	Restrictions . . . . .	966
16.97.5	Related commands . . . . .	966
16.97.6	Default . . . . .	966
16.98	pair_style command . . . . .	966
16.98.1	Syntax . . . . .	966
16.98.2	Examples . . . . .	966
16.98.3	Description . . . . .	966
16.98.4	Restrictions . . . . .	975
16.98.5	Related commands . . . . .	975
16.98.6	Default . . . . .	975
16.99	pair_write command . . . . .	975
16.99.1	Syntax . . . . .	975
16.99.2	Examples . . . . .	976
16.99.3	Description . . . . .	976
16.99.4	Restrictions . . . . .	976
16.99.5	Related commands . . . . .	976
16.99.6	Default . . . . .	977
16.100	partition command . . . . .	977
16.100.1	Syntax . . . . .	977
16.100.2	Examples . . . . .	977
16.100.3	Description . . . . .	977
16.100.4	Restrictions . . . . .	978
16.100.5	Related commands . . . . .	978
16.100.6	Default . . . . .	978
16.101	plugin command . . . . .	978
16.101.1	Syntax . . . . .	978
16.101.2	Examples . . . . .	978
16.101.3	Description . . . . .	978
16.101.4	Restrictions . . . . .	979
16.101.5	Related commands . . . . .	979

16.101.6Default	979
16.102prd command	979
16.102.1Syntax	979
16.102.2Examples	980
16.102.3Description	980
16.102.4Restrictions	983
16.102.5Related commands	983
16.102.6Default	983
16.103print command	983
16.103.1Syntax	983
16.103.2Examples	984
16.103.3Description	984
16.103.4Restrictions	985
16.103.5Related commands	985
16.103.6Default	985
16.104processors command	985
16.104.1Syntax	985
16.104.2Examples	986
16.104.3Description	986
16.104.4Restrictions	989
16.104.5Related commands	989
16.104.6Default	989
16.105python command	989
16.105.1Syntax	989
16.105.2Examples	990
16.105.3Description	991
16.105.4Restrictions	996
16.105.5Related commands	996
16.105.6Default	996
16.106quit command	996
16.106.1Syntax	996
16.106.2Examples	997
16.106.3Description	997
16.106.4Restrictions	997
16.106.5Related commands	997
16.106.6Default	997
16.107read_data command	997
16.107.1Syntax	997
16.107.2Examples	998
16.107.3Description	998
16.107.4Reading multiple data files	999
16.107.5Format of a data file	1000
16.107.6Format of the header of a data file	1001
16.107.7Format of the body of a data file	1004
16.107.8Restrictions	1017
16.107.9Related commands	1018
16.107.10Default	1018
16.108read_dump command	1018
16.108.1Syntax	1018
16.108.2Examples	1019
16.108.3Description	1019
16.108.4Restrictions	1022
16.108.5Related commands	1023
16.108.6Default	1023



16.109	read_restart command	1023
16.109.1	Syntax	1023
16.109.2	Examples	1023
16.109.3	Description	1023
16.109.4	Restrictions	1026
16.109.5	Related commands	1026
16.109.6	Default	1026
16.110	region command	1026
16.110.1	Syntax	1026
16.110.2	Examples	1028
16.110.3	Description	1028
16.110.4	Restrictions	1032
16.110.5	Related commands	1032
16.110.6	Default	1032
16.111	replicate command	1032
16.111.1	Syntax	1032
16.111.2	Examples	1032
16.111.3	Description	1032
16.111.4	Restrictions	1033
16.111.5	Related commands	1033
16.111.6	Default	1033
16.112	rerun command	1033
16.112.1	Syntax	1033
16.112.2	Examples	1034
16.112.3	Description	1034
16.112.4	Restrictions	1036
16.112.5	Related commands	1036
16.112.6	Default	1036
16.113	reset_atom_ids command	1036
16.113.1	Syntax	1036
16.113.2	Examples	1037
16.113.3	Description	1037
16.113.4	Restrictions	1037
16.113.5	Related commands	1037
16.113.6	Default	1038
16.114	reset_mol_ids command	1038
16.114.1	Syntax	1038
16.114.2	Examples	1038
16.114.3	Description	1038
16.114.4	Restrictions	1039
16.114.5	Related commands	1039
16.114.6	Default	1039
16.115	reset_timestep command	1039
16.115.1	Syntax	1039
16.115.2	Examples	1040
16.115.3	Description	1040
16.115.4	Restrictions	1040
16.115.5	Related commands	1040
16.115.6	Default	1040
16.116	restart command	1040
16.116.1	Syntax	1040
16.116.2	Examples	1041
16.116.3	Description	1041
16.116.4	Restrictions	1043

16.116.5	Related commands	1043
16.116.6	Default	1043
16.117	run command	1043
16.117.1	Syntax	1043
16.117.2	Examples	1043
16.117.3	Description	1044
16.117.4	Restrictions	1046
16.117.5	Related commands	1046
16.117.6	Default	1046
16.118	run_style command	1046
16.118.1	Syntax	1046
16.118.2	Examples	1047
16.118.3	Description	1047
16.118.4	Restrictions	1050
16.118.5	Related commands	1050
16.118.6	Default	1050
16.119	set command	1050
16.119.1	Syntax	1050
16.119.2	Examples	1053
16.119.3	Description	1053
16.119.4	Restrictions	1057
16.119.5	Related commands	1057
16.119.6	Default	1058
16.120	shell command	1058
16.120.1	Syntax	1058
16.120.2	Examples	1058
16.120.3	Description	1058
16.120.4	Restrictions	1059
16.120.5	Related commands	1059
16.120.6	Default	1060
16.121	special_bonds command	1060
16.121.1	Syntax	1060
16.121.2	Examples	1060
16.121.3	Description	1060
16.121.4	Restrictions	1063
16.121.5	Related commands	1063
16.121.6	Default	1063
16.122	suffix command	1063
16.122.1	Syntax	1063
16.122.2	Examples	1063
16.122.3	Description	1064
16.122.4	Restrictions	1065
16.122.5	Related commands	1065
16.122.6	Default	1065
16.123	ad command	1065
16.123.1	Syntax	1065
16.123.2	Examples	1066
16.123.3	Description	1066
16.123.4	Restrictions	1068
16.123.5	Related commands	1069
16.123.6	Default	1069
16.124	temper command	1069
16.124.1	Syntax	1069
16.124.2	Examples	1069

16.124.3	Description	1069
16.124.4	Restrictions	1071
16.124.5	Related commands	1071
16.124.6	Default	1071
16.125	temper/grem command	1071
16.125.1	Syntax	1071
16.125.2	Examples	1072
16.125.3	Description	1072
16.125.4	Restrictions	1073
16.125.5	Related commands	1073
16.125.6	Default	1073
16.126	temper/npt command	1073
16.126.1	Syntax	1073
16.126.2	Examples	1073
16.126.3	Description	1074
16.126.4	Restrictions	1074
16.126.5	Related commands	1074
16.126.6	Default	1074
16.127	thermo command	1074
16.127.1	Syntax	1074
16.127.2	Examples	1075
16.127.3	Description	1075
16.127.4	Restrictions	1075
16.127.5	Related commands	1075
16.127.6	Default	1075
16.128	thermo_modify command	1075
16.128.1	Syntax	1075
16.128.2	Examples	1076
16.128.3	Description	1076
16.128.4	Restrictions	1079
16.128.5	Related commands	1079
16.128.6	Default	1079
16.129	thermo_style command	1079
16.129.1	Syntax	1079
16.129.2	Examples	1081
16.129.3	Description	1081
16.129.4	Restrictions	1085
16.129.5	Related commands	1085
16.129.6	Default	1085
16.130	third_order command	1085
16.130.1	Syntax	1085
16.130.2	Examples	1085
16.130.3	Description	1086
16.130.4	Restrictions	1086
16.130.5	Related commands	1086
16.130.6	Default	1086
16.131	timer command	1087
16.131.1	Syntax	1087
16.131.2	Examples	1087
16.131.3	Description	1087
16.131.4	Restrictions	1088
16.131.5	Related commands	1088
16.131.6	Default	1088
16.132	timestep command	1088

16.132.1	Syntax . . . . .	1088
16.132.2	Examples . . . . .	1089
16.132.3	Description . . . . .	1089
16.132.4	Restrictions . . . . .	1089
16.132.5	Related commands . . . . .	1089
16.132.6	Default . . . . .	1089
16.133	uncompute command . . . . .	1089
16.133.1	Syntax . . . . .	1089
16.133.2	Examples . . . . .	1090
16.133.3	Description . . . . .	1090
16.133.4	Restrictions . . . . .	1090
16.133.5	Related commands . . . . .	1090
16.133.6	Default . . . . .	1090
16.134	undump command . . . . .	1090
16.134.1	Syntax . . . . .	1090
16.134.2	Examples . . . . .	1090
16.134.3	Description . . . . .	1090
16.134.4	Restrictions . . . . .	1091
16.134.5	Related commands . . . . .	1091
16.134.6	Default . . . . .	1091
16.135	unfix command . . . . .	1091
16.135.1	Syntax . . . . .	1091
16.135.2	Examples . . . . .	1091
16.135.3	Description . . . . .	1091
16.135.4	Restrictions . . . . .	1091
16.135.5	Related commands . . . . .	1091
16.135.6	Default . . . . .	1092
16.136	units command . . . . .	1092
16.136.1	Syntax . . . . .	1092
16.136.2	Examples . . . . .	1092
16.136.3	Description . . . . .	1092
16.136.4	Restrictions . . . . .	1096
16.136.5	Related commands . . . . .	1096
16.136.6	Default . . . . .	1096
16.137	variable command . . . . .	1097
16.137.1	Syntax . . . . .	1097
16.137.2	Examples . . . . .	1098
16.137.3	Description . . . . .	1099
16.137.4	Immediate Evaluation of Variables . . . . .	1113
16.137.5	Variable Accuracy . . . . .	1114
16.137.6	Restrictions . . . . .	1116
16.137.7	Related commands . . . . .	1116
16.137.8	Default . . . . .	1116
16.138	velocity command . . . . .	1116
16.138.1	Syntax . . . . .	1116
16.138.2	Examples . . . . .	1117
16.138.3	Description . . . . .	1117
16.138.4	Restrictions . . . . .	1119
16.138.5	Related commands . . . . .	1119
16.138.6	Default . . . . .	1119
16.139	write_coeff command . . . . .	1120
16.139.1	Syntax . . . . .	1120
16.139.2	Examples . . . . .	1120
16.139.3	Description . . . . .	1120

16.139.4	Restrictions	1120
16.139.5	Related commands	1120
16.140	write_data command	1120
16.140.1	Syntax	1120
16.140.2	Examples	1121
16.140.3	Description	1121
16.140.4	Restrictions	1122
16.140.5	Related commands	1122
16.140.6	Default	1123
16.141	write_dump command	1123
16.141.1	Syntax	1123
16.141.2	Examples	1123
16.141.3	Description	1123
16.141.4	Restrictions	1124
16.141.5	Related commands	1124
16.141.6	Default	1124
16.142	write_restart command	1124
16.142.1	Syntax	1124
16.142.2	Examples	1125
16.142.3	Description	1125
16.142.4	Restrictions	1126
16.142.5	Related commands	1126
16.142.6	Default	1126
<b>17</b>	<b>Fixes</b>	<b>1127</b>
17.1	fix accelerate/cos command	1127
17.1.1	Syntax	1127
17.1.2	Examples	1127
17.1.3	Description	1127
17.1.4	Restart, fix_modify, output, run start/stop, minimize info	1128
17.1.5	Restrictions	1128
17.1.6	Related commands	1128
17.1.7	Default	1128
17.2	fix acks2/reaxff command	1128
17.2.1	Syntax	1128
17.2.2	Examples	1129
17.2.3	Description	1129
17.2.4	Restart, fix_modify, output, run start/stop, minimize info	1130
17.2.5	Restrictions	1130
17.2.6	Related commands	1130
17.2.7	Default	1130
17.3	fix adapt command	1131
17.3.1	Syntax	1131
17.3.2	Examples	1132
17.3.3	Description	1132
17.3.4	Restart, fix_modify, output, run start/stop, minimize info	1136
17.3.5	Restrictions	1136
17.3.6	Related commands	1136
17.3.7	Default	1136
17.4	fix adapt/fep command	1136
17.4.1	Syntax	1136
17.4.2	Examples	1137
17.4.3	Description	1137
17.4.4	Restart, fix_modify, output, run start/stop, minimize info	1140

17.4.5	Restrictions	1140
17.4.6	Related commands	1140
17.4.7	Default	1140
17.5	fix addforce command	1140
17.5.1	Syntax	1140
17.5.2	Examples	1141
17.5.3	Description	1141
17.5.4	Restart, fix_modify, output, run start/stop, minimize info	1142
17.5.5	Restrictions	1143
17.5.6	Related commands	1143
17.5.7	Default	1143
17.6	fix addtorque command	1143
17.6.1	Syntax	1143
17.6.2	Examples	1143
17.6.3	Description	1143
17.6.4	Restart, fix_modify, output, run start/stop, minimize info	1144
17.6.5	Restrictions	1144
17.6.6	Related commands	1144
17.6.7	Default	1144
17.7	fix amoeba/bitorsion command	1144
17.7.1	Syntax	1144
17.7.2	Examples	1145
17.7.3	Description	1145
17.7.4	Restart, fix_modify, output, run start/stop, minimize info	1146
17.7.5	Restrictions	1146
17.7.6	Related commands	1146
17.7.7	Default	1146
17.8	fix amoeba/pitorsion command	1147
17.8.1	Syntax	1147
17.8.2	Examples	1147
17.8.3	Description	1147
17.8.4	Restart, fix_modify, output, run start/stop, minimize info	1148
17.8.5	Restrictions	1149
17.8.6	Related commands	1149
17.8.7	Default	1149
17.9	fix append/atoms command	1149
17.9.1	Syntax	1149
17.9.2	Examples	1150
17.9.3	Description	1150
17.9.4	Restart, fix_modify, output, run start/stop, minimize info	1150
17.9.5	Restrictions	1150
17.9.6	Related commands	1151
17.9.7	Default	1151
17.10	fix atc command	1151
17.10.1	Syntax	1151
17.10.2	Examples	1151
17.10.3	Description	1151
17.10.4	Restart, fix_modify, output, run start/stop, minimize info	1153
17.10.5	Restrictions	1154
17.10.6	Related commands	1154
17.10.7	Default	1156
17.11	fix atom/swap command	1157
17.11.1	Syntax	1157
17.11.2	Examples	1157

17.11.3	Description	1157
17.11.4	Restart, fix_modify, output, run start/stop, minimize info	1159
17.11.5	Restrictions	1159
17.11.6	Related commands	1159
17.11.7	Default	1159
17.12	fix ave/atom command	1159
17.12.1	Syntax	1159
17.12.2	Examples	1160
17.12.3	Description	1160
17.12.4	Restart, fix_modify, output, run start/stop, minimize info	1162
17.12.5	Restrictions	1162
17.12.6	Related commands	1162
17.12.7	Default	1162
17.13	fix ave/chunk command	1162
17.13.1	Syntax	1162
17.13.2	Examples	1163
17.13.3	Description	1164
17.13.4	Restart, fix_modify, output, run start/stop, minimize info	1168
17.13.5	Restrictions	1168
17.13.6	Related commands	1169
17.13.7	Default	1169
17.14	fix ave/correlate command	1169
17.14.1	Syntax	1169
17.14.2	Examples	1170
17.14.3	Description	1170
17.14.4	Restart, fix_modify, output, run start/stop, minimize info	1173
17.14.5	Restrictions	1174
17.14.6	Related commands	1174
17.14.7	Default	1174
17.15	fix ave/correlate/long command	1174
17.15.1	Syntax	1174
17.15.2	Examples	1175
17.15.3	Description	1175
17.15.4	Restart, fix_modify, output, run start/stop, minimize info	1176
17.15.5	Restrictions	1176
17.15.6	Related commands	1176
17.15.7	Default	1176
17.16	fix ave/histo command	1176
17.17	fix ave/histo/weight command	1176
17.17.1	Syntax	1176
17.17.2	Examples	1177
17.17.3	Description	1178
17.17.4	Restart, fix_modify, output, run start/stop, minimize info	1181
17.17.5	Restrictions	1181
17.17.6	Related commands	1181
17.17.7	Default	1181
17.18	fix ave/time command	1181
17.18.1	Syntax	1181
17.18.2	Examples	1182
17.18.3	Description	1183
17.18.4	Restart, fix_modify, output, run start/stop, minimize info	1186
17.18.5	Restrictions	1186
17.18.6	Related commands	1186
17.18.7	Default	1186

17.19	fix aveforce command	1186
17.19.1	Syntax	1186
17.19.2	Examples	1187
17.19.3	Description	1187
17.19.4	Restart, fix_modify, output, run start/stop, minimize info	1188
17.19.5	Restrictions	1188
17.19.6	Related commands	1188
17.19.7	Default	1188
17.20	fix balance command	1188
17.20.1	Syntax	1188
17.20.2	Examples	1189
17.20.3	Description	1189
17.20.4	Restart, fix_modify, output, run start/stop, minimize info	1193
17.20.5	Restrictions	1194
17.20.6	Related commands	1194
17.20.7	Default	1194
17.21	fix bocs command	1194
17.21.1	Syntax	1194
17.21.2	Examples	1194
17.21.3	Description	1194
17.21.4	Restart, fix_modify, output, run start/stop, minimize info	1195
17.21.5	Restrictions	1195
17.21.6	Further information	1196
17.22	fix bond/break command	1196
17.22.1	Syntax	1196
17.22.2	Examples	1196
17.22.3	Description	1196
17.22.4	Restart, fix_modify, output, run start/stop, minimize info	1197
17.22.5	Restrictions	1198
17.22.6	Related commands	1198
17.22.7	Default	1198
17.23	fix bond/create command	1198
17.24	fix bond/create/angle command	1198
17.24.1	Syntax	1198
17.24.2	Examples	1199
17.24.3	Description	1199
17.24.4	Restart, fix_modify, output, run start/stop, minimize info	1201
17.24.5	Restrictions	1201
17.24.6	Related commands	1201
17.24.7	Default	1201
17.25	fix bond/react command	1202
17.25.1	Syntax	1202
17.25.2	Examples	1203
17.25.3	Description	1203
17.25.4	Restart, fix_modify, output, run start/stop, minimize info	1210
17.25.5	Restrictions	1210
17.25.6	Related commands	1210
17.25.7	Default	1210
17.26	fix bond/swap command	1210
17.26.1	Syntax	1210
17.26.2	Examples	1211
17.26.3	Description	1211
17.26.4	Restart, fix_modify, output, run start/stop, minimize info	1213
17.26.5	Restrictions	1213



17.26.6	Related commands	1213
17.26.7	Default	1214
17.27	fix box/relax command	1214
17.27.1	Syntax	1214
17.27.2	Examples	1214
17.27.3	Description	1214
17.27.4	Restart, fix_modify, output, run start/stop, minimize info	1218
17.27.5	Restrictions	1218
17.27.6	Related commands	1218
17.27.7	Default	1219
17.28	fix brownian command	1219
17.29	fix brownian/sphere command	1219
17.30	fix brownian/asphere command	1219
17.30.1	Syntax	1219
17.30.2	Examples	1220
17.30.3	Description	1220
17.30.4	Restart, fix_modify, output, run start/stop, minimize info	1222
17.30.5	Restrictions	1222
17.30.6	Related commands	1222
17.30.7	Default	1222
17.31	fix charge/regulation command	1222
17.31.1	Syntax	1222
17.31.2	Examples	1223
17.31.3	Description	1223
17.31.4	Output	1225
17.31.5	Restrictions	1226
17.31.6	Related commands	1226
17.31.7	Default	1226
17.32	fix cmap command	1226
17.32.1	Syntax	1226
17.32.2	Examples	1227
17.32.3	Description	1227
17.32.4	Restart, fix_modify, output, run start/stop, minimize info	1228
17.32.5	Restrictions	1228
17.32.6	Related commands	1228
17.32.7	Default	1228
17.33	fix colvars command	1229
17.33.1	Syntax	1229
17.33.2	Examples	1229
17.33.3	Description	1229
17.33.4	Restart, fix_modify, output, run start/stop, minimize info	1230
17.33.5	Restrictions	1230
17.33.6	Related commands	1230
17.33.7	Default	1231
17.34	fix controller command	1231
17.34.1	Syntax	1231
17.34.2	Examples	1231
17.34.3	Description	1232
17.34.4	Restart, fix_modify, output, run start/stop, minimize info	1233
17.34.5	Restrictions	1234
17.34.6	Related commands	1234
17.34.7	Default	1234
17.35	fix damping/cundall command	1234
17.35.1	Syntax	1234

17.35.2	Examples	1234
17.35.3	Description	1235
17.35.4	Restart, fix_modify, output, run start/stop, minimize info	1235
17.35.5	Restrictions	1236
17.35.6	Related commands	1236
17.35.7	Default	1236
17.35.8	References	1236
17.36	fix deform command	1236
17.36.1	Syntax	1236
17.36.2	Examples	1238
17.36.3	Description	1238
17.36.4	Restart, fix_modify, output, run start/stop, minimize info	1243
17.36.5	Restrictions	1243
17.36.6	Related commands	1244
17.36.7	Default	1244
17.37	fix deposit command	1244
17.37.1	Syntax	1244
17.37.2	Examples	1245
17.37.3	Description	1245
17.37.4	Restart, fix_modify, output, run start/stop, minimize info	1247
17.37.5	Restrictions	1248
17.37.6	Related commands	1248
17.37.7	Default	1248
17.38	fix dpd/energy command	1248
17.38.1	Syntax	1248
17.38.2	Examples	1248
17.38.3	Description	1248
17.38.4	Restrictions	1249
17.38.5	Related commands	1249
17.38.6	Default	1249
17.39	fix edpd/source command	1249
17.40	fix tdpd/source command	1249
17.40.1	Syntax	1249
17.40.2	Examples	1250
17.40.3	Description	1250
17.40.4	Restart, fix_modify, output, run start/stop, minimize info	1250
17.40.5	Restrictions	1251
17.40.6	Related commands	1251
17.40.7	Default	1251
17.41	fix drag command	1251
17.41.1	Syntax	1251
17.41.2	Examples	1251
17.41.3	Description	1252
17.41.4	Restart, fix_modify, output, run start/stop, minimize info	1252
17.41.5	Restrictions	1252
17.41.6	Related commands	1252
17.41.7	Default	1252
17.42	fix drude command	1252
17.42.1	Syntax	1252
17.42.2	Examples	1253
17.42.3	Description	1253
17.42.4	Restrictions	1253
17.42.5	Related commands	1253
17.42.6	Default	1253

17.43	fix drude/transform/direct command	1253
17.44	fix drude/transform/inverse command	1253
17.44.1	Syntax	1253
17.44.2	Examples	1254
17.44.3	Description	1254
17.44.4	Restart, fix_modify, output, run start/stop, minimize info	1256
17.44.5	Restrictions	1256
17.44.6	Related commands	1256
17.44.7	Default	1256
17.45	fix dt/reset command	1256
17.45.1	Syntax	1256
17.45.2	Examples	1257
17.45.3	Description	1257
17.45.4	Restart, fix_modify, output, run start/stop, minimize info	1257
17.45.5	Restrictions	1258
17.45.6	Related commands	1258
17.45.7	Default	1258
17.46	fix efield command	1258
17.46.1	Syntax	1258
17.46.2	Examples	1258
17.46.3	Description	1258
17.46.4	Restart, fix_modify, output, run start/stop, minimize info	1259
17.46.5	Restrictions	1260
17.46.6	Related commands	1260
17.46.7	Default	1260
17.47	fix ehex command	1260
17.47.1	Syntax	1260
17.47.2	Examples	1261
17.47.3	Description	1261
17.47.4	Restart, fix_modify, output, run start/stop, minimize info	1263
17.47.5	Restrictions	1263
17.47.6	Related commands	1263
17.47.7	Default	1263
17.48	fix electrode/conp command	1263
17.49	fix electrode/conq command	1263
17.50	fix electrode/thermo command	1263
17.50.1	Syntax	1263
17.50.2	Examples	1264
17.50.3	Description	1264
17.51	fix electron/stopping command	1266
17.52	fix electron/stopping/fit command	1266
17.52.1	Syntax	1266
17.52.2	Examples	1267
17.52.3	Description	1267
17.52.4	Restart, fix_modify, output, run start/stop, minimize info	1268
17.52.5	Restrictions	1269
17.52.6	Default	1269
17.53	fix enforce2d command	1269
17.53.1	Syntax	1269
17.53.2	Examples	1269
17.53.3	Description	1270
17.53.4	Restart, fix_modify, output, run start/stop, minimize info	1270
17.53.5	Restrictions	1270
17.53.6	Related commands	1270

17.53.7	Default	1270
17.54	fix eos/cv command	1270
17.54.1	Syntax	1270
17.54.2	Examples	1271
17.54.3	Description	1271
17.54.4	Restrictions	1271
17.54.5	Related commands	1271
17.54.6	Default	1271
17.55	fix eos/table command	1271
17.55.1	Syntax	1271
17.55.2	Examples	1272
17.55.3	Description	1272
17.55.4	Restrictions	1273
17.55.5	Related commands	1273
17.55.6	Default	1273
17.56	fix eos/table/rx command	1273
17.56.1	Syntax	1273
17.56.2	Examples	1274
17.56.3	Description	1274
17.56.4	Restrictions	1276
17.56.5	Related commands	1276
17.56.6	Default	1276
17.57	fix evaporate command	1276
17.57.1	Syntax	1276
17.57.2	Examples	1277
17.57.3	Description	1277
17.57.4	Restart, fix_modify, output, run start/stop, minimize info	1277
17.57.5	Restrictions	1277
17.57.6	Related commands	1278
17.57.7	Default	1278
17.58	fix external command	1278
17.58.1	Syntax	1278
17.58.2	Examples	1278
17.58.3	Description	1278
17.58.4	Restart, fix_modify, output, run start/stop, minimize info	1280
17.58.5	Restrictions	1280
17.58.6	Related commands	1280
17.58.7	Default	1280
17.59	fix ffl command	1280
17.59.1	Syntax	1280
17.59.2	Examples	1281
17.59.3	Description	1281
17.59.4	Restart, fix_modify, output, run start/stop, minimize info	1281
17.59.5	Restrictions	1282
17.59.6	Related commands	1282
17.60	fix filter/corotate command	1282
17.60.1	Syntax	1282
17.60.2	Examples	1282
17.60.3	Description	1283
17.60.4	Restart, fix_modify, output, run start/stop, minimize info	1283
17.60.5	Restrictions	1283
17.60.6	Related commands	1283
17.60.7	Default	1283
17.61	fix flow/gauss command	1283

17.61.1	Syntax	1283
17.61.2	Examples	1284
17.61.3	Description	1284
17.61.4	Restart, fix_modify, output, run start/stop, minimize info	1285
17.61.5	Restrictions	1285
17.61.6	Related commands	1285
17.61.7	Default	1286
17.62	fix freeze command	1286
17.62.1	Syntax	1286
17.62.2	Examples	1286
17.62.3	Description	1286
17.62.4	Restart, fix_modify, output, run start/stop, minimize info	1287
17.62.5	Restrictions	1287
17.62.6	Related commands	1287
17.62.7	Default	1287
17.63	fix gcmc command	1287
17.63.1	Syntax	1287
17.63.2	Examples	1288
17.63.3	Description	1288
17.63.4	Restart, fix_modify, output, run start/stop, minimize info	1292
17.63.5	Restrictions	1293
17.63.6	Related commands	1293
17.63.7	Default	1293
17.64	fix gld command	1294
17.64.1	Syntax	1294
17.64.2	Examples	1294
17.64.3	Description	1294
17.64.4	Restart, fix_modify, output, run start/stop, minimize info	1296
17.64.5	Restrictions	1296
17.64.6	Related commands	1296
17.64.7	Default	1296
17.65	fix gle command	1296
17.65.1	Syntax	1296
17.65.2	Examples	1297
17.65.3	Description	1297
17.65.4	Restart, fix_modify, output, run start/stop, minimize info	1298
17.65.5	Restrictions	1298
17.65.6	Related commands	1298
17.66	fix gravity command	1298
17.66.1	Syntax	1299
17.66.2	Examples	1299
17.66.3	Description	1299
17.66.4	Restart, fix_modify, output, run start/stop, minimize info	1300
17.66.5	Restrictions	1300
17.66.6	Related commands	1300
17.66.7	Default	1301
17.67	fix grem command	1301
17.67.1	Syntax	1301
17.67.2	Examples	1301
17.67.3	Description	1301
17.67.4	Restart, fix_modify, output, run start/stop, minimize info	1302
17.67.5	Restrictions	1302
17.67.6	Related commands	1302
17.67.7	Default	1302

17.68	fix halt command . . . . .	1302
17.68.1	Syntax . . . . .	1302
17.68.2	Examples . . . . .	1303
17.68.3	Description . . . . .	1303
17.68.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1304
17.68.5	Restrictions . . . . .	1305
17.68.6	Related commands . . . . .	1305
17.68.7	Default . . . . .	1305
17.69	fix heat command . . . . .	1305
17.69.1	Syntax . . . . .	1305
17.69.2	Examples . . . . .	1305
17.69.3	Description . . . . .	1305
17.69.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1306
17.69.5	Restrictions . . . . .	1307
17.69.6	Related commands . . . . .	1307
17.69.7	Default . . . . .	1307
17.70	fix hyper/global command . . . . .	1307
17.70.1	Syntax . . . . .	1307
17.70.2	Examples . . . . .	1307
17.70.3	Description . . . . .	1307
17.70.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1309
17.70.5	Restrictions . . . . .	1310
17.70.6	Related commands . . . . .	1310
17.70.7	Default . . . . .	1310
17.71	fix hyper/local command . . . . .	1311
17.71.1	Syntax . . . . .	1311
17.71.2	Examples . . . . .	1311
17.71.3	Description . . . . .	1311
17.71.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1315
17.71.5	Restrictions . . . . .	1317
17.71.6	Related commands . . . . .	1317
17.71.7	Default . . . . .	1317
17.72	fix imd command . . . . .	1318
17.72.1	Syntax . . . . .	1318
17.72.2	Examples . . . . .	1318
17.72.3	Description . . . . .	1318
17.72.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1319
17.72.5	Restrictions . . . . .	1320
17.72.6	Related commands . . . . .	1320
17.72.7	Default . . . . .	1320
17.73	fix indent command . . . . .	1320
17.73.1	Syntax . . . . .	1320
17.73.2	Examples . . . . .	1321
17.73.3	Description . . . . .	1321
17.73.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1322
17.73.5	Restrictions . . . . .	1323
17.73.6	Related commands . . . . .	1323
17.73.7	Default . . . . .	1323
17.74	fix ipi command . . . . .	1323
17.74.1	Syntax . . . . .	1323
17.74.2	Examples . . . . .	1323
17.74.3	Description . . . . .	1324
17.74.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1324
17.74.5	Restrictions . . . . .	1324

17.74.6	Related commands	1324
17.75	fix langevin command	1325
17.75.1	Syntax	1325
17.75.2	Examples	1325
17.75.3	Description	1326
17.75.4	Restart, fix_modify, output, run start/stop, minimize info	1328
17.75.5	Restrictions	1329
17.75.6	Related commands	1329
17.75.7	Default	1329
17.76	fix langevin/drude command	1329
17.76.1	Syntax	1329
17.76.2	Examples	1330
17.76.3	Description	1330
17.76.4	Restart, fix_modify, output, run start/stop, minimize info	1332
17.76.5	Restrictions	1333
17.76.6	Related commands	1333
17.76.7	Default	1333
17.77	fix langevin/eff command	1333
17.77.1	Syntax	1333
17.77.2	Examples	1334
17.77.3	Description	1334
17.77.4	Restart, fix_modify, output, run start/stop, minimize info	1334
17.77.5	Restrictions	1335
17.77.6	Related commands	1335
17.77.7	Default	1335
17.78	fix langevin/spin command	1335
17.78.1	Syntax	1335
17.78.2	Examples	1335
17.78.3	Description	1335
17.78.4	Restart, fix_modify, output, run start/stop, minimize info	1336
17.78.5	Restrictions	1336
17.78.6	Related commands	1336
17.78.7	Default	1336
17.79	fix latte command	1337
17.79.1	Syntax	1337
17.79.2	Examples	1337
17.79.3	Description	1337
17.79.4	Restart, fix_modify, output, run start/stop, minimize info	1338
17.79.5	Restrictions	1339
17.79.6	Related commands	1339
17.79.7	Default	1339
17.80	fix lb/fluid command	1340
17.80.1	Syntax	1340
17.80.2	Examples	1341
17.80.3	Description	1341
17.80.4	Restart, fix_modify, output, run start/stop, minimize info	1344
17.80.5	Restrictions	1345
17.80.6	Related commands	1345
17.80.7	Default	1345
17.81	fix lb/momentum command	1345
17.81.1	Syntax	1345
17.81.2	Examples	1346
17.81.3	Description	1346
17.81.4	Restart, fix_modify, output, run start/stop, minimize info	1346

17.81.5	Restrictions	1346
17.81.6	Related commands	1346
17.81.7	Default	1347
17.82	fix lb/viscous command	1347
17.82.1	Syntax	1347
17.82.2	Examples	1347
17.82.3	Description	1347
17.82.4	Restart, fix_modify, output, run start/stop, minimize info	1347
17.82.5	Restrictions	1348
17.82.6	Related commands	1348
17.82.7	Default	1348
17.83	fix lineforce command	1348
17.83.1	Syntax	1348
17.83.2	Examples	1348
17.83.3	Description	1348
17.83.4	Restart, fix_modify, output, run start/stop, minimize info	1349
17.83.5	Restrictions	1349
17.83.6	Related commands	1349
17.83.7	Default	1349
17.84	fix manifoldforce command	1349
17.84.1	Syntax	1349
17.84.2	Examples	1349
17.84.3	Description	1349
17.84.4	Restart, fix_modify, output, run start/stop, minimize info	1350
17.84.5	Restrictions	1350
17.84.6	Related commands	1350
17.85	fix mdi/qm command	1350
17.85.1	Syntax	1350
17.85.2	Examples	1351
17.85.3	Description	1351
17.85.4	Restart, fix_modify, output, run start/stop, minimize info	1353
17.85.5	Restrictions	1353
17.85.6	Related commands	1354
17.85.7	Default	1354
17.86	fix meso/move command	1354
17.86.1	Syntax	1354
17.86.2	Examples	1354
17.86.3	Description	1355
17.86.4	Restart, fix_modify, output, run start/stop, minimize info	1357
17.86.5	Restrictions	1357
17.86.6	Related commands	1357
17.86.7	Default	1357
17.87	fix_modify AtC commands	1357
17.87.1	fix_modify AtC add_molecule command	1357
17.87.2	fix_modify AtC add_species command	1358
17.87.3	fix_modify AtC atom_element_map command	1359
17.87.4	fix_modify AtC atom_weight command	1360
17.87.5	fix_modify AtC atomic_charge command	1361
17.87.6	fix_modify AtC boundary_dynamics command	1362
17.87.7	fix_modify AtC boundary_faceset command	1363
17.87.8	fix_modify AtC boundary_type command	1364
17.87.9	fix_modify AtC consistent_fe_initialization command	1364
17.87.10	fix_modify AtC control localized_lambda command	1365
17.87.11	fix_modify AtC control momentum command	1366



17.87.12	fix_modify AtC control thermal command . . . . .	1367
17.87.13	fix_modify AtC decomposition command . . . . .	1369
17.87.14	fix_modify AtC extrinsic electron_integration command . . . . .	1370
17.87.15	fix_modify AtC equilibrium_start command . . . . .	1371
17.87.16	fix_modify AtC extrinsic exchange command . . . . .	1371
17.87.17	fix_modify AtC fe_md_boundary command . . . . .	1372
17.87.18	fix_modify AtC filter scale command . . . . .	1373
17.87.19	fix_modify AtC filter type command . . . . .	1374
17.87.20	fix_modify AtC fix command . . . . .	1375
17.87.21	fix_modify AtC fix_flux command . . . . .	1376
17.87.22	fix_modify AtC computes command . . . . .	1376
17.87.23	fix_modify AtC fields command . . . . .	1377
17.87.24	fix_modify AtC gradients command . . . . .	1379
17.87.25	fix_modify AtC kernel command . . . . .	1381
17.87.26	fix_modify AtC on_the_fly command . . . . .	1382
17.87.27	fix_modify AtC rates command . . . . .	1383
17.87.28	fix_modify AtC initial command . . . . .	1384
17.87.29	fix_modify AtC internal_element_set command . . . . .	1385
17.87.30	fix_modify AtC internal_quadrature command . . . . .	1386
17.87.31	fix_modify AtC kernel_bandwidth command . . . . .	1387
17.87.32	fix_modify AtC control lumped_lambda_solve command . . . . .	1388
17.87.33	fix_modify AtC control mask_direction command . . . . .	1388
17.87.34	fix_modify AtC mass_matrix command . . . . .	1389
17.87.35	fix_modify AtC material command . . . . .	1390
17.87.36	fix_modify AtC mesh add_to_nodeset command . . . . .	1391
17.87.37	fix_modify AtC mesh create command . . . . .	1391
17.87.38	fix_modify AtC mesh create_elementset command . . . . .	1392
17.87.39	fix_modify AtC mesh create_faceset box command . . . . .	1393
17.87.40	fix_modify AtC mesh create_faceset plane command . . . . .	1394
17.87.41	fix_modify AtC mesh create_nodeset command . . . . .	1395
17.87.42	fix_modify AtC mesh delete_elements command . . . . .	1396
17.87.43	fix_modify AtC mesh nodeset_to_elementset command . . . . .	1397
17.87.44	fix_modify AtC mesh output command . . . . .	1398
17.87.45	fix_modify AtC mesh quadrature command . . . . .	1398
17.87.46	fix_modify AtC mesh read command . . . . .	1399
17.87.47	fix_modify AtC mesh write command . . . . .	1400
17.87.48	fix_modify AtC output command . . . . .	1401
17.87.49	fix_modify AtC output boundary_integral command . . . . .	1402
17.87.50	fix_modify AtC output contour_integral command . . . . .	1403
17.87.51	fix_modify AtC output nodeset command . . . . .	1404
17.87.52	fix_modify AtC output volume_integral command . . . . .	1405
17.87.53	fix_modify AtC pair_interactions command . . . . .	1405
17.87.54	fix_modify AtC bond_interactions command . . . . .	1405
17.87.55	fix_modify AtC poisson_solver command . . . . .	1406
17.87.56	fix_modify AtC read_restart command . . . . .	1407
17.87.57	fix_modify AtC remove_molecule command . . . . .	1408
17.87.58	fix_modify AtC remove_source command . . . . .	1409
17.87.59	fix_modify AtC remove_species command . . . . .	1409
17.87.60	fix_modify AtC reset_atomic_reference_positions command . . . . .	1410
17.87.61	fix_modify AtC reset_time command . . . . .	1411
17.87.62	fix_modify AtC sample_frequency command . . . . .	1412
17.87.63	fix_modify AtC set reference_potential_energy command . . . . .	1413
17.87.64	fix_modify AtC source command . . . . .	1413
17.87.65	fix_modify AtC source_integration command . . . . .	1414

17.87.66	fix_modify AtC temperature_definition command . . . . .	1415
17.87.67	fix_modify AtC filter command . . . . .	1416
17.87.68	fix_modify AtC time_integration command . . . . .	1417
17.87.69	fix_modify AtC track_displacement command . . . . .	1418
17.87.70	fix_modify AtC unfix command . . . . .	1419
17.87.71	fix_modify AtC unfix_flux command . . . . .	1419
17.87.72	fix_modify AtC write_atom_weights command . . . . .	1420
17.87.73	fix_modify AtC write_restart command . . . . .	1421
17.88	fix mol/swap command . . . . .	1422
17.88.1	Syntax . . . . .	1422
17.88.2	Examples . . . . .	1422
17.88.3	Description . . . . .	1422
17.88.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1424
17.88.5	Restrictions . . . . .	1424
17.88.6	Related commands . . . . .	1424
17.88.7	Default . . . . .	1424
17.89	fix momentum command . . . . .	1424
17.90	fix momentum/chunk command . . . . .	1425
17.90.1	Syntax . . . . .	1425
17.90.2	Examples . . . . .	1425
17.90.3	Description . . . . .	1425
17.90.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1426
17.90.5	Restrictions . . . . .	1426
17.90.6	Related commands . . . . .	1426
17.90.7	Default . . . . .	1426
17.91	fix move command . . . . .	1426
17.91.1	Syntax . . . . .	1426
17.91.2	Examples . . . . .	1427
17.91.3	Description . . . . .	1427
17.91.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1429
17.91.5	Restrictions . . . . .	1430
17.91.6	Related commands . . . . .	1430
17.91.7	Default . . . . .	1430
17.92	fix mscg command . . . . .	1430
17.92.1	Syntax . . . . .	1430
17.92.2	Examples . . . . .	1430
17.92.3	Description . . . . .	1431
17.92.4	Restrictions . . . . .	1432
17.92.5	Related commands . . . . .	1432
17.92.6	Default . . . . .	1432
17.93	fix msst command . . . . .	1432
17.93.1	Syntax . . . . .	1432
17.93.2	Examples . . . . .	1433
17.93.3	Description . . . . .	1433
17.93.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1434
17.93.5	Restrictions . . . . .	1435
17.93.6	Related commands . . . . .	1435
17.93.7	Default . . . . .	1435
17.94	fix mvv/dpd command . . . . .	1435
17.95	fix mvv/edpd command . . . . .	1435
17.96	fix mvv/tdpd command . . . . .	1435
17.96.1	Syntax . . . . .	1435
17.96.2	Examples . . . . .	1436
17.96.3	Description . . . . .	1436

17.96.4	Restart, fix_modify, output, run start/stop, minimize info	1436
17.96.5	Restrictions	1437
17.96.6	Related commands	1437
17.96.7	Default	1437
17.97	fix neb command	1437
17.97.1	Syntax	1437
17.97.2	Examples	1438
17.97.3	Description	1438
17.97.4	Restart, fix_modify, output, run start/stop, minimize info	1440
17.97.5	Restrictions	1440
17.97.6	Related commands	1440
17.97.7	Default	1440
17.98	fix neb/spin command	1440
17.98.1	Syntax	1440
17.98.2	Examples	1441
17.98.3	Description	1441
17.98.4	Restart, fix_modify, output, run start/stop, minimize info	1441
17.98.5	Restrictions	1441
17.98.6	Related commands	1441
17.98.7	Default	1441
17.99	fix nvt command	1442
17.100	fix npt command	1442
17.101	fix nph command	1442
17.101.1	Syntax	1442
17.101.2	Examples	1443
17.101.3	Description	1443
17.101.4	Restart, fix_modify, output, run start/stop, minimize info	1449
17.101.5	Restrictions	1450
17.101.6	Related commands	1450
17.101.7	Default	1450
17.102	fix nvt/eff command	1451
17.103	fix npt/eff command	1451
17.104	fix nph/eff command	1451
17.104.1	Syntax	1451
17.104.2	Examples	1451
17.104.3	Description	1452
17.104.4	Restart, fix_modify, output, run start/stop, minimize info	1452
17.104.5	Restrictions	1452
17.104.6	Related commands	1453
17.104.7	Default	1453
17.105	fix nvt/uef command	1453
17.106	fix npt/uef command	1453
17.106.1	Syntax	1453
17.106.2	Examples	1454
17.106.3	Description	1454
17.106.4	Restart, fix_modify, output, run start/stop, minimize info	1456
17.106.5	Restrictions	1456
17.106.6	Related commands	1456
17.106.7	Default	1456
17.107	fix nph/asphere command	1457
17.107.1	Syntax	1457
17.107.2	Examples	1457
17.107.3	Description	1457
17.107.4	Restart, fix_modify, output, run start/stop, minimize info	1458

17.107.5	Restrictions	1458
17.107.6	Related commands	1459
17.107.7	Default	1459
17.108	fix nph/body command	1459
17.108.1	Syntax	1459
17.108.2	Examples	1459
17.108.3	Description	1459
17.108.4	Restart, fix_modify, output, run start/stop, minimize info	1460
17.108.5	Restrictions	1461
17.108.6	Related commands	1461
17.108.7	Default	1461
17.109	fix nph/sphere command	1461
17.109.1	Syntax	1461
17.109.2	Examples	1461
17.109.3	Description	1462
17.109.4	Restart, fix_modify, output, run start/stop, minimize info	1463
17.109.5	Restrictions	1463
17.109.6	Related commands	1463
17.109.7	Default	1463
17.110	fix nphug command	1463
17.110.1	Syntax	1463
17.110.2	Examples	1464
17.110.3	Description	1464
17.110.4	Restart, fix_modify, output, run start/stop, minimize info	1466
17.110.5	Restrictions	1466
17.110.6	Related commands	1466
17.110.7	Default	1466
17.111	fix npt/asphere command	1467
17.111.1	Syntax	1467
17.111.2	Examples	1467
17.111.3	Description	1467
17.111.4	Restart, fix_modify, output, run start/stop, minimize info	1468
17.111.5	Restrictions	1469
17.111.6	Related commands	1469
17.111.7	Default	1469
17.112	fix npt/body command	1469
17.112.1	Syntax	1469
17.112.2	Examples	1469
17.112.3	Description	1469
17.112.4	Restart, fix_modify, output, run start/stop, minimize info	1471
17.112.5	Restrictions	1471
17.112.6	Related commands	1471
17.112.7	Default	1471
17.113	fix npt/cauchy command	1471
17.113.1	Syntax	1471
17.113.2	Examples	1472
17.113.3	Description	1472
17.113.4	Restart, fix_modify, output, run start/stop, minimize info	1477
17.113.5	Restrictions	1478
17.113.6	Related commands	1479
17.113.7	Default	1479
17.114	fix npt/sphere command	1479
17.114.1	Syntax	1479
17.114.2	Examples	1479

17.114.3	Description	1480
17.114.4	Restart, fix_modify, output, run start/stop, minimize info	1481
17.114.5	Restrictions	1481
17.114.6	Related commands	1481
17.114.7	Default	1481
17.115	fix numdiff command	1482
17.115.1	Syntax	1482
17.115.2	Examples	1482
17.115.3	Description	1482
17.115.4	Restart, fix_modify, output, run start/stop, minimize info	1483
17.115.5	Restrictions	1483
17.115.6	Related commands	1483
17.115.7	Default	1483
17.116	fix numdiff/virial command	1483
17.116.1	Syntax	1483
17.116.2	Examples	1484
17.116.3	Description	1484
17.116.4	Restart, fix_modify, output, run start/stop, minimize info	1484
17.116.5	Restrictions	1485
17.116.6	Related commands	1485
17.116.7	Default	1485
17.117	fix nve command	1485
17.117.1	Syntax	1485
17.117.2	Examples	1485
17.117.3	Description	1485
17.117.4	Restart, fix_modify, output, run start/stop, minimize info	1486
17.117.5	Restrictions	1486
17.117.6	Related commands	1486
17.117.7	Default	1486
17.118	fix nve/asphere command	1486
17.118.1	Syntax	1486
17.118.2	Examples	1486
17.118.3	Description	1487
17.118.4	Restart, fix_modify, output, run start/stop, minimize info	1487
17.118.5	Restrictions	1487
17.118.6	Related commands	1487
17.118.7	Default	1487
17.119	fix nve/asphere/noforce command	1488
17.119.1	Syntax	1488
17.119.2	Examples	1488
17.119.3	Description	1488
17.119.4	Restart, fix_modify, output, run start/stop, minimize info	1488
17.119.5	Restrictions	1488
17.119.6	Related commands	1489
17.119.7	Default	1489
17.120	fix nve/awpmd command	1489
17.120.1	Syntax	1489
17.120.2	Examples	1489
17.120.3	Description	1489
17.120.4	Restart, fix_modify, output, run start/stop, minimize info	1489
17.120.5	Restrictions	1490
17.120.6	Related commands	1490
17.120.7	Default	1490
17.121	fix nve/body command	1490

17.121.1	Syntax . . . . .	1490
17.121.2	Examples . . . . .	1490
17.121.3	Description . . . . .	1490
17.121.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1490
17.121.5	Restrictions . . . . .	1491
17.121.6	Related commands . . . . .	1491
17.121.7	Default . . . . .	1491
17.122	fix nve/bpm/sphere command . . . . .	1491
17.122.1	Syntax . . . . .	1491
17.122.2	Examples . . . . .	1491
17.122.3	Description . . . . .	1491
17.122.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1492
17.122.5	Restrictions . . . . .	1492
17.122.6	Related commands . . . . .	1492
17.122.7	Default . . . . .	1492
17.123	fix nve/dot command . . . . .	1492
17.123.1	Syntax . . . . .	1492
17.123.2	Examples . . . . .	1492
17.123.3	Description . . . . .	1493
17.123.4	Restrictions . . . . .	1493
17.123.5	Related commands . . . . .	1493
17.123.6	Default . . . . .	1493
17.124	fix nve/dotc/langevin command . . . . .	1493
17.124.1	Syntax . . . . .	1493
17.124.2	Examples . . . . .	1494
17.124.3	Description . . . . .	1494
17.124.4	Restrictions . . . . .	1495
17.124.5	Related commands . . . . .	1495
17.124.6	Default . . . . .	1495
17.125	fix nve/eff command . . . . .	1496
17.125.1	Syntax . . . . .	1496
17.125.2	Examples . . . . .	1496
17.125.3	Description . . . . .	1496
17.125.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1496
17.125.5	Restrictions . . . . .	1496
17.125.6	Related commands . . . . .	1496
17.125.7	Default . . . . .	1496
17.126	fix nve/limit command . . . . .	1497
17.126.1	Syntax . . . . .	1497
17.126.2	Examples . . . . .	1497
17.126.3	Description . . . . .	1497
17.126.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1497
17.126.5	Restrictions . . . . .	1498
17.126.6	Related commands . . . . .	1498
17.126.7	Default . . . . .	1498
17.127	fix nve/line command . . . . .	1498
17.127.1	Syntax . . . . .	1498
17.127.2	Examples . . . . .	1498
17.127.3	Description . . . . .	1498
17.127.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1498
17.127.5	Restrictions . . . . .	1499
17.127.6	Related commands . . . . .	1499
17.127.7	Default . . . . .	1499
17.128	fix nve/manifold/rattle command . . . . .	1499

17.128.1	Syntax . . . . .	1499
17.128.2	Examples . . . . .	1499
17.128.3	Description . . . . .	1500
17.128.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1500
17.128.5	Restrictions . . . . .	1500
17.128.6	Related commands . . . . .	1500
17.128.7	Default . . . . .	1500
17.129	fix nve/noforce command . . . . .	1501
17.129.1	Syntax . . . . .	1501
17.129.2	Examples . . . . .	1501
17.129.3	Description . . . . .	1501
17.129.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1501
17.129.5	Restrictions . . . . .	1501
17.129.6	Related commands . . . . .	1501
17.129.7	Default . . . . .	1502
17.130	fix nve/sphere command . . . . .	1502
17.130.1	Syntax . . . . .	1502
17.130.2	Examples . . . . .	1502
17.130.3	Description . . . . .	1502
17.130.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1503
17.130.5	Restrictions . . . . .	1503
17.130.6	Related commands . . . . .	1503
17.130.7	Default . . . . .	1503
17.131	fix nve/spin command . . . . .	1503
17.131.1	Syntax . . . . .	1503
17.131.2	Examples . . . . .	1504
17.131.3	Description . . . . .	1504
17.131.4	Restrictions . . . . .	1505
17.131.5	Related commands . . . . .	1505
17.131.6	Default . . . . .	1505
17.132	fix nve/tri command . . . . .	1505
17.132.1	Syntax . . . . .	1505
17.132.2	Examples . . . . .	1505
17.132.3	Description . . . . .	1505
17.132.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1506
17.132.5	Restrictions . . . . .	1506
17.132.6	Related commands . . . . .	1506
17.132.7	Default . . . . .	1506
17.133	fix nvk command . . . . .	1506
17.133.1	Syntax . . . . .	1506
17.133.2	Examples . . . . .	1506
17.133.3	Description . . . . .	1506
17.133.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1507
17.133.5	Restrictions . . . . .	1507
17.133.6	Related commands . . . . .	1507
17.133.7	Default . . . . .	1507
17.134	fix nvt/asphere command . . . . .	1507
17.134.1	Syntax . . . . .	1507
17.134.2	Examples . . . . .	1508
17.134.3	Description . . . . .	1508
17.134.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1509
17.134.5	Restrictions . . . . .	1509
17.134.6	Related commands . . . . .	1509
17.134.7	Default . . . . .	1509

17.135	fix nvt/body command	1509
17.135.1	Syntax	1509
17.135.2	Examples	1510
17.135.3	Description	1510
17.135.4	Restart, fix_modify, output, run start/stop, minimize info	1511
17.135.5	Restrictions	1511
17.135.6	Related commands	1511
17.135.7	Default	1511
17.136	fix nvt/manifold/rattle command	1511
17.136.1	Syntax	1511
17.136.2	Examples	1512
17.136.3	Description	1512
17.136.4	Restart, fix_modify, output, run start/stop, minimize info	1512
17.136.5	Restrictions	1512
17.136.6	Related commands	1512
17.137	fix nvt/sllod command	1513
17.137.1	Syntax	1513
17.137.2	Examples	1513
17.137.3	Description	1513
17.137.4	Restart, fix_modify, output, run start/stop, minimize info	1514
17.137.5	Restrictions	1515
17.137.6	Related commands	1515
17.137.7	Default	1515
17.138	fix nvt/sllod/eff command	1515
17.138.1	Syntax	1515
17.138.2	Examples	1515
17.138.3	Description	1515
17.138.4	Restart, fix_modify, output, run start/stop, minimize info	1516
17.138.5	Restrictions	1516
17.138.6	Related commands	1516
17.138.7	Default	1516
17.139	fix nvt/sphere command	1516
17.139.1	Syntax	1516
17.139.2	Examples	1517
17.139.3	Description	1517
17.139.4	Restart, fix_modify, output, run start/stop, minimize info	1518
17.139.5	Restrictions	1518
17.139.6	Related commands	1518
17.139.7	Default	1519
17.140	fix oneway command	1519
17.140.1	Syntax	1519
17.140.2	Examples	1519
17.140.3	Description	1519
17.140.4	Restart, fix_modify, output, run start/stop, minimize info	1519
17.140.5	Restrictions	1520
17.140.6	Related commands	1520
17.140.7	Default	1520
17.141	fix orient/fcc command	1520
17.142	fix orient/bcc command	1520
17.142.1	Syntax	1520
17.142.2	Examples	1520
17.142.3	Description	1521
17.142.4	Restart, fix_modify, output, run start/stop, minimize info	1522
17.142.5	Restrictions	1522



17.142.6	Related commands	1523
17.142.7	Default	1523
17.143	fix orient/eco command	1523
17.143.1	Examples	1524
17.143.2	Description	1524
17.143.3	Restart, fix_modify, output, run start/stop, minimize info	1525
17.143.4	Restrictions	1525
17.143.5	Related commands	1525
17.143.6	Default	1525
17.144	fix pafi command	1526
17.144.1	Syntax	1526
17.144.2	Examples	1526
17.144.3	Description	1526
17.144.4	Restart, fix_modify, output, run start/stop, minimize info	1527
17.144.5	Restrictions	1527
17.144.6	Default	1527
17.145	fix pair command	1527
17.145.1	Syntax	1527
17.145.2	Examples	1528
17.145.3	Description	1528
17.145.4	Restart, fix_modify, output, run start/stop, minimize info	1528
17.145.5	Restrictions	1529
17.145.6	Related commands	1529
17.145.7	Default	1529
17.146	fix phonon command	1529
17.146.1	Syntax	1529
17.146.2	Examples	1530
17.146.3	Description	1530
17.146.4	Restart, fix_modify, output, run start/stop, minimize info	1531
17.146.5	Restrictions	1531
17.146.6	Related commands	1531
17.146.7	Default	1531
17.147	fix pimd command	1532
17.147.1	Syntax	1532
17.147.2	Examples	1532
17.147.3	Description	1532
17.147.4	Restrictions	1535
17.147.5	Default	1535
17.148	fix planeforce command	1535
17.148.1	Syntax	1535
17.148.2	Examples	1535
17.148.3	Description	1536
17.148.4	Restart, fix_modify, output, run start/stop, minimize info	1536
17.148.5	Restrictions	1536
17.148.6	Related commands	1536
17.148.7	Default	1536
17.149	fix plumed command	1536
17.149.1	Syntax	1536
17.149.2	Examples	1537
17.149.3	Description	1537
17.149.4	Restart, fix_modify, output, run start/stop, minimize info	1537
17.149.5	Restrictions	1538
17.149.6	Related commands	1538
17.149.7	Default	1538

17.150	fix poems command	1538
17.150.1	Syntax	1538
17.150.2	Examples	1538
17.150.3	Description	1539
17.150.4	Restart, fix_modify, output, run start/stop, minimize info	1540
17.150.5	Restrictions	1540
17.150.6	Related commands	1540
17.150.7	Default	1540
17.151	fix polarize/bem/gmres command	1540
17.152	fix polarize/bem/icc command	1540
17.153	fix polarize/functional command	1540
17.153.1	Syntax	1540
17.153.2	Examples	1541
17.153.3	Description	1541
17.153.4	Restart, fix_modify, output, run start/stop, minimize info	1542
17.153.5	Restrictions	1542
17.153.6	Related commands	1542
17.153.7	Default	1543
17.154	fix pour command	1543
17.154.1	Syntax	1543
17.154.2	Examples	1544
17.154.3	Description	1544
17.154.4	Restart, fix_modify, output, run start/stop, minimize info	1546
17.154.5	Restrictions	1546
17.154.6	Related commands	1546
17.154.7	Default	1546
17.155	fix precession/spin command	1547
17.155.1	Syntax	1547
17.155.2	Examples	1547
17.155.3	Description	1547
17.155.4	Restart, fix_modify, output, run start/stop, minimize info	1549
17.155.5	Restrictions	1549
17.155.6	Related commands	1549
17.155.7	Default	1549
17.156	fix press/berendsen command	1550
17.156.1	Syntax	1550
17.156.2	Examples	1550
17.156.3	Description	1550
17.156.4	Restart, fix_modify, output, run start/stop, minimize info	1552
17.156.5	Restrictions	1552
17.156.6	Related commands	1552
17.156.7	Default	1553
17.157	fix print command	1553
17.157.1	Syntax	1553
17.157.2	Examples	1553
17.157.3	Description	1553
17.157.4	Restart, fix_modify, output, run start/stop, minimize info	1554
17.157.5	Restrictions	1554
17.157.6	Related commands	1554
17.157.7	Default	1555
17.158	fix propel/self command	1555
17.158.1	Syntax	1555
17.158.2	Examples	1555
17.158.3	Description	1555

17.158.4Restart, fix_modify, output, run start/stop, minimize info	1556
17.158.5Restrictions	1556
17.158.6Related commands	1557
17.158.7Default	1557
17.159fix property/atom command	1557
17.159.1Syntax	1557
17.159.2Examples	1557
17.159.3Description	1558
17.159.4Restart, fix_modify, output, run start/stop, minimize info	1561
17.159.5Restrictions	1561
17.159.6Related commands	1561
17.159.7Default	1561
17.160fix python/invoke command	1561
17.160.1Syntax	1561
17.160.2Examples	1562
17.160.3Description	1562
17.160.4Restrictions	1562
17.160.5Related commands	1563
17.161fix python/move command	1563
17.161.1Syntax	1563
17.161.2Examples	1563
17.161.3Description	1563
17.161.4Restart, fix_modify, output, run start/stop, minimize info	1564
17.161.5Restrictions	1564
17.161.6Related commands	1564
17.161.7Default	1564
17.162fix qbmsst command	1564
17.162.1Syntax	1564
17.162.2Examples	1565
17.162.3Description	1566
17.162.4Restart, fix_modify, output, run start/stop, minimize info	1567
17.162.5Restrictions	1568
17.162.6Related commands	1568
17.162.7Default	1568
17.163fix qeq/point command	1568
17.164fix qeq/shielded command	1568
17.165fix qeq/slater command	1568
17.166fix qeq/dynamic command	1568
17.167fix qeq/fire command	1568
17.167.1Syntax	1568
17.167.2Examples	1569
17.167.3Description	1569
17.167.4Restart, fix_modify, output, run start/stop, minimize info	1571
17.167.5Restrictions	1571
17.167.6Related commands	1571
17.167.7Default	1572
17.168fix qeq/comb command	1572
17.168.1Syntax	1572
17.168.2Examples	1572
17.168.3Description	1572
17.168.4Restart, fix_modify, output, run start/stop, minimize info	1573
17.168.5Restrictions	1573
17.168.6Related commands	1574
17.168.7Default	1574

17.169	fix qeq/reaxff command	1574
17.169.1	Syntax	1574
17.169.2	Examples	1574
17.169.3	Description	1575
17.169.4	Restart, fix_modify, output, run start/stop, minimize info	1575
17.169.5	Restrictions	1576
17.169.6	Related commands	1576
17.169.7	Default	1576
17.170	fix qmmm command	1576
17.170.1	Syntax	1576
17.170.2	Examples	1577
17.170.3	Description	1577
17.170.4	Restart, fix_modify, output, run start/stop, minimize info	1577
17.170.5	Restrictions	1577
17.170.6	Related commands	1577
17.170.7	Default	1577
17.171	fix qtb command	1578
17.171.1	Syntax	1578
17.171.2	Examples	1578
17.171.3	Description	1578
17.171.4	Restart, fix_modify, output, run start/stop, minimize info	1579
17.171.5	Restrictions	1580
17.171.6	Related commands	1580
17.171.7	Default	1580
17.172	fix reaxff/bonds command	1580
17.172.1	Syntax	1580
17.172.2	Examples	1580
17.172.3	Description	1581
17.172.4	Restart, fix_modify, output, run start/stop, minimize info	1581
17.172.5	Restrictions	1582
17.172.6	Related commands	1582
17.172.7	Default	1582
17.173	fix reaxff/species command	1582
17.173.1	Syntax	1582
17.173.2	Examples	1583
17.173.3	Description	1583
17.173.4	Restart, fix_modify, output, run start/stop, minimize info	1584
17.173.5	Restrictions	1585
17.173.6	Related commands	1585
17.173.7	Default	1585
17.174	fix recenter command	1585
17.174.1	Syntax	1585
17.174.2	Examples	1585
17.174.3	Description	1586
17.174.4	Restart, fix_modify, output, run start/stop, minimize info	1586
17.174.5	Restrictions	1587
17.174.6	Related commands	1587
17.174.7	Default	1587
17.175	fix restrain command	1587
17.175.1	Syntax	1587
17.175.2	Examples	1588
17.175.3	Description	1588
17.175.4	Restart, fix_modify, output, run start/stop, minimize info	1590
17.175.5	Restrictions	1591

17.175.6	Related commands	1591
17.175.7	Default	1591
17.176	fix rhok command	1591
17.176.1	Syntax	1591
17.176.2	Examples	1591
17.176.3	Description	1591
17.176.4	Restart, fix_modify, output, run start/stop, minimize info	1592
17.176.5	Restrictions	1592
17.176.6	Related commands	1592
17.176.7	Default	1592
17.177	fix rigid command	1592
17.178	fix rigid/nve command	1592
17.179	fix rigid/nvt command	1592
17.180	fix rigid/npt command	1593
17.181	fix rigid/nph command	1593
17.182	fix rigid/small command	1593
17.183	fix rigid/nve/small command	1593
17.184	fix rigid/nvt/small command	1593
17.185	fix rigid/npt/small command	1593
17.186	fix rigid/nph/small command	1593
17.186.1	Syntax	1593
17.186.2	Examples	1594
17.186.3	Description	1595
17.186.4	Restart, fix_modify, output, run start/stop, minimize info	1602
17.186.5	Restrictions	1603
17.186.6	Related commands	1603
17.186.7	Default	1603
17.187	fix rigid/meso command	1604
17.187.1	Syntax	1604
17.187.2	Examples	1604
17.187.3	Description	1604
17.187.4	Restart, fix_modify, output, run start/stop, minimize info	1607
17.187.5	Restrictions	1608
17.187.6	Related commands	1608
17.187.7	Default	1608
17.188	fix rx command	1608
17.188.1	Syntax	1608
17.188.2	Examples	1609
17.188.3	Description	1609
17.188.4	Restrictions	1611
17.188.5	Related commands	1611
17.188.6	Default	1611
17.189	fix saed/vtk command	1611
17.189.1	Syntax	1611
17.189.2	Examples	1612
17.189.3	Description	1612
17.189.4	Restart, fix_modify, output, run start/stop, minimize info	1614
17.189.5	Restrictions	1614
17.189.6	Related commands	1614
17.189.7	Default	1614
17.190	fix setforce command	1614
17.191	fix setforce/spin command	1614
17.191.1	Syntax	1614
17.191.2	Examples	1615

17.191.3	Description	1615
17.191.4	Restart, fix_modify, output, run start/stop, minimize info	1616
17.191.5	Restrictions	1616
17.191.6	Related commands	1616
17.191.7	Default	1616
17.192	fix shake command	1616
17.193	fix rattle command	1616
17.193.1	Syntax	1616
17.193.2	Examples	1617
17.193.3	Description	1617
17.193.4	Restart, fix_modify, output, run start/stop, minimize info	1619
17.193.5	Restrictions	1620
17.193.6	Related commands	1620
17.193.7	Default	1620
17.194	fix shardlow command	1620
17.194.1	Syntax	1620
17.194.2	Examples	1621
17.194.3	Description	1621
17.194.4	Restrictions	1621
17.194.5	Related commands	1622
17.194.6	Default	1622
17.195	fix smd command	1622
17.195.1	Syntax	1622
17.195.2	Examples	1623
17.195.3	Description	1623
17.195.4	Restart, fix_modify, output, run start/stop, minimize info	1623
17.195.5	Restrictions	1624
17.195.6	Related commands	1624
17.195.7	Default	1624
17.196	fix smd/adjust_dt command	1624
17.196.1	Syntax	1624
17.196.2	Examples	1625
17.196.3	Description	1625
17.196.4	Restart, fix_modify, output, run start/stop, minimize info	1625
17.196.5	Restrictions	1625
17.196.6	Related commands	1625
17.196.7	Default	1625
17.197	fix smd/integrate_tlsph command	1625
17.197.1	Syntax	1625
17.197.2	Examples	1626
17.197.3	Description	1626
17.197.4	Restart, fix_modify, output, run start/stop, minimize info	1626
17.197.5	Restrictions	1626
17.197.6	Related commands	1626
17.197.7	Default	1626
17.198	fix smd/integrate_ulsph command	1626
17.198.1	Syntax	1626
17.198.2	Examples	1627
17.198.3	Description	1627
17.198.4	Restart, fix_modify, output, run start/stop, minimize info	1627
17.198.5	Restrictions	1627
17.198.6	Related commands	1627
17.198.7	Default	1627
17.199	fix smd/move_tri_surf command	1628

17.199.1	Syntax . . . . .	1628
17.199.2	Examples . . . . .	1628
17.199.3	Description . . . . .	1628
17.199.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1629
17.199.5	Restrictions . . . . .	1629
17.199.6	Related commands . . . . .	1629
17.199.7	Default . . . . .	1629
17.200	fix smd/setvel command . . . . .	1629
17.200.1	Syntax . . . . .	1629
17.200.2	Examples . . . . .	1629
17.200.3	Description . . . . .	1630
17.200.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1630
17.200.5	Restrictions . . . . .	1630
17.200.6	Related commands . . . . .	1630
17.200.7	Default . . . . .	1630
17.201	fix smd/wall_surface command . . . . .	1631
17.201.1	Syntax . . . . .	1631
17.201.2	Examples . . . . .	1631
17.201.3	Description . . . . .	1631
17.201.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1631
17.201.5	Restrictions . . . . .	1631
17.201.6	Related commands . . . . .	1632
17.201.7	Default . . . . .	1632
17.202	fix sph command . . . . .	1632
17.202.1	Syntax . . . . .	1632
17.202.2	Examples . . . . .	1632
17.202.3	Description . . . . .	1632
17.202.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1632
17.202.5	Restrictions . . . . .	1632
17.202.6	Related commands . . . . .	1633
17.202.7	Default . . . . .	1633
17.203	fix sph/stationary command . . . . .	1633
17.203.1	Syntax . . . . .	1633
17.203.2	Examples . . . . .	1633
17.203.3	Description . . . . .	1633
17.203.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1633
17.203.5	Restrictions . . . . .	1633
17.203.6	Related commands . . . . .	1634
17.203.7	Default . . . . .	1634
17.204	fix spring command . . . . .	1634
17.204.1	Syntax . . . . .	1634
17.204.2	Examples . . . . .	1634
17.204.3	Description . . . . .	1634
17.204.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1635
17.204.5	Restrictions . . . . .	1636
17.204.6	Related commands . . . . .	1636
17.204.7	Default . . . . .	1636
17.205	fix spring/chunk command . . . . .	1636
17.205.1	Syntax . . . . .	1636
17.205.2	Examples . . . . .	1636
17.205.3	Description . . . . .	1636
17.205.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1637
17.205.5	Restrictions . . . . .	1637
17.205.6	Related commands . . . . .	1637

17.205.7Default	1637
17.206fix spring/rg command	1637
17.206.1Syntax	1637
17.206.2Examples	1638
17.206.3Description	1638
17.206.4Restart, fix_modify, output, run start/stop, minimize info	1638
17.206.5Restrictions	1639
17.206.6Related commands	1639
17.206.7Default	1639
17.207fix spring/self command	1639
17.207.1Syntax	1639
17.207.2Examples	1639
17.207.3Description	1639
17.207.4Restart, fix_modify, output, run start/stop, minimize info	1640
17.207.5Restrictions	1640
17.207.6Related commands	1640
17.207.7Default	1640
17.208fix srd command	1640
17.208.1Syntax	1640
17.208.2Examples	1641
17.208.3Description	1641
17.208.4Restart, fix_modify, output, run start/stop, minimize info	1645
17.208.5Restrictions	1645
17.208.6Related commands	1645
17.208.7Default	1645
17.209fix store/force command	1646
17.209.1Syntax	1646
17.209.2Examples	1646
17.209.3Description	1646
17.209.4Restart, fix_modify, output, run start/stop, minimize info	1646
17.209.5Restrictions	1647
17.209.6Related commands	1647
17.209.7Default	1647
17.210fix store/state command	1647
17.210.1Syntax	1647
17.210.2Examples	1648
17.210.3Description	1648
17.210.4Restart, fix_modify, output, run start/stop, minimize info	1649
17.210.5Restrictions	1649
17.210.6Related commands	1649
17.210.7Default	1649
17.211fix temp/berendsen command	1649
17.211.1Syntax	1649
17.211.2Examples	1650
17.211.3Description	1650
17.211.4Restart, fix_modify, output, run start/stop, minimize info	1651
17.211.5Restrictions	1651
17.211.6Related commands	1651
17.211.7Default	1651
17.212fix temp/csvr command	1652
17.213fix temp/csld command	1652
17.213.1Syntax	1652
17.213.2Examples	1652
17.213.3Description	1652



17.213.4	Restart, fix_modify, output, run start/stop, minimize info	1653
17.213.5	Restrictions	1654
17.213.6	Related commands	1654
17.213.7	Default	1654
17.214	fix temp/rescale command	1654
17.214.1	Syntax	1654
17.214.2	Examples	1655
17.214.3	Description	1655
17.214.4	Restart, fix_modify, output, run start/stop, minimize info	1656
17.214.5	Restrictions	1656
17.214.6	Related commands	1656
17.214.7	Default	1656
17.215	fix temp/rescale/eff command	1657
17.215.1	Syntax	1657
17.215.2	Examples	1657
17.215.3	Description	1657
17.215.4	Restart, fix_modify, output, run start/stop, minimize info	1657
17.215.5	Restrictions	1658
17.215.6	Related commands	1658
17.215.7	Default	1658
17.216	fix tfmc command	1658
17.216.1	Syntax	1658
17.216.2	Examples	1658
17.216.3	Description	1659
17.216.4	Restart, fix_modify, output, run start/stop, minimize info	1660
17.216.5	Restrictions	1660
17.216.6	Related commands	1660
17.216.7	Default	1660
17.217	fix tgnvt/drude command	1660
17.218	fix tgnpt/drude command	1660
17.218.1	Syntax	1660
17.218.2	Examples	1661
17.218.3	Description	1661
17.218.4	Restart, fix_modify, output, run start/stop, minimize info	1663
17.218.5	Restrictions	1664
17.218.6	Related commands	1664
17.218.7	Default	1664
17.219	fix thermal/conductivity command	1664
17.219.1	Syntax	1664
17.219.2	Examples	1665
17.219.3	Description	1665
17.219.4	Restart, fix_modify, output, run start/stop, minimize info	1666
17.219.5	Restrictions	1666
17.219.6	Related commands	1666
17.219.7	Default	1666
17.220	fix ti/spring command	1667
17.220.1	Syntax	1667
17.220.2	Example	1667
17.220.3	Description	1667
17.220.4	Restart, fix_modify, output, run start/stop, minimize info	1668
17.220.5	Related commands	1669
17.220.6	Restrictions	1669
17.220.7	Default	1669
17.221	fix tmd command	1669

17.221.1	Syntax . . . . .	1669
17.221.2	Examples . . . . .	1669
17.221.3	Description . . . . .	1669
17.221.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1670
17.221.5	Restrictions . . . . .	1671
17.221.6	Related commands . . . . .	1671
17.221.7	Default . . . . .	1671
17.222	fix ttm command . . . . .	1671
17.223	fix ttm/grid command . . . . .	1671
17.224	fix ttm/mod command . . . . .	1671
17.224.1	Syntax . . . . .	1671
17.224.2	Examples . . . . .	1672
17.224.3	Description . . . . .	1672
17.224.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1675
17.224.5	Restrictions . . . . .	1676
17.224.6	Related commands . . . . .	1676
17.224.7	Default . . . . .	1676
17.225	fix tune/kSPACE command . . . . .	1676
17.225.1	Syntax . . . . .	1676
17.225.2	Examples . . . . .	1677
17.225.3	Description . . . . .	1677
17.225.4	Restrictions . . . . .	1677
17.225.5	Related commands . . . . .	1678
17.225.6	Default . . . . .	1678
17.226	fix vector command . . . . .	1678
17.226.1	Syntax . . . . .	1678
17.226.2	Examples . . . . .	1678
17.226.3	Description . . . . .	1678
17.226.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1679
17.226.5	Restrictions . . . . .	1680
17.226.6	Related commands . . . . .	1680
17.226.7	Default . . . . .	1680
17.227	fix viscosity command . . . . .	1680
17.227.1	Syntax . . . . .	1680
17.227.2	Examples . . . . .	1681
17.227.3	Description . . . . .	1681
17.227.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1682
17.227.5	Restrictions . . . . .	1682
17.227.6	Related commands . . . . .	1682
17.227.7	Default . . . . .	1682
17.228	fix viscous command . . . . .	1683
17.228.1	Syntax . . . . .	1683
17.228.2	Examples . . . . .	1683
17.228.3	Description . . . . .	1683
17.228.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1684
17.228.5	Restrictions . . . . .	1684
17.228.6	Related commands . . . . .	1684
17.228.7	Default . . . . .	1684
17.229	fix viscous/sphere command . . . . .	1684
17.229.1	Syntax . . . . .	1684
17.229.2	Examples . . . . .	1685
17.229.3	Description . . . . .	1685
17.229.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	1685
17.229.5	Restrictions . . . . .	1685

17.229.6	Related commands	1686
17.229.7	Default	1686
17.230	fix wall/lj93 command	1686
17.231	fix wall/lj126 command	1686
17.232	fix wall/lj1043 command	1686
17.233	fix wall/colloid command	1686
17.234	fix wall/harmonic command	1686
17.235	fix wall/morse command	1686
17.235.1	Syntax	1686
17.235.2	Examples	1687
17.235.3	Description	1687
17.235.4	Restart, fix_modify, output, run start/stop, minimize info	1690
17.235.5	Restrictions	1691
17.235.6	Related commands	1691
17.235.7	Default	1691
17.236	fix wall/body/polygon command	1691
17.236.1	Syntax	1691
17.236.2	Examples	1692
17.236.3	Description	1692
17.236.4	Restart, fix_modify, output, run start/stop, minimize info	1692
17.236.5	Restrictions	1692
17.236.6	Related commands	1693
17.236.7	Default	1693
17.237	fix wall/body/polyhedron command	1693
17.237.1	Syntax	1693
17.237.2	Examples	1693
17.237.3	Description	1694
17.237.4	Restart, fix_modify, output, run start/stop, minimize info	1694
17.237.5	Restrictions	1694
17.237.6	Related commands	1694
17.237.7	Default	1695
17.238	fix wall/ees command	1695
17.239	fix wall/region/ees command	1695
17.239.1	Syntax	1695
17.239.2	Examples	1695
17.239.3	Description	1696
17.239.4	Restart, fix_modify, output, run start/stop, minimize info	1697
17.239.5	Restrictions	1697
17.239.6	Related commands	1697
17.239.7	Default	1697
17.240	fix wall/gran command	1697
17.240.1	Syntax	1697
17.240.2	Examples	1698
17.240.3	Description	1699
17.240.4	Restart, fix_modify, output, run start/stop, minimize info	1700
17.240.5	Restrictions	1700
17.240.6	Related commands	1700
17.240.7	Default	1701
17.241	fix wall/gran/region command	1701
17.241.1	Syntax	1701
17.241.2	Examples	1701
17.241.3	Description	1702
17.241.4	Restart, fix_modify, output, run start/stop, minimize info	1704
17.241.5	Restrictions	1704

17.241.6	Related commands	1704
17.241.7	Default	1705
17.242	fix wall/piston command	1705
17.242.1	Syntax	1705
17.242.2	Examples	1705
17.242.3	Description	1705
17.242.4	Restart, fix_modify, output, run start/stop, minimize info	1706
17.242.5	Restrictions	1706
17.242.6	Related commands	1706
17.242.7	Default	1706
17.243	fix wall/reflect command	1707
17.243.1	Syntax	1707
17.243.2	Examples	1707
17.243.3	Description	1707
17.243.4	Restart, fix_modify, output, run start/stop, minimize info	1709
17.243.5	Restrictions	1709
17.243.6	Related commands	1709
17.243.7	Default	1709
17.244	fix wall/reflect/stochastic command	1709
17.244.1	Syntax	1709
17.244.2	Examples	1710
17.244.3	Description	1710
17.244.4	Restrictions	1711
17.244.5	Related commands	1711
17.244.6	Default	1711
17.245	fix wall/region command	1711
17.245.1	Syntax	1711
17.245.2	Examples	1712
17.245.3	Description	1712
17.245.4	Restart, fix_modify, output, run start/stop, minimize info	1713
17.245.5	Restrictions	1714
17.245.6	Related commands	1714
17.245.7	Default	1714
17.246	fix wall/srd command	1714
17.246.1	Syntax	1714
17.246.2	Examples	1715
17.246.3	Description	1715
17.246.4	Restart, fix_modify, output, run start/stop, minimize info	1717
17.246.5	Restrictions	1717
17.246.6	Related commands	1717
17.246.7	Default	1717
17.247	fix widom command	1717
17.247.1	Syntax	1717
17.247.2	Examples	1718
17.247.3	Description	1718
17.247.4	Restart, fix_modify, output, run start/stop, minimize info	1719
17.247.5	Restrictions	1720
17.247.6	Related commands	1720
17.247.7	Default	1720
<b>18</b>	<b>Computes</b>	<b>1721</b>
18.1	compute ackland/atom command	1721
18.1.1	Syntax	1721
18.1.2	Examples	1721

18.1.3	Description	1721
18.1.4	Output info	1722
18.1.5	Restrictions	1722
18.1.6	Related commands	1722
18.1.7	Default	1722
18.2	compute adf command	1722
18.2.1	Syntax	1722
18.2.2	Examples	1723
18.2.3	Description	1723
18.2.4	Output info	1725
18.2.5	Restrictions	1725
18.2.6	Related commands	1725
18.2.7	Default	1725
18.3	compute angle command	1725
18.3.1	Syntax	1725
18.3.2	Examples	1726
18.3.3	Description	1726
18.3.4	Output info	1726
18.3.5	Restrictions	1726
18.3.6	Related commands	1726
18.3.7	Default	1726
18.4	compute angle/local command	1726
18.4.1	Syntax	1726
18.4.2	Examples	1727
18.4.3	Description	1727
18.4.4	Output info	1728
18.4.5	Restrictions	1728
18.4.6	Related commands	1728
18.4.7	Default	1728
18.5	compute angmom/chunk command	1729
18.5.1	Syntax	1729
18.5.2	Examples	1729
18.5.3	Description	1729
18.5.4	Output info	1730
18.5.5	Restrictions	1730
18.5.6	Related commands	1730
18.5.7	Default	1730
18.6	compute ave/sphere/atom command	1730
18.6.1	Syntax	1730
18.6.2	Examples	1730
18.6.3	Description	1731
18.6.4	Output info	1731
18.6.5	Restrictions	1732
18.6.6	Related commands	1732
18.6.7	Default	1732
18.7	compute basal/atom command	1732
18.7.1	Syntax	1732
18.7.2	Examples	1732
18.7.3	Description	1732
18.7.4	Output info	1733
18.7.5	Restrictions	1733
18.7.6	Related commands	1733
18.7.7	Default	1733
18.8	compute body/local command	1733

18.8.1	Syntax . . . . .	1733
18.8.2	Examples . . . . .	1734
18.8.3	Description . . . . .	1734
18.8.4	Output info . . . . .	1734
18.8.5	Restrictions . . . . .	1735
18.8.6	Related commands . . . . .	1735
18.8.7	Default . . . . .	1735
18.9	compute bond command . . . . .	1735
18.9.1	Syntax . . . . .	1735
18.9.2	Examples . . . . .	1735
18.9.3	Description . . . . .	1735
18.9.4	Output info . . . . .	1735
18.9.5	Restrictions . . . . .	1736
18.9.6	Related commands . . . . .	1736
18.9.7	Default . . . . .	1736
18.10	compute bond/local command . . . . .	1736
18.10.1	Syntax . . . . .	1736
18.10.2	Examples . . . . .	1737
18.10.3	Description . . . . .	1737
18.10.4	Output info . . . . .	1738
18.10.5	Restrictions . . . . .	1739
18.10.6	Related commands . . . . .	1739
18.10.7	Default . . . . .	1739
18.11	compute born/matrix command . . . . .	1739
18.11.1	Syntax . . . . .	1739
18.11.2	Examples . . . . .	1739
18.11.3	Description . . . . .	1740
18.11.4	Restrictions . . . . .	1741
18.11.5	Default . . . . .	1742
18.12	compute centro/atom command . . . . .	1742
18.12.1	Syntax . . . . .	1742
18.12.2	Examples . . . . .	1742
18.12.3	Description . . . . .	1742
18.12.4	Output info . . . . .	1743
18.12.5	Restrictions . . . . .	1744
18.12.6	Related commands . . . . .	1744
18.12.7	Default . . . . .	1744
18.13	compute chunk/atom command . . . . .	1744
18.13.1	Syntax . . . . .	1744
18.13.2	Examples . . . . .	1746
18.13.3	Description . . . . .	1746
18.13.4	Output info . . . . .	1752
18.13.5	Restrictions . . . . .	1752
18.13.6	Related commands . . . . .	1752
18.13.7	Default . . . . .	1752
18.14	compute chunk/spread/atom command . . . . .	1753
18.14.1	Syntax . . . . .	1753
18.14.2	Examples . . . . .	1753
18.14.3	Description . . . . .	1753
18.14.4	Output info . . . . .	1755
18.14.5	Restrictions . . . . .	1755
18.14.6	Related commands . . . . .	1756
18.14.7	Default . . . . .	1756
18.15	compute cluster/atom command . . . . .	1756

18.16	compute fragment/atom command . . . . .	1756
18.17	compute aggregate/atom command . . . . .	1756
	18.17.1 Syntax . . . . .	1756
	18.17.2 Examples . . . . .	1756
	18.17.3 Description . . . . .	1756
	18.17.4 Output info . . . . .	1757
	18.17.5 Restrictions . . . . .	1757
	18.17.6 Related commands . . . . .	1757
	18.17.7 Default . . . . .	1758
18.18	compute cna/atom command . . . . .	1758
	18.18.1 Syntax . . . . .	1758
	18.18.2 Examples . . . . .	1758
	18.18.3 Description . . . . .	1758
	18.18.4 Output info . . . . .	1759
	18.18.5 Restrictions . . . . .	1759
	18.18.6 Related commands . . . . .	1759
	18.18.7 Default . . . . .	1759
18.19	compute cnp/atom command . . . . .	1759
	18.19.1 Syntax . . . . .	1759
	18.19.2 Examples . . . . .	1760
	18.19.3 Description . . . . .	1760
	18.19.4 Output info . . . . .	1761
	18.19.5 Restrictions . . . . .	1761
	18.19.6 Related commands . . . . .	1761
	18.19.7 Default . . . . .	1761
18.20	compute com command . . . . .	1761
	18.20.1 Syntax . . . . .	1761
	18.20.2 Examples . . . . .	1762
	18.20.3 Description . . . . .	1762
	18.20.4 Output info . . . . .	1762
	18.20.5 Restrictions . . . . .	1762
	18.20.6 Related commands . . . . .	1762
	18.20.7 Default . . . . .	1762
18.21	compute com/chunk command . . . . .	1762
	18.21.1 Syntax . . . . .	1762
	18.21.2 Examples . . . . .	1763
	18.21.3 Description . . . . .	1763
	18.21.4 Output info . . . . .	1763
	18.21.5 Restrictions . . . . .	1764
	18.21.6 Related commands . . . . .	1764
	18.21.7 Default . . . . .	1764
18.22	compute contact/atom command . . . . .	1764
	18.22.1 Syntax . . . . .	1764
	18.22.2 Examples . . . . .	1764
	18.22.3 Description . . . . .	1764
	18.22.4 Output info . . . . .	1764
	18.22.5 Restrictions . . . . .	1765
	18.22.6 Related commands . . . . .	1765
	18.22.7 Default . . . . .	1765
18.23	compute coord/atom command . . . . .	1765
	18.23.1 Syntax . . . . .	1765
	18.23.2 Examples . . . . .	1765
	18.23.3 Description . . . . .	1766
	18.23.4 Output info . . . . .	1767

18.23.5	Restrictions	1767
18.23.6	Related commands	1767
18.23.7	Default	1767
18.24	compute damage/atom command	1767
18.24.1	Syntax	1767
18.24.2	Examples	1768
18.24.3	Description	1768
18.24.4	Output info	1768
18.24.5	Restrictions	1768
18.24.6	Related commands	1768
18.24.7	Default	1768
18.25	compute dihedral command	1768
18.25.1	Syntax	1768
18.25.2	Examples	1769
18.25.3	Description	1769
18.25.4	Output info	1769
18.25.5	Restrictions	1769
18.25.6	Related commands	1769
18.25.7	Default	1769
18.26	compute dihedral/local command	1769
18.26.1	Syntax	1769
18.26.2	Examples	1770
18.26.3	Description	1770
18.26.4	Output info	1771
18.26.5	Restrictions	1771
18.26.6	Related commands	1771
18.26.7	Default	1771
18.27	compute dilatation/atom command	1771
18.27.1	Syntax	1771
18.27.2	Examples	1772
18.27.3	Description	1772
18.27.4	Output info	1772
18.27.5	Restrictions	1772
18.27.6	Related commands	1772
18.27.7	Default	1772
18.28	compute dipole command	1772
18.28.1	Syntax	1772
18.28.2	Examples	1773
18.28.3	Description	1773
18.28.4	Output info	1773
18.28.5	Restrictions	1773
18.28.6	Related commands	1773
18.28.7	Default	1773
18.29	compute dipole/chunk command	1774
18.29.1	Syntax	1774
18.29.2	Examples	1774
18.29.3	Description	1774
18.29.4	Output info	1775
18.29.5	Restrictions	1775
18.29.6	Related commands	1775
18.29.7	Default	1775
18.30	compute displace/atom command	1775
18.30.1	Syntax	1775
18.30.2	Examples	1775



18.30.3	Description	1776
18.30.4	Output info	1777
18.30.5	Restrictions	1777
18.30.6	Related commands	1777
18.30.7	Default	1777
18.31	compute dpd command	1777
18.31.1	Syntax	1777
18.31.2	Examples	1778
18.31.3	Description	1778
18.31.4	Output info	1778
18.31.5	Restrictions	1778
18.31.6	Related commands	1779
18.31.7	Default	1779
18.32	compute dpd/atom command	1779
18.32.1	Syntax	1779
18.32.2	Examples	1779
18.32.3	Description	1779
18.32.4	Output info	1779
18.32.5	Restrictions	1780
18.32.6	Related commands	1780
18.32.7	Default	1780
18.33	compute edpd/temp/atom command	1780
18.33.1	Syntax	1780
18.33.2	Examples	1780
18.33.3	Description	1780
18.33.4	Output info	1781
18.33.5	Restrictions	1781
18.33.6	Related commands	1781
18.33.7	Default	1781
18.34	compute efield/atom command	1781
18.34.1	Syntax	1781
18.34.2	Examples	1781
18.34.3	Description	1782
18.34.4	Output info	1782
18.34.5	Restrictions	1782
18.34.6	Related commands	1782
18.34.7	Default	1782
18.35	compute entropy/atom command	1782
18.35.1	Syntax	1782
18.35.2	Examples	1783
18.35.3	Description	1783
18.35.4	Output info	1784
18.35.5	Restrictions	1784
18.35.6	Related commands	1784
18.35.7	Default	1784
18.36	compute erotate/asphere command	1784
18.36.1	Syntax	1784
18.36.2	Examples	1785
18.36.3	Description	1785
18.36.4	Output info	1785
18.36.5	Restrictions	1785
18.36.6	Related commands	1785
18.36.7	Default	1786
18.37	compute erotate/rigid command	1786

18.37.1	Syntax . . . . .	1786
18.37.2	Examples . . . . .	1786
18.37.3	Description . . . . .	1786
18.37.4	Output info . . . . .	1786
18.37.5	Restrictions . . . . .	1786
18.37.6	Related commands . . . . .	1787
18.37.7	Default . . . . .	1787
18.38	compute erotate/sphere command . . . . .	1787
18.38.1	Syntax . . . . .	1787
18.38.2	Examples . . . . .	1787
18.38.3	Description . . . . .	1787
18.38.4	Output info . . . . .	1787
18.38.5	Restrictions . . . . .	1788
18.38.6	Related commands . . . . .	1788
18.38.7	Default . . . . .	1788
18.39	compute erotate/sphere/atom command . . . . .	1788
18.39.1	Syntax . . . . .	1788
18.39.2	Examples . . . . .	1788
18.39.3	Description . . . . .	1788
18.39.4	Output info . . . . .	1789
18.39.5	Restrictions . . . . .	1789
18.39.6	Related commands . . . . .	1789
18.39.7	Default . . . . .	1789
18.40	compute event/displace command . . . . .	1789
18.40.1	Syntax . . . . .	1789
18.40.2	Examples . . . . .	1789
18.40.3	Description . . . . .	1789
18.40.4	Output info . . . . .	1790
18.40.5	Restrictions . . . . .	1790
18.40.6	Related commands . . . . .	1790
18.40.7	Default . . . . .	1790
18.41	compute fabric command . . . . .	1790
18.41.1	Syntax . . . . .	1790
18.41.2	Examples . . . . .	1791
18.41.3	Description . . . . .	1791
18.41.4	Output info . . . . .	1792
18.41.5	Restrictions . . . . .	1792
18.41.6	Related commands . . . . .	1792
18.41.7	Default . . . . .	1792
18.42	compute fep command . . . . .	1793
18.42.1	Syntax . . . . .	1793
18.42.2	Examples . . . . .	1793
18.42.3	Description . . . . .	1793
18.42.4	Output info . . . . .	1796
18.42.5	Restrictions . . . . .	1796
18.42.6	Related commands . . . . .	1796
18.42.7	Default . . . . .	1796
18.43	compute fep/ta command . . . . .	1797
18.43.1	Syntax . . . . .	1797
18.43.2	Examples . . . . .	1797
18.43.3	Description . . . . .	1797
18.43.4	Output info . . . . .	1797
18.43.5	Restrictions . . . . .	1798
18.43.6	Related commands . . . . .	1798

18.43.7	Default	1798
18.44	compute global/atom command	1798
18.44.1	Syntax	1798
18.44.2	Examples	1799
18.44.3	Description	1799
18.44.4	Output info	1801
18.44.5	Restrictions	1801
18.44.6	Related commands	1801
18.44.7	Default	1801
18.45	compute group/group command	1801
18.45.1	Syntax	1801
18.45.2	Examples	1802
18.45.3	Description	1802
18.45.4	Output info	1803
18.45.5	Restrictions	1803
18.45.6	Related commands	1803
18.45.7	Default	1803
18.46	compute gyration command	1804
18.46.1	Syntax	1804
18.46.2	Examples	1804
18.46.3	Description	1804
18.46.4	Output info	1804
18.46.5	Restrictions	1805
18.46.6	Related commands	1805
18.46.7	Default	1805
18.47	compute gyration/chunk command	1805
18.47.1	Syntax	1805
18.47.2	Examples	1805
18.47.3	Description	1805
18.47.4	Output info	1806
18.47.5	Restrictions	1806
18.47.6	Related commands	1806
18.47.7	Default	1807
18.48	compute gyration/shape command	1807
18.48.1	Syntax	1807
18.48.2	Examples	1807
18.48.3	Description	1807
18.48.4	Output info	1808
18.48.5	Restrictions	1808
18.48.6	Related commands	1808
18.48.7	Default	1808
18.49	compute gyration/shape/chunk command	1808
18.49.1	Syntax	1808
18.49.2	Examples	1808
18.49.3	Description	1809
18.49.4	Output info	1809
18.49.5	Restrictions	1809
18.49.6	Related commands	1810
18.49.7	Default	1810
18.50	compute heat/flux command	1810
18.50.1	Syntax	1810
18.50.2	Examples	1810
18.50.3	Description	1810
18.50.4	Output info	1812

18.50.5	Restrictions	1812
18.50.6	Related commands	1812
18.50.7	Default	1812
18.51	compute hexorder/atom command	1814
18.51.1	Syntax	1814
18.51.2	Examples	1814
18.51.3	Description	1814
18.51.4	Output info	1815
18.51.5	Restrictions	1815
18.51.6	Related commands	1815
18.51.7	Default	1815
18.52	compute hma command	1815
18.52.1	Syntax	1815
18.52.2	Examples	1816
18.52.3	Description	1816
18.52.4	Output info	1818
18.52.5	Restrictions	1818
18.52.6	Related commands	1818
18.52.7	Default	1818
18.53	compute improper command	1818
18.53.1	Syntax	1818
18.53.2	Examples	1818
18.53.3	Description	1819
18.53.4	Output info	1819
18.53.5	Restrictions	1819
18.53.6	Related commands	1819
18.53.7	Default	1819
18.54	compute improper/local command	1819
18.54.1	Syntax	1819
18.54.2	Examples	1820
18.54.3	Description	1820
18.54.4	Output info	1820
18.54.5	Restrictions	1820
18.54.6	Related commands	1820
18.54.7	Default	1821
18.55	compute inertia/chunk command	1821
18.55.1	Syntax	1821
18.55.2	Examples	1821
18.55.3	Description	1821
18.55.4	Output info	1822
18.55.5	Restrictions	1822
18.55.6	Related commands	1822
18.55.7	Default	1822
18.56	compute ke command	1822
18.56.1	Syntax	1822
18.56.2	Examples	1822
18.56.3	Description	1822
18.56.4	Output info	1823
18.56.5	Restrictions	1823
18.56.6	Related commands	1823
18.56.7	Default	1823
18.57	compute ke/atom command	1823
18.57.1	Syntax	1823
18.57.2	Examples	1823

18.57.3	Description	1823
18.57.4	Output info	1824
18.57.5	Restrictions	1824
18.57.6	Related commands	1824
18.57.7	Default	1824
18.58	compute ke/atom/eff command	1824
18.58.1	Syntax	1824
18.58.2	Examples	1824
18.58.3	Description	1824
18.58.4	Output info	1825
18.58.5	Restrictions	1825
18.58.6	Related commands	1825
18.58.7	Default	1825
18.59	compute ke/eff command	1825
18.59.1	Syntax	1825
18.59.2	Examples	1826
18.59.3	Description	1826
18.59.4	Output info	1826
18.59.5	Restrictions	1826
18.59.6	Related commands	1827
18.59.7	Default	1827
18.60	compute ke/rigid command	1827
18.60.1	Syntax	1827
18.60.2	Examples	1827
18.60.3	Description	1827
18.60.4	Output info	1827
18.60.5	Restrictions	1828
18.60.6	Related commands	1828
18.60.7	Default	1828
18.61	compute mesont command	1828
18.61.1	Syntax	1828
18.61.2	Examples	1828
18.61.3	Description	1828
18.61.4	Output info	1828
18.61.5	Restrictions	1829
18.61.6	Related commands	1829
18.61.7	Default	1829
18.62	compute mliap command	1829
18.62.1	Syntax	1829
18.62.2	Examples	1829
18.62.3	Description	1829
18.62.4	Output info	1831
18.62.5	Restrictions	1831
18.62.6	Related commands	1831
18.62.7	Default	1831
18.63	compute momentum command	1831
18.63.1	Syntax	1831
18.63.2	Examples	1832
18.63.3	Description	1832
18.63.4	Output info	1832
18.63.5	Restrictions	1832
18.63.6	Related commands	1832
18.63.7	Default	1832
18.64	compute msd command	1832

18.64.1	Syntax	1832
18.64.2	Examples	1833
18.64.3	Description	1833
18.64.4	Output info	1834
18.64.5	Restrictions	1834
18.64.6	Related commands	1834
18.64.7	Default	1834
18.65	compute msd/chunk command	1834
18.65.1	Syntax	1834
18.65.2	Examples	1834
18.65.3	Description	1834
18.65.4	Output info	1835
18.65.5	Restrictions	1836
18.65.6	Related commands	1836
18.65.7	Default	1836
18.66	compute msd/nongauss command	1836
18.66.1	Syntax	1836
18.66.2	Examples	1836
18.66.3	Description	1836
18.66.4	Output info	1837
18.66.5	Restrictions	1837
18.66.6	Related commands	1837
18.66.7	Default	1837
18.67	compute nbond/atom command	1837
18.67.1	Syntax	1837
18.67.2	Examples	1837
18.67.3	Description	1838
18.67.4	Output info	1838
18.67.5	Restrictions	1838
18.67.6	Related commands	1838
18.67.7	Default	1838
18.68	compute omega/chunk command	1838
18.68.1	Syntax	1838
18.68.2	Examples	1838
18.68.3	Description	1838
18.68.4	Output info	1839
18.68.5	Restrictions	1839
18.68.6	Related commands	1839
18.68.7	Default	1839
18.69	compute orientorder/atom command	1840
18.69.1	Syntax	1840
18.69.2	Examples	1840
18.69.3	Description	1840
18.69.4	Output info	1842
18.69.5	Restrictions	1842
18.69.6	Related commands	1842
18.69.7	Default	1842
18.70	compute pair command	1843
18.70.1	Syntax	1843
18.70.2	Examples	1843
18.70.3	Description	1843
18.70.4	Output info	1844
18.70.5	Restrictions	1844
18.70.6	Related commands	1844

18.70.7	Default	1844
18.71	compute pair/local command	1844
18.71.1	Syntax	1844
18.71.2	Examples	1845
18.71.3	Description	1845
18.71.4	Output info	1846
18.71.5	Restrictions	1846
18.71.6	Related commands	1846
18.71.7	Default	1846
18.72	compute pe command	1846
18.72.1	Syntax	1846
18.72.2	Examples	1847
18.72.3	Description	1847
18.72.4	Output info	1847
18.72.5	Restrictions	1848
18.72.6	Related commands	1848
18.72.7	Default	1848
18.73	compute pe/atom command	1848
18.73.1	Syntax	1848
18.73.2	Examples	1848
18.73.3	Description	1848
18.73.4	Output info	1849
18.73.5	Restrictions	1849
18.73.6	Related commands	1849
18.73.7	Default	1849
18.74	compute plasticity/atom command	1850
18.74.1	Syntax	1850
18.74.2	Examples	1850
18.74.3	Description	1850
18.74.4	Output info	1850
18.74.5	Restrictions	1850
18.74.6	Related commands	1850
18.74.7	Default	1851
18.75	compute pressure command	1851
18.75.1	Syntax	1851
18.75.2	Examples	1851
18.75.3	Description	1851
18.75.4	Output info	1853
18.75.5	Restrictions	1853
18.75.6	Related commands	1853
18.75.7	Default	1853
18.76	compute pressure/uef command	1853
18.76.1	Syntax	1853
18.76.2	Examples	1853
18.76.3	Description	1854
18.76.4	Restrictions	1854
18.76.5	Related commands	1854
18.76.6	Default	1854
18.77	compute property/atom command	1854
18.77.1	Syntax	1854
18.77.2	Examples	1856
18.77.3	Description	1856
18.77.4	Output info	1857
18.77.5	Restrictions	1857

18.77.6	Related commands	1857
18.77.7	Default	1857
18.78	compute property/chunk command	1857
18.78.1	Syntax	1857
18.78.2	Examples	1858
18.78.3	Description	1858
18.78.4	Output info	1859
18.78.5	Restrictions	1859
18.78.6	Related commands	1859
18.78.7	Default	1859
18.79	compute property/local command	1859
18.79.1	Syntax	1859
18.79.2	Examples	1860
18.79.3	Description	1860
18.79.4	Output info	1861
18.79.5	Restrictions	1861
18.79.6	Related commands	1862
18.79.7	Default	1862
18.80	compute ptm/atom command	1862
18.80.1	Syntax	1862
18.80.2	Examples	1862
18.80.3	Description	1862
18.80.4	Output info	1863
18.80.5	Restrictions	1864
18.80.6	Related commands	1864
18.80.7	Default	1864
18.81	compute rdf command	1864
18.81.1	Syntax	1864
18.81.2	Examples	1864
18.81.3	Description	1865
18.81.4	Output info	1866
18.81.5	Restrictions	1866
18.81.6	Related commands	1867
18.81.7	Default	1867
18.82	compute reduce command	1867
18.83	compute reduce/region command	1867
18.83.1	Syntax	1867
18.83.2	Examples	1868
18.83.3	Description	1868
18.83.4	Output info	1870
18.83.5	Restrictions	1870
18.83.6	Related commands	1870
18.83.7	Default	1870
18.84	compute reduce/chunk command	1870
18.84.1	Syntax	1870
18.84.2	Examples	1871
18.84.3	Description	1871
18.84.4	Output info	1872
18.84.5	Restrictions	1873
18.84.6	Related commands	1873
18.84.7	Default	1873
18.85	compute rigid/local command	1873
18.85.1	Syntax	1873
18.85.2	Examples	1874



18.85.3	Description	1874
18.85.4	Output info	1875
18.85.5	Restrictions	1875
18.85.6	Related commands	1876
18.85.7	Default	1876
18.86	compute saed command	1876
18.86.1	Syntax	1876
18.86.2	Examples	1876
18.86.3	Description	1877
18.86.4	Output info	1879
18.86.5	Restrictions	1879
18.86.6	Related commands	1879
18.86.7	Default	1879
18.87	compute slice command	1879
18.87.1	Syntax	1879
18.87.2	Examples	1880
18.87.3	Description	1880
18.87.4	Output info	1881
18.87.5	Restrictions	1881
18.87.6	Related commands	1881
18.87.7	Default	1881
18.88	compute smd/contact/radius command	1881
18.88.1	Syntax	1881
18.88.2	Examples	1881
18.88.3	Description	1882
18.88.4	Output info	1882
18.88.5	Restrictions	1882
18.88.6	Related commands	1882
18.88.7	Default	1882
18.89	compute smd/damage command	1882
18.89.1	Syntax	1882
18.89.2	Examples	1883
18.89.3	Description	1883
18.89.4	Restrictions	1883
18.89.5	Related commands	1883
18.89.6	Default	1883
18.90	compute smd/hourglass/error command	1883
18.90.1	Syntax	1883
18.90.2	Examples	1884
18.90.3	Description	1884
18.90.4	Restrictions	1884
18.90.5	Related commands	1884
18.90.6	Default	1884
18.91	compute smd/internal/energy command	1884
18.91.1	Syntax	1884
18.91.2	Examples	1885
18.91.3	Description	1885
18.91.4	Output Info	1885
18.91.5	Restrictions	1885
18.91.6	Related commands	1885
18.91.7	Default	1885
18.92	compute smd/plastic/strain command	1885
18.92.1	Syntax	1885
18.92.2	Examples	1886

18.92.3	Description	1886
18.92.4	Restrictions	1886
18.92.5	Related commands	1886
18.92.6	Default	1886
18.93	compute smd/plastic/strain/rate command	1886
18.93.1	Syntax	1886
18.93.2	Examples	1887
18.93.3	Description	1887
18.93.4	Restrictions	1887
18.93.5	Related commands	1887
18.93.6	Default	1887
18.94	compute smd/rho command	1887
18.94.1	Syntax	1887
18.94.2	Examples	1888
18.94.3	Description	1888
18.94.4	Output info	1888
18.94.5	Restrictions	1888
18.94.6	Related commands	1888
18.94.7	Default	1888
18.95	compute smd/tlsph/defgrad command	1888
18.95.1	Syntax	1888
18.95.2	Examples	1889
18.95.3	Description	1889
18.95.4	Output info	1889
18.95.5	Restrictions	1889
18.95.6	Related commands	1889
18.95.7	Default	1889
18.96	compute smd/tlsph/dt command	1889
18.96.1	Syntax	1889
18.96.2	Examples	1890
18.96.3	Description	1890
18.96.4	Output info	1890
18.96.5	Restrictions	1890
18.96.6	Related commands	1890
18.96.7	Default	1890
18.97	compute smd/tlsph/num/neighs command	1890
18.97.1	Syntax	1890
18.97.2	Examples	1891
18.97.3	Description	1891
18.97.4	Output info	1891
18.97.5	Restrictions	1891
18.97.6	Related commands	1891
18.97.7	Default	1891
18.98	compute smd/tlsph/shape command	1891
18.98.1	Syntax	1891
18.98.2	Examples	1892
18.98.3	Description	1892
18.98.4	Output info	1892
18.98.5	Restrictions	1892
18.98.6	Related commands	1892
18.98.7	Default	1892
18.99	compute smd/tlsph/strain command	1892
18.99.1	Syntax	1892
18.99.2	Examples	1893

18.99.3	Description	1893
18.99.4	Output info	1893
18.99.5	Restrictions	1893
18.99.6	Related commands	1893
18.99.7	Default	1893
18.100	compute smd/tlsph/strain/rate command	1893
18.100.1	Syntax	1893
18.100.2	Examples	1894
18.100.3	Description	1894
18.100.4	Output info	1894
18.100.5	Restrictions	1894
18.100.6	Related commands	1894
18.100.7	Default	1894
18.101	compute smd/tlsph/stress command	1894
18.101.1	Syntax	1894
18.101.2	Examples	1895
18.101.3	Description	1895
18.101.4	Output info	1895
18.101.5	Restrictions	1895
18.101.6	Related commands	1895
18.101.7	Default	1895
18.102	compute smd/triangle/vertices command	1895
18.102.1	Syntax	1895
18.102.2	Examples	1896
18.102.3	Description	1896
18.102.4	Output info	1896
18.102.5	Restrictions	1896
18.102.6	Related commands	1896
18.102.7	Default	1896
18.103	compute smd/ulsph/effm command	1896
18.103.1	Syntax	1896
18.103.2	Examples	1897
18.103.3	Description	1897
18.103.4	Output info	1897
18.103.5	Restrictions	1897
18.103.6	Related commands	1897
18.103.7	Default	1897
18.104	compute smd/ulsph/num/neighs command	1897
18.104.1	Syntax	1897
18.104.2	Examples	1898
18.104.3	Description	1898
18.104.4	Output info	1898
18.104.5	Restrictions	1898
18.104.6	Related commands	1898
18.104.7	Default	1898
18.105	compute smd/ulsph/strain command	1898
18.105.1	Syntax	1898
18.105.2	Examples	1899
18.105.3	Description	1899
18.105.4	Output info	1899
18.105.5	Restrictions	1899
18.105.6	Related commands	1899
18.105.7	Default	1899
18.106	compute smd/ulsph/strain/rate command	1899

18.106.1	Syntax . . . . .	1899
18.106.2	Examples . . . . .	1900
18.106.3	Description . . . . .	1900
18.106.4	Output info . . . . .	1900
18.106.5	Restrictions . . . . .	1900
18.106.6	Related commands . . . . .	1900
18.106.7	Default . . . . .	1900
18.107	compute smd/ulsph/stress command . . . . .	1900
18.107.1	Syntax . . . . .	1900
18.107.2	Examples . . . . .	1901
18.107.3	Description . . . . .	1901
18.107.4	Output info . . . . .	1901
18.107.5	Restrictions . . . . .	1901
18.107.6	Related commands . . . . .	1901
18.107.7	Default . . . . .	1901
18.108	compute smd/vol command . . . . .	1901
18.108.1	Syntax . . . . .	1901
18.108.2	Examples . . . . .	1902
18.108.3	Description . . . . .	1902
18.108.4	Output info . . . . .	1902
18.108.5	Restrictions . . . . .	1902
18.108.6	Related commands . . . . .	1902
18.108.7	Default . . . . .	1902
18.109	compute sna/atom command . . . . .	1903
18.110	compute snad/atom command . . . . .	1903
18.111	compute snav/atom command . . . . .	1903
18.112	compute snap command . . . . .	1903
18.113	compute sna/grid command . . . . .	1903
18.114	compute sna/grid/local command . . . . .	1903
18.114.1	Syntax . . . . .	1903
18.114.2	Examples . . . . .	1904
18.114.3	Description . . . . .	1904
18.114.4	Output info . . . . .	1908
18.114.5	Restrictions . . . . .	1910
18.114.6	Related commands . . . . .	1910
18.114.7	Default . . . . .	1910
18.115	compute sph/e/atom command . . . . .	1910
18.115.1	Syntax . . . . .	1910
18.115.2	Examples . . . . .	1910
18.115.3	Description . . . . .	1911
18.115.4	Output info . . . . .	1911
18.115.5	Restrictions . . . . .	1911
18.115.6	Related commands . . . . .	1911
18.115.7	Default . . . . .	1911
18.116	compute sph/rho/atom command . . . . .	1911
18.116.1	Syntax . . . . .	1911
18.116.2	Examples . . . . .	1912
18.116.3	Description . . . . .	1912
18.116.4	Output info . . . . .	1912
18.116.5	Restrictions . . . . .	1912
18.116.6	Related commands . . . . .	1912
18.116.7	Default . . . . .	1912
18.117	compute sph/t/atom command . . . . .	1912
18.117.1	Syntax . . . . .	1912

18.117.2	Examples	1913
18.117.3	Description	1913
18.117.4	Output info	1913
18.117.5	Restrictions	1913
18.117.6	Related commands	1913
18.117.7	Default	1913
18.118	compute spin command	1913
18.118.1	Syntax	1913
18.118.2	Examples	1914
18.118.3	Description	1914
18.118.4	Output info	1914
18.118.5	Restrictions	1914
18.118.6	Default	1915
18.119	compute stress/atom command	1915
18.120	compute centroid/stress/atom command	1915
18.120.1	Syntax	1915
18.120.2	Examples	1915
18.120.3	Description	1915
18.120.4	Output info	1918
18.120.5	Restrictions	1918
18.120.6	Related commands	1918
18.120.7	Default	1918
18.121	compute stress/mop command	1919
18.122	compute stress/mop/profile command	1919
18.122.1	Syntax	1919
18.122.2	Examples	1919
18.122.3	Description	1919
18.122.4	Output info	1920
18.122.5	Restrictions	1920
18.122.6	Related commands	1920
18.122.7	Default	1921
18.123	compute stress/cartesian command	1921
18.124	compute stress/cylinder command	1921
18.125	compute stress/spherical command	1921
18.125.1	Syntax	1921
18.125.2	Examples	1921
18.125.3	Description	1922
18.125.4	Output info	1922
18.125.5	Restrictions	1923
18.125.6	Related commands	1923
18.125.7	Default	1923
18.126	compute force/tally command	1923
18.127	compute heat/flux/tally command	1923
18.128	compute heat/flux/virial/tally command	1923
18.129	compute pe/tally command	1923
18.130	compute pe/mol/tally command	1923
18.131	compute stress/tally command	1923
18.131.1	Syntax	1923
18.131.2	Examples	1924
18.131.3	Description	1924
18.131.4	Output info	1925
18.131.5	Restrictions	1926
18.131.6	Related commands	1926
18.131.7	Default	1926

18.132	compute tdpd/cc/atom command . . . . .	1926
18.132.1	Syntax . . . . .	1926
18.132.2	Examples . . . . .	1926
18.132.3	Description . . . . .	1926
18.132.4	Output info . . . . .	1927
18.132.5	Restrictions . . . . .	1927
18.132.6	Related commands . . . . .	1927
18.132.7	Default . . . . .	1927
18.133	compute temp command . . . . .	1927
18.133.1	Syntax . . . . .	1927
18.133.2	Examples . . . . .	1927
18.133.3	Description . . . . .	1928
18.133.4	Output info . . . . .	1928
18.133.5	Restrictions . . . . .	1929
18.133.6	Related commands . . . . .	1929
18.133.7	Default . . . . .	1929
18.134	compute temp/asphere command . . . . .	1929
18.134.1	Syntax . . . . .	1929
18.134.2	Examples . . . . .	1929
18.134.3	Description . . . . .	1929
18.134.4	Output info . . . . .	1931
18.134.5	Restrictions . . . . .	1931
18.134.6	Related commands . . . . .	1931
18.134.7	Default . . . . .	1931
18.135	compute temp/body command . . . . .	1931
18.135.1	Syntax . . . . .	1931
18.135.2	Examples . . . . .	1932
18.135.3	Description . . . . .	1932
18.135.4	Output info . . . . .	1933
18.135.5	Restrictions . . . . .	1933
18.135.6	Related commands . . . . .	1933
18.135.7	Default . . . . .	1933
18.136	compute temp/chunk command . . . . .	1933
18.136.1	Syntax . . . . .	1933
18.136.2	Examples . . . . .	1934
18.136.3	Description . . . . .	1934
18.136.4	Output info . . . . .	1936
18.136.5	Restrictions . . . . .	1936
18.136.6	Related commands . . . . .	1936
18.136.7	Default . . . . .	1936
18.137	compute temp/com command . . . . .	1936
18.137.1	Syntax . . . . .	1936
18.137.2	Examples . . . . .	1937
18.137.3	Description . . . . .	1937
18.137.4	Output info . . . . .	1937
18.137.5	Restrictions . . . . .	1938
18.137.6	Related commands . . . . .	1938
18.137.7	Default . . . . .	1938
18.138	compute temp/cs command . . . . .	1938
18.138.1	Syntax . . . . .	1938
18.138.2	Examples . . . . .	1938
18.138.3	Description . . . . .	1938
18.138.4	Output info . . . . .	1939
18.138.5	Restrictions . . . . .	1939

18.138.6	Related commands	1939
18.138.7	Default	1940
18.139	compute temp/deform command	1940
18.139.1	Syntax	1940
18.139.2	Examples	1940
18.139.3	Description	1940
18.139.4	Output info	1941
18.139.5	Restrictions	1941
18.139.6	Related commands	1942
18.139.7	Default	1942
18.140	compute temp/deform/eff command	1942
18.140.1	Syntax	1942
18.140.2	Examples	1942
18.140.3	Description	1942
18.140.4	Output info	1942
18.140.5	Restrictions	1943
18.140.6	Related commands	1943
18.140.7	Default	1943
18.141	compute temp/drude command	1943
18.141.1	Syntax	1943
18.141.2	Examples	1943
18.141.3	Description	1943
18.141.4	Output info	1944
18.141.5	Restrictions	1944
18.141.6	Related commands	1944
18.141.7	Default	1944
18.142	compute temp/eff command	1944
18.142.1	Syntax	1944
18.142.2	Examples	1945
18.142.3	Description	1945
18.142.4	Output info	1946
18.142.5	Restrictions	1946
18.142.6	Related commands	1946
18.142.7	Default	1946
18.143	compute temp/partial command	1946
18.143.1	Syntax	1946
18.143.2	Examples	1946
18.143.3	Description	1946
18.143.4	Output info	1947
18.143.5	Restrictions	1947
18.143.6	Related commands	1948
18.143.7	Default	1948
18.144	compute temp/profile command	1948
18.144.1	Syntax	1948
18.144.2	Examples	1948
18.144.3	Description	1949
18.144.4	Output info	1950
18.144.5	Restrictions	1950
18.144.6	Related commands	1950
18.144.7	Default	1951
18.145	compute temp/ramp command	1951
18.145.1	Syntax	1951
18.145.2	Examples	1951
18.145.3	Description	1951

18.145.4	Output info	1952
18.145.5	Restrictions	1952
18.145.6	Related commands	1952
18.145.7	Default	1952
18.146	compute temp/region command	1953
18.146.1	Syntax	1953
18.146.2	Examples	1953
18.146.3	Description	1953
18.146.4	Output info	1954
18.146.5	Restrictions	1954
18.146.6	Related commands	1954
18.146.7	Default	1954
18.147	compute temp/region/eff command	1954
18.147.1	Syntax	1954
18.147.2	Examples	1954
18.147.3	Description	1955
18.147.4	Output info	1955
18.147.5	Restrictions	1955
18.147.6	Related commands	1955
18.147.7	Default	1955
18.148	compute temp/rotate command	1955
18.148.1	Syntax	1955
18.148.2	Examples	1956
18.148.3	Description	1956
18.148.4	Output info	1956
18.148.5	Restrictions	1957
18.148.6	Related commands	1957
18.148.7	Default	1957
18.149	compute temp/sphere command	1957
18.149.1	Syntax	1957
18.149.2	Examples	1957
18.149.3	Description	1957
18.149.4	Output info	1958
18.149.5	Restrictions	1959
18.149.6	Related commands	1959
18.149.7	Default	1959
18.150	compute temp/uef command	1959
18.150.1	Syntax	1959
18.150.2	Examples	1959
18.150.3	Description	1959
18.150.4	Restrictions	1959
18.150.5	Related commands	1960
18.150.6	Default	1960
18.151	compute ti command	1960
18.151.1	Syntax	1960
18.151.2	Examples	1960
18.151.3	Description	1961
18.151.4	Output info	1961
18.151.5	Restrictions	1962
18.151.6	Related commands	1962
18.151.7	Default	1962
18.152	compute torque/chunk command	1962
18.152.1	Syntax	1962
18.152.2	Examples	1962



18.152.3	Description	1962
18.152.4	Output info	1963
18.152.5	Restrictions	1963
18.152.6	Related commands	1963
18.152.7	Default	1963
18.153	compute vacf command	1963
18.153.1	Syntax	1963
18.153.2	Examples	1964
18.153.3	Description	1964
18.153.4	Output info	1964
18.153.5	Restrictions	1964
18.153.6	Related commands	1965
18.153.7	Default	1965
18.154	compute vcm/chunk command	1965
18.154.1	Syntax	1965
18.154.2	Examples	1965
18.154.3	Description	1965
18.154.4	Output info	1966
18.154.5	Restrictions	1966
18.154.6	Related commands	1966
18.154.7	Default	1966
18.155	compute viscosity/cos command	1966
18.155.1	Syntax	1966
18.155.2	Examples	1966
18.155.3	Description	1967
18.155.4	Output info	1968
18.155.5	Restrictions	1968
18.155.6	Related commands	1968
18.155.7	Default	1968
18.156	compute voronoi/atom command	1968
18.156.1	Syntax	1968
18.156.2	Examples	1969
18.156.3	Description	1969
18.156.4	Output info	1971
18.156.5	Restrictions	1971
18.156.6	Related commands	1971
18.156.7	Default	1971
18.157	compute xrd command	1971
18.157.1	Syntax	1971
18.157.2	Examples	1972
18.157.3	Description	1972
18.157.4	Output info	1974
18.157.5	Restrictions	1974
18.157.6	Related commands	1974
18.157.7	Default	1974
<b>19</b>	<b>Pair Styles</b>	<b>1975</b>
19.1	pair_style adp command	1975
19.1.1	Syntax	1975
19.1.2	Examples	1975
19.1.3	Description	1975
19.1.4	Mixing, shift, table, tail correction, restart, rRESPA info	1977
19.1.5	Restrictions	1977
19.1.6	Related commands	1977

19.1.7	Default	1977
19.2	pair_style agni command	1978
19.2.1	Syntax	1978
19.2.2	Examples	1978
19.2.3	Description	1978
19.2.4	Mixing, shift, table, tail correction, restart, rRESPA info	1979
19.2.5	Restrictions	1979
19.2.6	Related commands	1979
19.2.7	Default	1979
19.3	pair_style airebo command	1980
19.4	pair_style airebo/morse command	1980
19.5	pair_style rebo command	1980
19.5.1	Syntax	1980
19.5.2	Examples	1980
19.5.3	Description	1980
19.5.4	Mixing, shift, table, tail correction, restart, rRESPA info	1982
19.5.5	Restrictions	1983
19.5.6	Related commands	1983
19.5.7	Default	1983
19.6	pair_style amoeba command	1983
19.7	pair_style hippo command	1983
19.7.1	Syntax	1983
19.7.2	Examples	1983
19.7.3	Additional info	1984
19.7.4	Description	1984
19.7.5	Mixing, shift, table, tail correction, restart, rRESPA info	1985
19.7.6	Restrictions	1986
19.7.7	Related commands	1986
19.7.8	Default	1986
19.8	pair_style atm command	1987
19.8.1	Syntax	1987
19.8.2	Examples	1987
19.8.3	Description	1987
19.8.4	Mixing, shift, table, tail correction, restart, rRESPA info	1989
19.8.5	Restrictions	1989
19.8.6	Related commands	1989
19.8.7	Default	1989
19.9	pair_style awpmd/cut command	1989
19.9.1	Syntax	1989
19.9.2	Examples	1990
19.9.3	Description	1990
19.9.4	Mixing, shift, table, tail correction, restart, rRESPA info	1991
19.9.5	Restrictions	1991
19.9.6	Related commands	1991
19.9.7	Default	1991
19.10	pair_style beck command	1991
19.10.1	Syntax	1991
19.10.2	Examples	1991
19.10.3	Description	1992
19.10.4	Mixing, shift, table, tail correction, restart, rRESPA info	1992
19.10.5	Restrictions	1993
19.10.6	Related commands	1993
19.10.7	Default	1993
19.11	pair_style body/nparticle command	1993

19.11.1	Syntax	1993
19.11.2	Examples	1993
19.11.3	Description	1993
19.11.4	Mixing, shift, table, tail correction, restart, rRESPA info	1994
19.11.5	Restrictions	1994
19.11.6	Related commands	1995
19.11.7	Default	1995
19.12	pair_style body/rounded/polygon command	1995
19.12.1	Syntax	1995
19.12.2	Examples	1995
19.12.3	Description	1995
19.12.4	Mixing, shift, table, tail correction, restart, rRESPA info	1997
19.12.5	Restrictions	1997
19.12.6	Related commands	1997
19.12.7	Default	1997
19.13	pair_style body/rounded/polyhedron command	1998
19.13.1	Syntax	1998
19.13.2	Examples	1998
19.13.3	Description	1998
19.13.4	Mixing, shift, table, tail correction, restart, rRESPA info	2000
19.13.5	Restrictions	2000
19.13.6	Related commands	2000
19.13.7	Default	2000
19.14	pair_style bop command	2000
19.14.1	Syntax	2000
19.14.2	Examples	2001
19.14.3	Description	2001
19.14.4	Mixing, shift, table, tail correction, restart, rRESPA info	2005
19.14.5	Restrictions	2005
19.14.6	Related commands	2006
19.14.7	Default	2006
19.15	pair_style born command	2006
19.16	pair_style born/coul/long command	2006
19.17	pair_style born/coul/msm command	2006
19.18	pair_style born/coul/wolf command	2006
19.19	pair_style born/coul/dsf command	2006
19.19.1	Syntax	2006
19.19.2	Examples	2007
19.19.3	Description	2008
19.19.4	Mixing, shift, table, tail correction, restart, rRESPA info	2009
19.19.5	Restrictions	2009
19.19.6	Related commands	2009
19.19.7	Default	2009
19.20	pair_style bpm/spring command	2009
19.20.1	Syntax	2009
19.20.2	Examples	2010
19.20.3	Description	2010
19.20.4	Mixing, shift, table, tail correction, restart, rRESPA info	2010
19.20.5	Restrictions	2011
19.20.6	Related commands	2011
19.20.7	Default	2011
19.21	pair_style brownian command	2011
19.22	pair_style brownian/poly command	2011
19.22.1	Syntax	2011

19.22.2	Examples	2012
19.22.3	Description	2012
19.22.4	Mixing, shift, table, tail correction, restart, rRESPA info	2013
19.22.5	Restrictions	2013
19.22.6	Related commands	2013
19.22.7	Default	2013
19.23	pair_style buck command	2013
19.24	pair_style buck/coul/cut command	2013
19.25	pair_style buck/coul/long command	2014
19.26	pair_style buck/coul/msm command	2014
19.26.1	Syntax	2014
19.26.2	Examples	2014
19.26.3	Description	2015
19.26.4	Mixing, shift, table, tail correction, restart, rRESPA info	2016
19.26.5	Restrictions	2016
19.26.6	Related commands	2016
19.26.7	Default	2016
19.27	pair_style buck6d/coul/gauss/dsf command	2016
19.28	pair_style buck6d/coul/gauss/long command	2016
19.28.1	Syntax	2016
19.28.2	Examples	2017
19.28.3	Description	2017
19.28.4	Mixing, shift, table, tail correction, restart, rRESPA info	2018
19.28.5	Restrictions	2018
19.28.6	Related commands	2018
19.28.7	Default	2018
19.29	pair_style buck/long/coul/long command	2019
19.29.1	Syntax	2019
19.29.2	Examples	2019
19.29.3	Description	2019
19.29.4	Mixing, shift, table, tail correction, restart, rRESPA info	2020
19.29.5	Restrictions	2021
19.29.6	Related commands	2021
19.29.7	Default	2021
19.30	pair_style lj/charmm/coul/charmm command	2021
19.31	pair_style lj/charmm/coul/charmm/implicit command	2021
19.32	pair_style lj/charmm/coul/long command	2021
19.33	pair_style lj/charmm/coul/msm command	2021
19.34	pair_style lj/charmmfsw/coul/charmmfsh command	2021
19.35	pair_style lj/charmmfsw/coul/long command	2021
19.35.1	Syntax	2021
19.35.2	Examples	2022
19.35.3	Description	2023
19.35.4	Mixing, shift, table, tail correction, restart, rRESPA info	2025
19.35.5	Restrictions	2025
19.35.6	Related commands	2025
19.35.7	Default	2025
19.36	pair_style lj/class2 command	2026
19.37	pair_style lj/class2/coul/cut command	2026
19.38	pair_style lj/class2/coul/long command	2026
19.38.1	Syntax	2026
19.38.2	Examples	2026
19.38.3	Description	2027
19.38.4	Mixing, shift, table, tail correction, restart, rRESPA info	2028

19.38.5	Restrictions	2028
19.38.6	Related commands	2028
19.38.7	Default	2028
19.39	pair_style colloid command	2028
19.39.1	Syntax	2028
19.39.2	Examples	2029
19.39.3	Description	2029
19.39.4	Mixing, shift, table, tail correction, restart, rRESPA info	2031
19.39.5	Restrictions	2031
19.39.6	Related commands	2031
19.39.7	Default	2031
19.40	pair_style comb command	2031
19.41	pair_style comb3 command	2032
19.41.1	Syntax	2032
19.41.2	Examples	2032
19.41.3	Description	2032
19.41.4	Mixing, shift, table, tail correction, restart, rRESPA info	2034
19.41.5	Restrictions	2034
19.41.6	Related commands	2034
19.41.7	Default	2034
19.42	pair_style cosine/squared command	2034
19.42.1	Syntax	2034
19.42.2	Examples	2035
19.42.3	Description	2035
19.42.4	Mixing, shift, table, tail correction, restart, rRESPA info	2036
19.42.5	Restrictions	2036
19.42.6	Related commands	2036
19.42.7	Default	2036
19.43	pair_style coul/cut command	2037
19.44	pair_style coul/debye command	2037
19.45	pair_style coul/dsf command	2037
19.46	pair_style coul/exclude command	2037
19.47	pair_style coul/cut/global command	2037
19.48	pair_style coul/long command	2037
19.49	pair_style coul/msm command	2037
19.50	pair_style coul/streitz command	2037
19.51	pair_style coul/wolf command	2037
19.52	pair_style tip4p/cut command	2038
19.53	pair_style tip4p/long command	2038
19.53.1	Syntax	2038
19.53.2	Examples	2038
19.53.3	Description	2039
19.53.4	Mixing, shift, table, tail correction, restart, rRESPA info	2041
19.53.5	Restrictions	2042
19.53.6	Related commands	2042
19.53.7	Default	2042
19.54	pair_style coul/diel command	2042
19.54.1	Syntax	2042
19.54.2	Examples	2042
19.54.3	Description	2043
19.54.4	Mixing, shift, table, tail correction, restart, rRESPA info	2043
19.54.5	Restrictions	2044
19.54.6	Related commands	2044
19.54.7	Default	2044

19.55	pair_style coul/shield command	2044
19.55.1	Syntax	2044
19.55.2	Examples	2044
19.55.3	Description	2044
19.55.4	Mixing, shift, table, tail correction, restart, rRESPA info	2045
19.55.5	Restrictions	2045
19.55.6	Related commands	2045
19.55.7	Default	2045
19.56	pair_style coul/slatter command	2046
19.57	pair_style coul/slatter/cut command	2046
19.58	pair_style coul/slatter/long command	2046
19.58.1	Syntax	2046
19.58.2	Examples	2046
19.58.3	Description	2046
19.58.4	Mixing, shift, table, tail correction, restart, rRESPA info	2047
19.58.5	Restrictions	2047
19.58.6	Related commands	2047
19.58.7	Default	2047
19.59	pair_style coul/tt command	2047
19.59.1	Syntax	2047
19.59.2	Examples	2048
19.59.3	Description	2048
19.59.4	Mixing, shift, table, tail correction, restart, rRESPA info	2049
19.59.5	Restrictions	2049
19.59.6	Related commands	2049
19.59.7	Default	2049
19.60	pair_style born/coul/dsf/cs command	2049
19.61	pair_style born/coul/long/cs command	2049
19.62	pair_style born/coul/wolf/cs command	2049
19.63	pair_style buck/coul/long/cs command	2049
19.64	pair_style coul/long/cs command	2049
19.65	pair_style coul/wolf/cs command	2050
19.66	pair_style lj/cut/coul/long/cs command	2050
19.67	pair_style lj/class2/coul/long/cs command	2050
19.67.1	Syntax	2050
19.67.2	Examples	2050
19.67.3	Description	2051
19.67.4	Mixing, shift, table, tail correction, restart, rRESPA info	2052
19.67.5	Restrictions	2052
19.67.6	Related commands	2052
19.67.7	Default	2052
19.68	pair_style coul/cut/dielectric command	2053
19.69	pair_style coul/long/dielectric command	2053
19.70	pair_style lj/cut/coul/cut/dielectric command	2053
19.71	pair_style lj/cut/coul/debye/dielectric command	2053
19.72	pair_style lj/cut/coul/long/dielectric command	2053
19.73	pair_style lj/cut/coul/msm/dielectric command	2053
19.74	pair_style lj/long/coul/long/dielectric command	2053
19.74.1	Syntax	2053
19.74.2	Examples	2053
19.74.3	Description	2054
19.74.4	Mixing, shift, table, tail correction, restart, rRESPA info	2054
19.74.5	Restrictions	2054
19.74.6	Related commands	2055

19.74.7	Default	2055
19.75	pair_style lj/cut/dipole/cut command	2055
19.76	pair_style lj/sf/dipole/sf command	2055
19.77	pair_style lj/cut/dipole/long command	2055
19.78	pair_style lj/long/dipole/long command	2055
19.78.1	Syntax	2055
19.78.2	Examples	2056
19.78.3	Description	2056
19.78.4	Mixing, shift, table, tail correction, restart, rRESPA info	2059
19.78.5	Restrictions	2060
19.78.6	Related commands	2060
19.78.7	Default	2060
19.79	pair_style dpd command	2060
19.80	pair_style dpd/tstat command	2060
19.80.1	Syntax	2060
19.80.2	Examples	2061
19.80.3	Description	2061
19.80.4	Mixing, shift, table, tail correction, restart, rRESPA info	2062
19.80.5	Restrictions	2063
19.80.6	Related commands	2063
19.80.7	Default	2063
19.81	pair_style dpd/ext command	2063
19.82	pair_style dpd/ext/tstat command	2063
19.82.1	Syntax	2063
19.82.2	Examples	2064
19.82.3	Description	2064
19.82.4	Restrictions	2066
19.82.5	Related commands	2066
19.83	pair_style dpd/fdt command	2066
19.84	pair_style dpd/fdt/energy command	2066
19.84.1	Syntax	2066
19.84.2	Examples	2067
19.84.3	Description	2067
19.84.4	Restrictions	2068
19.84.5	Related commands	2069
19.84.6	Default	2069
19.85	pair_style drip command	2069
19.85.1	Syntax	2069
19.85.2	Examples	2069
19.85.3	Description	2069
19.85.4	Mixing, shift, table, tail correction, restart, rRESPA info	2070
19.85.5	Restrictions	2071
19.85.6	Related commands	2071
19.86	pair_style dsmc command	2071
19.86.1	Syntax	2071
19.86.2	Examples	2071
19.86.3	Description	2072
19.86.4	Mixing, shift, table, tail correction, restart, rRESPA info	2073
19.86.5	Restrictions	2073
19.86.6	Related commands	2073
19.86.7	Default	2073
19.87	pair_style e3b command	2073
19.87.1	Syntax	2073
19.87.2	Examples	2074

19.87.3	Description	2074
19.87.4	Mixing, shift, table, tail correction, restart, rRESPA info	2075
19.87.5	Restrictions	2075
19.87.6	Related commands	2076
19.87.7	Default	2076
19.88	pair_style eam command	2076
19.89	pair_style eam/alloy command	2076
19.90	pair_style eam/cd command	2076
19.91	pair_style eam/cd/old command	2076
19.92	pair_style eam/fs command	2076
19.93	pair_style eam/he command	2076
19.93.1	Syntax	2076
19.93.2	Examples	2077
19.93.3	Description	2077
19.93.4	Mixing, shift, table, tail correction, restart, rRESPA info	2082
19.93.5	Restrictions	2082
19.93.6	Related commands	2082
19.93.7	Default	2082
19.94	pair_style edip command	2083
19.95	pair_style edip/multi command	2083
19.95.1	Syntax	2083
19.95.2	Examples	2083
19.95.3	Description	2083
19.95.4	Mixing, shift, table, tail correction, restart, rRESPA info	2085
19.95.5	Restrictions	2085
19.95.6	Related commands	2086
19.95.7	Default	2086
19.96	pair_style eff/cut command	2086
19.96.1	Syntax	2086
19.96.2	Examples	2086
19.96.3	Description	2086
19.96.4	Mixing, shift, table, tail correction, restart, rRESPA info	2089
19.96.5	Restrictions	2090
19.96.6	Related commands	2090
19.96.7	Default	2090
19.97	pair_style eim command	2090
19.97.1	Syntax	2090
19.97.2	Examples	2090
19.97.3	Description	2091
19.97.4	Restrictions	2093
19.97.5	Related commands	2093
19.97.6	Default	2093
19.98	pair_style exp6/rx command	2093
19.98.1	Syntax	2093
19.98.2	Examples	2093
19.98.3	Description	2094
19.98.4	Mixing, shift, table, tail correction, restart, rRESPA info	2095
19.98.5	Restrictions	2096
19.98.6	Related commands	2096
19.98.7	Default	2096
19.99	pair_style extep command	2096
19.99.1	Syntax	2096
19.99.2	Examples	2096
19.99.3	Description	2096



19.99.4	Restrictions	2096
19.99.5	Related commands	2096
19.99.6	Default	2097
19.100	pair_style lj/cut/soft command	2097
19.101	pair_style lj/cut/coul/cut/soft command	2097
19.102	pair_style lj/cut/coul/long/soft command	2097
19.103	pair_style lj/cut/tip4p/long/soft command	2097
19.104	pair_style lj/charmm/coul/long/soft command	2097
19.105	pair_style lj/class2/soft command	2097
19.106	pair_style lj/class2/coul/cut/soft command	2097
19.107	pair_style lj/class2/coul/long/soft command	2097
19.108	pair_style coul/cut/soft command	2097
19.109	pair_style coul/long/soft command	2098
19.110	pair_style tip4p/long/soft command	2098
19.111	pair_style morse/soft command	2098
19.111.1	Syntax	2098
19.111.2	Examples	2099
19.111.3	Description	2100
19.111.4	Mixing, shift, table, tail correction, restart, rRESPA info	2103
19.111.5	Restrictions	2103
19.111.6	Related commands	2104
19.111.7	Default	2104
19.112	pair_style gauss command	2104
19.113	pair_style gauss/cut command	2104
19.113.1	Syntax	2104
19.113.2	Examples	2104
19.113.3	Description	2104
19.113.4	Mixing, shift, table, tail correction, restart, rRESPA info	2105
19.113.5	Restrictions	2106
19.113.6	Related commands	2106
19.113.7	Default	2106
19.114	pair_style gayberne command	2107
19.114.1	Syntax	2107
19.114.2	Examples	2107
19.114.3	Description	2107
19.114.4	Mixing, shift, table, tail correction, restart, rRESPA info	2109
19.114.5	Restrictions	2109
19.114.6	Related commands	2109
19.114.7	Default	2109
19.115	pair_style gran/hooke command	2110
19.116	pair_style gran/hooke/history command	2110
19.117	pair_style gran/hertz/history command	2110
19.117.1	Syntax	2110
19.117.2	Examples	2111
19.117.3	Description	2111
19.117.4	Mixing, shift, table, tail correction, restart, rRESPA info	2113
19.117.5	Restrictions	2113
19.117.6	Related commands	2114
19.117.7	Default	2114
19.118	pair_style granular command	2114
19.118.1	Syntax	2114
19.118.2	Examples	2114
19.118.3	Description	2115
19.118.4	Mixing, shift, table, tail correction, restart, rRESPA info	2122

19.118.5	Restrictions	2123
19.118.6	Related commands	2123
19.118.7	Default	2123
19.118.8	References	2123
19.119	pair_style lj/gromacs command	2124
19.120	pair_style lj/gromacs/coul/gromacs command	2124
19.120.1	Syntax	2124
19.120.2	Examples	2124
19.120.3	Description	2125
19.120.4	Mixing, shift, table, tail correction, restart, rRESPA info	2126
19.120.5	Restrictions	2126
19.120.6	Related commands	2126
19.120.7	Default	2126
19.121	pair_style gw command	2126
19.122	pair_style gw/zbl command	2126
19.122.1	Syntax	2126
19.122.2	Examples	2127
19.122.3	Description	2127
19.122.4	Mixing, shift, table, tail correction, restart, rRESPA info	2128
19.122.5	Restrictions	2128
19.122.6	Related commands	2128
19.122.7	Default	2128
19.123	pair_style harmonic/cut command	2128
19.123.1	Syntax	2128
19.123.2	Examples	2129
19.123.3	Description	2129
19.123.4	Mixing, shift, table, tail correction, restart, rRESPA info	2129
19.123.5	Restrictions	2130
19.123.6	Related commands	2130
19.123.7	Default	2130
19.124	pair_style hbond/dreiding/lj command	2130
19.125	pair_style hbond/dreiding/morse command	2130
19.125.1	Syntax	2130
19.125.2	Examples	2130
19.125.3	Description	2131
19.125.4	Mixing, shift, table, tail correction, restart, rRESPA info	2133
19.125.5	Restrictions	2134
19.125.6	Related commands	2134
19.125.7	Default	2134
19.126	pair_style hdnp command	2134
19.126.1	Syntax	2134
19.126.2	Examples	2135
19.126.3	Description	2135
19.126.4	Mixing, shift, table, tail correction, restart, rRESPA info	2136
19.126.5	Restrictions	2137
19.127	pair_style hybrid command	2137
19.128	pair_style hybrid/overlay command	2137
19.129	pair_style hybrid/scaled command	2137
19.129.1	Syntax	2137
19.129.2	Examples	2138
19.129.3	Description	2138
19.129.4	Mixing, shift, table, tail correction, restart, rRESPA info	2143
19.129.5	Restrictions	2143
19.129.6	Related commands	2143

19.129.7Default	2143
19.130pair_style ilp/graphene/hbn command	2144
19.130.1Syntax	2144
19.130.2Examples	2144
19.130.3Description	2144
19.130.4Mixing, shift, table, tail correction, restart, rRESPA info	2146
19.130.5Restrictions	2146
19.130.6Related commands	2146
19.130.7Default	2146
19.131pair_style ilp/tmd command	2146
19.131.1Syntax	2147
19.131.2Examples	2147
19.131.3Description	2147
19.131.4Mixing, shift, table, tail correction, restart, rRESPA info	2148
19.131.5Restrictions	2148
19.131.6Related commands	2149
19.131.7Default	2149
19.132pair_style kim command	2149
19.132.1Syntax	2149
19.132.2Examples	2149
19.132.3Description	2149
19.132.4Mixing, shift, table, tail correction, restart, rRESPA info	2150
19.132.5Restrictions	2150
19.132.6Related commands	2150
19.132.7Default	2150
19.133pair_style kolmogorov/crespi/full command	2151
19.133.1Syntax	2151
19.133.2Examples	2151
19.133.3Description	2151
19.133.4Mixing, shift, table, tail correction, restart, rRESPA info	2152
19.133.5Restrictions	2152
19.133.6Related commands	2153
19.133.7Default	2153
19.134pair_style kolmogorov/crespi/z command	2153
19.134.1Syntax	2153
19.134.2Examples	2153
19.134.3Description	2153
19.134.4Restrictions	2154
19.134.5Related commands	2154
19.134.6Default	2154
19.135pair_style lcbop command	2154
19.135.1Syntax	2154
19.135.2Examples	2154
19.135.3Description	2155
19.135.4Mixing, shift, table, tail correction, restart, rRESPA info	2155
19.135.5Restrictions	2155
19.135.6Related commands	2156
19.135.7Default	2156
19.136pair_style lebedeva/z command	2156
19.136.1Syntax	2156
19.136.2Examples	2156
19.136.3Description	2156
19.136.4Restrictions	2157
19.136.5Related commands	2157

19.136.6	Default	2157
19.137	pair_style line/lj command	2157
19.137.1	Syntax	2157
19.137.2	Examples	2158
19.137.3	Description	2158
19.137.4	Mixing, shift, table, tail correction, restart, rRESPA info	2159
19.137.5	Restrictions	2159
19.137.6	Related commands	2159
19.137.7	Default	2159
19.138	pair_style list command	2159
19.138.1	Syntax	2159
19.138.2	Examples	2160
19.138.3	Description	2160
19.138.4	Mixing, shift, table, tail correction, restart, rRESPA info	2161
19.138.5	Restrictions	2161
19.138.6	Related commands	2162
19.138.7	Default	2162
19.139	pair_style lj/cut command	2162
19.139.1	Syntax	2162
19.139.2	Examples	2162
19.139.3	Description	2162
19.139.4	Coefficients	2163
19.139.5	Mixing, shift, table, tail correction, restart, rRESPA info	2163
19.139.6	Related commands	2164
19.139.7	Default	2164
19.140	pair_style lj96/cut command	2164
19.140.1	Syntax	2164
19.140.2	Examples	2164
19.140.3	Description	2164
19.140.4	Mixing, shift, table, tail correction, restart, rRESPA info	2165
19.140.5	Restrictions	2165
19.140.6	Related commands	2166
19.140.7	Default	2166
19.141	pair_style lj/cubic command	2166
19.141.1	Syntax	2166
19.141.2	Examples	2166
19.141.3	Description	2166
19.141.4	Mixing, shift, table, tail correction, restart, rRESPA info	2167
19.141.5	Restrictions	2167
19.141.6	Related commands	2167
19.141.7	Default	2168
19.142	pair_style lj/cut/coul/cut command	2168
19.143	pair_style lj/cut/coul/debye command	2168
19.144	pair_style lj/cut/coul/dsf command	2168
19.145	pair_style lj/cut/coul/long command	2168
19.146	pair_style lj/cut/coul/msm command	2168
19.147	pair_style lj/cut/coul/wolf command	2168
19.147.1	Syntax	2168
19.147.2	Examples	2169
19.147.3	Description	2170
19.147.4	Coefficients	2171
19.147.5	Mixing, shift, table, tail correction, restart, rRESPA info	2171
19.147.6	Restrictions	2172
19.147.7	Related commands	2172

19.147.8	Default	2172
19.148	pair_style lj/cut/tip4p/cut command	2172
19.149	pair_style lj/cut/tip4p/long command	2172
19.149.1	Syntax	2172
19.149.2	Examples	2173
19.149.3	Description	2173
19.149.4	Coefficients	2174
19.149.5	Mixing, shift, table, tail correction, restart, rRESPA info	2175
19.149.6	Restrictions	2175
19.149.7	Related commands	2175
19.149.8	Default	2175
19.150	pair_style lj/expand command	2175
19.151	pair_style lj/expand/coul/long command	2176
19.151.1	Syntax	2176
19.151.2	Examples	2176
19.151.3	Description	2176
19.151.4	Mixing, shift, table, tail correction, restart, rRESPA info	2177
19.151.5	Restrictions	2177
19.151.6	Related commands	2177
19.151.7	Default	2177
19.152	pair_style lj/long/coul/long command	2177
19.153	pair_style lj/long/tip4p/long command	2178
19.153.1	Syntax	2178
19.153.2	Examples	2178
19.153.3	Description	2179
19.153.4	Mixing, shift, table, tail correction, restart, rRESPA info	2180
19.153.5	Restrictions	2181
19.153.6	Related commands	2181
19.153.7	Default	2181
19.154	pair_style lj/relres command	2181
19.154.1	Syntax	2181
19.154.2	Examples	2181
19.154.3	Description	2181
19.154.4	Mixing, shift, table, tail correction, restart, rRESPA info	2184
19.154.5	Restrictions	2185
19.154.6	Related commands	2185
19.154.7	Default	2185
19.155	pair_style lj/smooth command	2185
19.155.1	Syntax	2185
19.155.2	Examples	2185
19.155.3	Description	2185
19.155.4	Mixing, shift, table, tail correction, restart, rRESPA info	2186
19.155.5	Restrictions	2187
19.155.6	Related commands	2187
19.155.7	Default	2187
19.156	pair_style lj/smooth/linear command	2187
19.156.1	Syntax	2187
19.156.2	Examples	2187
19.156.3	Description	2187
19.156.4	Mixing, shift, table, tail correction, restart, rRESPA info	2188
19.156.5	Restrictions	2188
19.156.6	Related commands	2188
19.156.7	Default	2189
19.157	pair_style lj/switch3/coulgauss/long command	2189

19.158	pair_style mm3/switch3/coulgauss/long command	2189
19.158.1	Syntax	2189
19.158.2	Examples	2189
19.158.3	Description	2190
19.158.4	Mixing, shift, table, tail correction, restart, rRESPA info	2190
19.158.5	Restrictions	2191
19.158.6	Related commands	2191
19.158.7	Default	2191
19.159	pair_style local/density command	2191
19.159.1	Syntax	2191
19.159.2	Examples	2191
19.159.3	Description	2191
19.159.4	Mixing, shift, table, tail correction, restart, rRESPA info	2194
19.159.5	Restrictions	2194
19.159.6	Related commands	2194
19.159.7	Default	2194
19.160	pair_style lubricate command	2195
19.161	pair_style lubricate/poly command	2195
19.161.1	Syntax	2195
19.161.2	Examples	2195
19.161.3	Description	2195
19.161.4	Mixing, shift, table, tail correction, restart, rRESPA info	2197
19.161.5	Restrictions	2197
19.161.6	Related commands	2198
19.161.7	Default	2198
19.162	pair_style lubricateU command	2198
19.163	pair_style lubricateU/poly command	2198
19.163.1	Syntax	2198
19.163.2	Examples	2198
19.163.3	Description	2199
19.163.4	Mixing, shift, table, tail correction, restart, rRESPA info	2200
19.163.5	Restrictions	2200
19.163.6	Related commands	2201
19.163.7	Default	2201
19.164	pair_style lj/mdf command	2201
19.165	pair_style buck/mdf command	2201
19.166	pair_style lennard/mdf command	2201
19.166.1	Syntax	2201
19.166.2	Examples	2202
19.166.3	Description	2202
19.166.4	Mixing, shift, table, tail correction, restart, rRESPA info	2203
19.166.5	Restrictions	2203
19.166.6	Related commands	2204
19.166.7	Default	2204
19.167	pair_style meam command	2204
19.167.1	Syntax	2204
19.167.2	Examples	2204
19.167.3	Description	2204
19.167.4	Mixing, shift, table, tail correction, restart, rRESPA info	2209
19.167.5	Restrictions	2209
19.167.6	Related commands	2210
19.167.7	Default	2210
19.168	pair_style meam/spline command	2210
19.168.1	Syntax	2210

19.168.2	Examples	2210
19.168.3	Description	2210
19.168.4	Mixing, shift, table, tail correction, restart, rRESPA info	2212
19.168.5	Restrictions	2212
19.168.6	Related commands	2212
19.168.7	Default	2212
19.169	pair_style meam/sw/spline command	2213
19.169.1	Syntax	2213
19.169.2	Examples	2213
19.169.3	Description	2213
19.169.4	Mixing, shift, table, tail correction, restart, rRESPA info	2214
19.169.5	Restrictions	2214
19.169.6	Related commands	2214
19.169.7	Default	2214
19.170	pair_style mesocnt command	2215
19.171	pair_style mesocnt/viscous command	2215
19.171.1	Syntax	2215
19.171.2	Examples	2215
19.171.3	Description	2215
19.171.4	Mixing, shift, table, tail correction, restart, rRESPA info	2217
19.171.5	Restrictions	2217
19.171.6	Related commands	2218
19.171.7	Default	2218
19.172	pair_style edpd command	2218
19.173	pair_style mdpd command	2218
19.174	pair_style mdpd/rhosum command	2218
19.175	pair_style tdpd command	2218
19.175.1	Syntax	2218
19.175.2	Examples	2219
19.175.3	Description	2219
19.175.4	Example scripts	2222
19.175.5	Mixing, shift, table, tail correction, restart, rRESPA info	2223
19.175.6	Restrictions	2224
19.175.7	Related commands	2224
19.175.8	Default	2224
19.176	pair_style mesont/tpm command	2224
19.176.1	Syntax	2224
19.176.2	Examples	2225
19.176.3	Description	2225
19.176.4	Restrictions	2226
19.176.5	Related commands	2226
19.177	pair_style mgpt command	2227
19.177.1	Syntax	2227
19.177.2	Examples	2227
19.177.3	Description	2227
19.177.4	Mixing, shift, table, tail correction, restart, rRESPA info	2229
19.177.5	Restrictions	2229
19.177.6	Related commands	2229
19.177.7	Default	2229
19.178	pair_style mie/cut command	2230
19.178.1	Syntax	2230
19.178.2	Examples	2230
19.178.3	Description	2230
19.178.4	Mixing, shift, table, tail correction, restart, rRESPA info	2231

19.178.5	Restrictions	2231
19.178.6	Related commands	2231
19.178.7	Default	2231
19.179	pair_style mliap command	2231
19.179.1	Syntax	2231
19.179.2	Examples	2232
19.179.3	Description	2232
19.179.4	Mixing, shift, table, tail correction, restart, rRESPA info	2234
19.179.5	Restrictions	2234
19.179.6	Related commands	2234
19.179.7	Default	2234
19.180	pair_style momb command	2234
19.180.1	Syntax	2234
19.180.2	Examples	2235
19.180.3	Description	2235
19.180.4	Restrictions	2235
19.180.5	Related commands	2235
19.180.6	Default	2235
19.181	pair_style morse command	2236
19.182	pair_style morse/smooth/linear command	2236
19.182.1	Syntax	2236
19.182.2	Examples	2236
19.182.3	Description	2236
19.182.4	Mixing, shift, table, tail correction, restart, rRESPA info	2237
19.182.5	Restrictions	2238
19.182.6	Related commands	2238
19.182.7	Default	2238
19.183	pair_style multi/lucy command	2238
19.183.1	Syntax	2238
19.183.2	Examples	2238
19.183.3	Description	2238
19.183.4	Mixing, shift, table, tail correction, restart, rRESPA info	2240
19.183.5	Restrictions	2240
19.183.6	Related commands	2240
19.183.7	Default	2241
19.184	pair_style multi/lucy/rx command	2241
19.184.1	Syntax	2241
19.184.2	Examples	2241
19.184.3	Description	2241
19.184.4	Mixing, shift, table, tail correction, restart, rRESPA info	2243
19.184.5	Restrictions	2244
19.184.6	Related commands	2244
19.184.7	Default	2244
19.185	pair_style nb3b/harmonic command	2244
19.185.1	Syntax	2244
19.185.2	Examples	2244
19.185.3	Description	2245
19.185.4	Restrictions	2246
19.185.5	Related commands	2246
19.185.6	Default	2246
19.186	pair_style nm/cut command	2246
19.187	pair_style nm/cut/split command	2246
19.188	pair_style nm/cut/coul/cut command	2246
19.189	pair_style nm/cut/coul/long command	2246



19.189.1	Syntax	2246
19.189.2	Examples	2247
19.189.3	Description	2247
19.189.4	Mixing, shift, table, tail correction, restart, rRESPA info	2248
19.189.5	Restrictions	2248
19.189.6	Related commands	2249
19.189.7	Default	2249
19.190	pair_style none command	2249
19.190.1	Syntax	2249
19.190.2	Examples	2249
19.190.3	Description	2249
19.190.4	Restrictions	2250
19.190.5	Related commands	2250
19.190.6	Default	2250
19.191	pair_style oxdna/excv command	2250
19.192	pair_style oxdna/stk command	2250
19.193	pair_style oxdna/hbond command	2250
19.194	pair_style oxdna/xstk command	2250
19.195	pair_style oxdna/coaxstk command	2250
19.195.1	Syntax	2250
19.195.2	Examples	2251
19.195.3	Description	2251
19.195.4	Restrictions	2252
19.195.5	Related commands	2252
19.195.6	Default	2252
19.196	pair_style oxdna2/excv command	2252
19.197	pair_style oxdna2/stk command	2252
19.198	pair_style oxdna2/hbond command	2252
19.199	pair_style oxdna2/xstk command	2252
19.200	pair_style oxdna2/coaxstk command	2252
19.201	pair_style oxdna2/dh command	2252
19.201.1	Syntax	2252
19.201.2	Examples	2253
19.201.3	Description	2254
19.201.4	Restrictions	2254
19.201.5	Related commands	2255
19.201.6	Default	2255
19.202	pair_style oxrna2/excv command	2255
19.203	pair_style oxrna2/stk command	2255
19.204	pair_style oxrna2/hbond command	2255
19.205	pair_style oxrna2/xstk command	2255
19.206	pair_style oxrna2/coaxstk command	2255
19.207	pair_style oxrna2/dh command	2255
19.207.1	Syntax	2255
19.207.2	Examples	2256
19.207.3	Description	2256
19.207.4	Restrictions	2257
19.207.5	Related commands	2257
19.207.6	Default	2257
19.208	pair_style pace command	2258
19.208.1	Syntax	2258
19.208.2	Examples	2258
19.208.3	Description	2258
19.208.4	Mixing, shift, table, tail correction, restart, rRESPA info	2259

19.208.5	Restrictions	2259
19.208.6	Related commands	2259
19.208.7	Default	2259
19.209	pair_style peri/pmb command	2260
19.210	pair_style peri/lps command	2260
19.211	pair_style peri/ves command	2260
19.212	pair_style peri/eps command	2260
19.212.1	Syntax	2260
19.212.2	Examples	2260
19.212.3	Description	2260
19.212.4	Mixing, shift, table, tail correction, restart, rRESPA info	2262
19.212.5	Restrictions	2262
19.212.6	Related commands	2262
19.212.7	Default	2263
19.213	pair_style polymorphic command	2263
19.213.1	Syntax	2263
19.213.2	Examples	2263
19.213.3	Description	2263
19.213.4	Mixing, shift, table, tail correction, restart, rRESPA info	2267
19.213.5	Restrictions	2267
19.213.6	Related commands	2267
19.214	pair_style python command	2268
19.214.1	Syntax	2268
19.214.2	Examples	2268
19.214.3	Description	2268
19.214.4	Mixing, shift, table, tail correction, restart, rRESPA info	2271
19.214.5	Restrictions	2272
19.214.6	Related commands	2272
19.214.7	Default	2272
19.215	pair_style quip command	2272
19.215.1	Syntax	2272
19.215.2	Examples	2272
19.215.3	Description	2272
19.215.4	Mixing, shift, table, tail correction, restart, rRESPA info	2273
19.215.5	Restrictions	2273
19.215.6	Related commands	2273
19.216	pair_style rann command	2273
19.216.1	Syntax	2273
19.216.2	Examples	2274
19.216.3	Description	2274
19.216.4	Potential file syntax	2274
19.216.5	Formulation	2277
19.216.6	Restrictions	2279
19.216.7	Defaults	2279
19.217	pair_style reaxff command	2279
19.217.1	Syntax	2280
19.217.2	Examples	2280
19.217.3	Description	2280
19.217.4	Control file	2283
19.217.5	Mixing, shift, table, tail correction, restart, rRESPA info	2284
19.217.6	Restrictions	2284
19.217.7	Related commands	2285
19.217.8	Default	2285
19.218	pair_style resquared command	2285

19.218.1	Syntax . . . . .	2285
19.218.2	Examples . . . . .	2285
19.218.3	Description . . . . .	2285
19.218.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2287
19.218.5	Restrictions . . . . .	2287
19.218.6	Related commands . . . . .	2288
19.218.7	Default . . . . .	2288
19.219	pair_style saip/metal command . . . . .	2288
19.219.1	Syntax . . . . .	2288
19.219.2	Examples . . . . .	2288
19.219.3	Description . . . . .	2289
19.219.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2290
19.219.5	Restrictions . . . . .	2290
19.219.6	Related commands . . . . .	2290
19.219.7	Default . . . . .	2290
19.220	pair_style sdpd/taitwater/isothermal command . . . . .	2291
19.220.1	Syntax . . . . .	2291
19.220.2	Examples . . . . .	2291
19.220.3	Description . . . . .	2291
19.220.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2292
19.220.5	Restrictions . . . . .	2292
19.220.6	Related commands . . . . .	2292
19.220.7	Default . . . . .	2292
19.221	pair_style smatb command . . . . .	2292
19.222	pair_style smatb/single command . . . . .	2292
19.222.1	Syntax . . . . .	2292
19.222.2	Examples . . . . .	2293
19.222.3	Description . . . . .	2293
19.222.4	Coefficients . . . . .	2293
19.222.5	Mixing info . . . . .	2294
19.222.6	Restrictions . . . . .	2294
19.222.7	Related commands . . . . .	2294
19.222.8	Default . . . . .	2294
19.223	pair_style smd/hertz command . . . . .	2294
19.223.1	Syntax . . . . .	2294
19.223.2	Examples . . . . .	2294
19.223.3	Description . . . . .	2294
19.223.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2295
19.223.5	Restrictions . . . . .	2295
19.223.6	Related commands . . . . .	2295
19.223.7	Default . . . . .	2295
19.224	pair_style smd/tlsph command . . . . .	2295
19.224.1	Syntax . . . . .	2295
19.224.2	Examples . . . . .	2295
19.224.3	Description . . . . .	2295
19.224.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2296
19.224.5	Restrictions . . . . .	2296
19.224.6	Related commands . . . . .	2296
19.224.7	Default . . . . .	2296
19.225	pair_style smd/tri_surface command . . . . .	2296
19.225.1	Syntax . . . . .	2296
19.225.2	Examples . . . . .	2297
19.225.3	Description . . . . .	2297
19.225.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2297

19.225.5	Restrictions	2297
19.225.6	Related commands	2297
19.225.7	Default	2297
19.226	pair_style smd/ulsph command	2297
19.226.1	Syntax	2297
19.226.2	Examples	2298
19.226.3	Description	2298
19.226.4	Mixing, shift, table, tail correction, restart, rRESPA info	2298
19.226.5	Restrictions	2298
19.226.6	Related commands	2299
19.226.7	Default	2299
19.227	pair_style smtbq command	2299
19.227.1	Syntax	2299
19.227.2	Examples	2299
19.227.3	Description	2299
19.227.4	Mixing, shift, table, tail correction, restart, rRESPA info	2302
19.227.5	Restrictions	2303
19.227.6	Citing this work	2303
19.228	pair_style snap command	2303
19.228.1	Syntax	2303
19.228.2	Examples	2303
19.228.3	Description	2303
19.228.4	Mixing, shift, table, tail correction, restart, rRESPA info	2306
19.228.5	Restrictions	2307
19.228.6	Related commands	2307
19.228.7	Default	2307
19.229	pair_style soft command	2307
19.229.1	Syntax	2307
19.229.2	Examples	2307
19.229.3	Description	2308
19.229.4	Mixing, shift, table, tail correction, restart, rRESPA info	2309
19.229.5	Restrictions	2309
19.229.6	Related commands	2309
19.229.7	Default	2309
19.230	pair_style sph/heatconduction command	2309
19.230.1	Syntax	2309
19.230.2	Examples	2309
19.230.3	Description	2310
19.230.4	Mixing, shift, table, tail correction, restart, rRESPA info	2310
19.230.5	Restrictions	2310
19.230.6	Related commands	2310
19.230.7	Default	2310
19.231	pair_style sph/idealgas command	2310
19.231.1	Syntax	2310
19.231.2	Examples	2311
19.231.3	Description	2311
19.231.4	Mixing, shift, table, tail correction, restart, rRESPA info	2311
19.231.5	Restrictions	2311
19.231.6	Related commands	2311
19.231.7	Default	2312
19.232	pair_style sph/lj command	2312
19.232.1	Syntax	2312
19.232.2	Examples	2312
19.232.3	Description	2312

19.232.4	Mixing, shift, table, tail correction, restart, rRESPA info	2312
19.232.5	Restrictions	2313
19.232.6	Related commands	2313
19.232.7	Default	2313
19.233	pair_style sph/rhosome command	2313
19.233.1	Syntax	2313
19.233.2	Examples	2313
19.233.3	Description	2313
19.233.4	Mixing, shift, table, tail correction, restart, rRESPA info	2314
19.233.5	Restrictions	2314
19.233.6	Related commands	2314
19.233.7	Default	2314
19.234	pair_style sph/taitwater command	2314
19.234.1	Syntax	2314
19.234.2	Examples	2314
19.234.3	Description	2314
19.234.4	Mixing, shift, table, tail correction, restart, rRESPA info	2315
19.234.5	Restrictions	2315
19.234.6	Related commands	2315
19.234.7	Default	2315
19.235	pair_style sph/taitwater/morris command	2315
19.235.1	Syntax	2315
19.235.2	Examples	2316
19.235.3	Description	2316
19.235.4	Mixing, shift, table, tail correction, restart, rRESPA info	2316
19.235.5	Restrictions	2316
19.235.6	Related commands	2317
19.235.7	Default	2317
19.236	pair_style lj/spica command	2317
19.237	pair_style lj/spica/coul/long command	2317
19.238	pair_style lj/spica/coul/msm command	2317
19.238.1	Syntax	2317
19.238.2	Examples	2317
19.238.3	Description	2318
19.238.4	Mixing, shift, table, tail correction, restart, rRESPA info	2319
19.238.5	Restrictions	2319
19.238.6	Related commands	2319
19.238.7	Default	2319
19.239	pair_style spin/dipole/cut command	2320
19.240	pair_style spin/dipole/long command	2320
19.240.1	Syntax	2320
19.240.2	Examples	2320
19.240.3	Description	2320
19.240.4	Restrictions	2321
19.240.5	Related commands	2321
19.240.6	Default	2321
19.241	pair_style spin/dmi command	2321
19.241.1	Syntax	2321
19.241.2	Examples	2321
19.241.3	Description	2321
19.241.4	Restrictions	2322
19.241.5	Related commands	2322
19.241.6	Default	2322
19.242	pair_style spin/exchange command	2322

19.243	pair_style spin/exchange/biquadratic command	2322
19.243.1	Syntax	2322
19.243.2	Examples	2323
19.243.3	Description	2323
19.243.4	Restrictions	2325
19.243.5	Related commands	2325
19.243.6	Default	2325
19.244	pair_style spin/magelec command	2325
19.244.1	Syntax	2325
19.244.2	Examples	2325
19.244.3	Description	2325
19.244.4	Restrictions	2326
19.244.5	Related commands	2326
19.244.6	Default	2326
19.245	pair_style spin/neel command	2326
19.245.1	Syntax	2326
19.245.2	Examples	2327
19.245.3	Description	2327
19.245.4	Restrictions	2327
19.245.5	Related commands	2328
19.245.6	Default	2328
19.246	pair_style srp command	2328
19.247	pair_style srp/react command	2328
19.247.1	Syntax	2328
19.247.2	Examples	2328
19.247.3	Mixing, shift, table, tail correction, restart, rRESPA info	2330
19.247.4	Restrictions	2330
19.247.5	Related commands	2330
19.247.6	Default	2330
19.248	pair_style sw command	2331
19.249	pair_style sw/mod command	2331
19.249.1	Syntax	2331
19.249.2	Examples	2331
19.249.3	Description	2331
19.249.4	Mixing, shift, table, tail correction, restart, rRESPA info	2334
19.249.5	Restrictions	2335
19.249.6	Related commands	2335
19.249.7	Default	2335
19.250	pair_style sw/angle/table command	2335
19.250.1	Syntax	2335
19.250.2	Examples	2335
19.250.3	Description	2336
19.250.4	Mixing, shift, table, tail correction, restart, rRESPA info	2339
19.250.5	Restrictions	2339
19.250.6	Related commands	2339
19.251	pair_style table command	2339
19.251.1	Syntax	2339
19.251.2	Examples	2340
19.251.3	Description	2340
19.251.4	Mixing, shift, table, tail correction, restart, rRESPA info	2342
19.251.5	Restrictions	2343
19.251.6	Related commands	2343
19.251.7	Default	2343
19.252	pair_style table/rx command	2343

19.252.1	Syntax . . . . .	2343
19.252.2	Examples . . . . .	2343
19.252.3	Description . . . . .	2343
19.252.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2346
19.252.5	Restrictions . . . . .	2346
19.252.6	Related commands . . . . .	2346
19.252.7	Default . . . . .	2346
19.253	pair_style tersoff command . . . . .	2347
19.254	pair_style tersoff/table command . . . . .	2347
19.254.1	Syntax . . . . .	2347
19.254.2	Examples . . . . .	2347
19.254.3	Description . . . . .	2348
19.254.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2351
19.254.5	Restrictions . . . . .	2351
19.254.6	Related commands . . . . .	2351
19.254.7	Default . . . . .	2351
19.255	pair_style tersoff/mod command . . . . .	2352
19.256	pair_style tersoff/mod/c command . . . . .	2352
19.256.1	Syntax . . . . .	2352
19.256.2	Examples . . . . .	2352
19.256.3	Description . . . . .	2353
19.256.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2355
19.256.5	Restrictions . . . . .	2355
19.256.6	Related commands . . . . .	2355
19.256.7	Default . . . . .	2355
19.257	pair_style tersoff/zbl command . . . . .	2356
19.257.1	Syntax . . . . .	2356
19.257.2	Examples . . . . .	2356
19.257.3	Description . . . . .	2356
19.257.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2360
19.257.5	Restrictions . . . . .	2360
19.257.6	Related commands . . . . .	2360
19.257.7	Default . . . . .	2360
19.258	pair_style thole command . . . . .	2361
19.259	pair_style lj/cut/thole/long command . . . . .	2361
19.259.1	Syntax . . . . .	2361
19.259.2	Examples . . . . .	2361
19.259.3	Description . . . . .	2361
19.259.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2363
19.259.5	Restrictions . . . . .	2363
19.259.6	Related commands . . . . .	2363
19.259.7	Default . . . . .	2363
19.260	pair_style threebody/table command . . . . .	2363
19.260.1	Syntax . . . . .	2363
19.260.2	Examples . . . . .	2364
19.260.3	Description . . . . .	2364
19.260.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2366
19.260.5	Restrictions . . . . .	2367
19.260.6	Related commands . . . . .	2367
19.261	pair_style tracker command . . . . .	2367
19.261.1	Syntax . . . . .	2367
19.261.2	Examples . . . . .	2368
19.261.3	Description . . . . .	2368
19.261.4	Mixing, shift, table, tail correction, restart, rRESPA info . . . . .	2369

19.261.5	Restrictions	2369
19.261.6	Related commands	2369
19.261.7	Default	2369
19.262	pair_style tri/lj command	2369
19.262.1	Syntax	2369
19.262.2	Examples	2370
19.262.3	Description	2370
19.262.4	Mixing, shift, table, tail correction, restart, rRESPA info	2371
19.262.5	Restrictions	2371
19.262.6	Related commands	2371
19.262.7	Default	2371
19.263	pair_style ufm command	2371
19.263.1	Syntax	2371
19.263.2	Examples	2371
19.263.3	Description	2372
19.263.4	Mixing, shift, table, tail correction, restart, rRESPA info	2373
19.263.5	Restrictions	2373
19.263.6	Related commands	2373
19.263.7	Default	2373
19.264	pair_style vashishta command	2373
19.265	pair_style vashishta/table command	2373
19.265.1	Syntax	2374
19.265.2	Examples	2374
19.265.3	Description	2374
19.265.4	Mixing, shift, table, tail correction, restart, rRESPA info	2376
19.265.5	Restrictions	2377
19.265.6	Related commands	2377
19.265.7	Default	2377
19.266	pair_style wf/cut command	2377
19.266.1	Syntax	2377
19.266.2	Examples	2377
19.266.3	Description	2377
19.266.4	Restrictions	2379
19.266.5	Related commands	2379
19.267	pair_style yukawa command	2379
19.267.1	Syntax	2379
19.267.2	Examples	2379
19.267.3	Description	2379
19.267.4	Mixing, shift, table, tail correction, restart, rRESPA info	2380
19.267.5	Restrictions	2380
19.267.6	Related commands	2380
19.267.7	Default	2380
19.268	pair_style yukawa/colloid command	2381
19.268.1	Syntax	2381
19.268.2	Examples	2381
19.268.3	Description	2381
19.268.4	Mixing, shift, table, tail correction, restart, rRESPA info	2382
19.268.5	Restrictions	2382
19.268.6	Related commands	2383
19.268.7	Default	2383
19.269	pair_style zbl command	2383
19.269.1	Syntax	2383
19.269.2	Examples	2383
19.269.3	Description	2383



19.269.4	Mixing, shift, table, tail correction, restart, rRESPA info	2384
19.269.5	Restrictions	2385
19.269.6	Related commands	2385
19.269.7	Default	2385
19.270	pair_style zero command	2385
19.270.1	Syntax	2385
19.270.2	Examples	2385
19.270.3	Description	2385
19.270.4	Mixing, shift, table, tail correction, restart, rRESPA info	2386
19.270.5	Restrictions	2386
19.270.6	Related commands	2386
19.270.7	Default	2386
<b>20</b>	<b>Bond Styles</b>	<b>2387</b>
20.1	bond_style bpm/rotational command	2387
20.1.1	Syntax	2387
20.1.2	Examples	2387
20.1.3	Description	2388
20.1.4	Restart and other info	2389
20.1.5	Restrictions	2390
20.1.6	Related commands	2390
20.1.7	Default	2390
20.2	bond_style bpm/spring command	2390
20.2.1	Syntax	2390
20.2.2	Examples	2391
20.2.3	Description	2391
20.2.4	Restart and other info	2392
20.2.5	Restrictions	2392
20.2.6	Related commands	2393
20.2.7	Default	2393
20.3	bond_style class2 command	2393
20.3.1	Syntax	2393
20.3.2	Examples	2393
20.3.3	Description	2393
20.3.4	Restrictions	2394
20.3.5	Related commands	2394
20.3.6	Default	2394
20.4	bond_style fene command	2394
20.5	bond_style fene/nm command	2394
20.5.1	Syntax	2394
20.5.2	Examples	2394
20.5.3	Description	2395
20.5.4	Restrictions	2396
20.5.5	Related commands	2396
20.5.6	Default	2396
20.6	bond_style fene/expand command	2396
20.6.1	Syntax	2396
20.6.2	Examples	2396
20.6.3	Description	2396
20.6.4	Restrictions	2397
20.6.5	Related commands	2397
20.6.6	Default	2397
20.7	bond_style gaussian command	2397
20.7.1	Syntax	2397

20.7.2	Examples	2398
20.7.3	Description	2398
20.7.4	Restrictions	2398
20.7.5	Related commands	2398
20.7.6	Default	2398
20.8	bond_style gromos command	2399
20.8.1	Syntax	2399
20.8.2	Examples	2399
20.8.3	Description	2399
20.8.4	Restrictions	2400
20.8.5	Related commands	2400
20.8.6	Default	2400
20.9	bond_style harmonic command	2400
20.9.1	Syntax	2400
20.9.2	Examples	2400
20.9.3	Description	2400
20.9.4	Restrictions	2401
20.9.5	Related commands	2401
20.9.6	Default	2401
20.10	bond_style harmonic/shift command	2401
20.10.1	Syntax	2401
20.10.2	Examples	2401
20.10.3	Description	2401
20.10.4	Restrictions	2402
20.10.5	Related commands	2402
20.10.6	Default	2402
20.11	bond_style harmonic/shift/cut command	2402
20.11.1	Syntax	2402
20.11.2	Examples	2402
20.11.3	Description	2403
20.11.4	Restrictions	2403
20.11.5	Related commands	2403
20.11.6	Default	2403
20.12	bond_style hybrid command	2404
20.12.1	Syntax	2404
20.12.2	Examples	2404
20.12.3	Description	2404
20.12.4	Restrictions	2404
20.12.5	Related commands	2405
20.12.6	Default	2405
20.13	bond_style mesocnt command	2405
20.13.1	Syntax	2405
20.13.2	Examples	2405
20.13.3	Description	2405
20.13.4	Restrictions	2406
20.13.5	Related commands	2406
20.13.6	Default	2406
20.14	bond_style mm3 command	2406
20.14.1	Syntax	2406
20.14.2	Examples	2406
20.14.3	Description	2406
20.14.4	Restrictions	2407
20.14.5	Related commands	2407
20.14.6	Default	2407

20.15	bond_style morse command	2407
20.15.1	Syntax	2407
20.15.2	Examples	2407
20.15.3	Description	2407
20.15.4	Restrictions	2408
20.15.5	Related commands	2408
20.15.6	Default	2408
20.16	bond_style none command	2408
20.16.1	Syntax	2408
20.16.2	Examples	2408
20.16.3	Description	2409
20.16.4	Restrictions	2409
20.16.5	Related commands	2409
20.16.6	Default	2409
20.17	bond_style nonlinear command	2409
20.17.1	Syntax	2409
20.17.2	Examples	2409
20.17.3	Description	2409
20.17.4	Restrictions	2410
20.17.5	Related commands	2410
20.17.6	Default	2410
20.18	bond_style oxdna/fene command	2410
20.19	bond_style oxdna2/fene command	2410
20.20	bond_style oxrna2/fene command	2410
20.20.1	Syntax	2410
20.20.2	Examples	2411
20.20.3	Description	2411
20.20.4	Restrictions	2412
20.20.5	Related commands	2412
20.20.6	Default	2412
20.21	bond_style quartic command	2412
20.21.1	Syntax	2413
20.21.2	Examples	2413
20.21.3	Description	2413
20.21.4	Restrictions	2414
20.21.5	Related commands	2414
20.21.6	Default	2414
20.22	bond_style special command	2414
20.22.1	Syntax	2414
20.22.2	Examples	2415
20.22.3	Description	2415
20.22.4	Restrictions	2416
20.22.5	Related commands	2416
20.22.6	Default	2416
20.23	bond_style table command	2416
20.23.1	Syntax	2416
20.23.2	Examples	2416
20.23.3	Description	2416
20.23.4	Restart, fix_modify, output, run start/stop, minimize info	2418
20.23.5	Restrictions	2418
20.23.6	Related commands	2418
20.23.7	Default	2418
20.24	bond_style zero command	2418
20.24.1	Syntax	2418

20.24.2	Examples	2419
20.24.3	Description	2419
20.24.4	Restrictions	2419
20.24.5	Related commands	2419
20.24.6	Default	2419
<b>21</b>	<b>Angle Styles</b>	<b>2421</b>
21.1	angle_style amoeba command	2421
21.1.1	Syntax	2421
21.1.2	Examples	2421
21.1.3	Description	2421
21.1.4	Restrictions	2423
21.1.5	Related commands	2423
21.1.6	Default	2423
21.2	angle_style charmm command	2423
21.2.1	Syntax	2423
21.2.2	Examples	2423
21.2.3	Description	2423
21.2.4	Restrictions	2424
21.2.5	Related commands	2424
21.2.6	Default	2424
21.3	angle_style class2 command	2424
21.4	angle_style class2/p6 command	2424
21.4.1	Syntax	2424
21.4.2	Examples	2425
21.4.3	Description	2425
21.4.4	Restrictions	2426
21.4.5	Related commands	2427
21.4.6	Default	2427
21.5	angle_style cosine command	2427
21.5.1	Syntax	2427
21.5.2	Examples	2427
21.5.3	Description	2427
21.5.4	Restrictions	2428
21.5.5	Related commands	2428
21.5.6	Default	2428
21.6	angle_style cosine/buck6d command	2428
21.6.1	Syntax	2428
21.6.2	Examples	2428
21.6.3	Description	2428
21.6.4	Restrictions	2429
21.6.5	Related commands	2429
21.6.6	Default	2429
21.7	angle_style cosine/delta command	2429
21.7.1	Syntax	2429
21.7.2	Examples	2429
21.7.3	Description	2429
21.7.4	Restrictions	2430
21.7.5	Related commands	2430
21.7.6	Default	2430
21.8	angle_style cosine/periodic command	2430
21.8.1	Syntax	2430
21.8.2	Examples	2430
21.8.3	Description	2431

21.8.4	Restrictions	2431
21.8.5	Related commands	2431
21.8.6	Default	2431
21.9	angle_style cosine/shift command	2432
21.9.1	Syntax	2432
21.9.2	Examples	2432
21.9.3	Description	2432
21.9.4	Restrictions	2433
21.9.5	Related commands	2433
21.9.6	Default	2433
21.10	angle_style cosine/shift/exp command	2433
21.10.1	Syntax	2433
21.10.2	Examples	2433
21.10.3	Description	2433
21.10.4	Restrictions	2434
21.10.5	Related commands	2434
21.10.6	Default	2434
21.11	angle_style cosine/squared command	2434
21.11.1	Syntax	2434
21.11.2	Examples	2434
21.11.3	Description	2435
21.11.4	Restrictions	2435
21.11.5	Related commands	2435
21.11.6	Default	2435
21.12	angle_style cross command	2436
21.12.1	Syntax	2436
21.12.2	Examples	2436
21.12.3	Description	2436
21.12.4	Restrictions	2436
21.12.5	Related commands	2437
21.12.6	Default	2437
21.13	angle_style dipole command	2437
21.13.1	Syntax	2437
21.13.2	Examples	2437
21.13.3	Description	2437
21.13.4	Restrictions	2438
21.13.5	Related commands	2439
21.13.6	Default	2439
21.14	angle_style fourier command	2439
21.14.1	Syntax	2439
21.14.2	Examples	2439
21.14.3	Description	2439
21.14.4	Restrictions	2440
21.14.5	Related commands	2440
21.14.6	Default	2440
21.15	angle_style fourier/simple command	2440
21.15.1	Syntax	2440
21.15.2	Examples	2440
21.15.3	Description	2440
21.15.4	Restrictions	2441
21.15.5	Related commands	2441
21.15.6	Default	2441
21.16	angle_style gaussian command	2441
21.16.1	Syntax	2441

21.16.2	Examples	2441
21.16.3	Description	2442
21.16.4	Restrictions	2442
21.16.5	Related commands	2442
21.16.6	Default	2442
21.17	angle_style harmonic command	2442
21.17.1	Syntax	2443
21.17.2	Examples	2443
21.17.3	Description	2443
21.17.4	Restrictions	2443
21.17.5	Related commands	2444
21.17.6	Default	2444
21.18	angle_style hybrid command	2444
21.18.1	Syntax	2444
21.18.2	Examples	2444
21.18.3	Description	2444
21.18.4	Restrictions	2445
21.18.5	Related commands	2445
21.18.6	Default	2445
21.19	angle_style mesocnt command	2445
21.19.1	Syntax	2445
21.19.2	Examples	2445
21.19.3	Description	2446
21.19.4	Restrictions	2447
21.19.5	Related commands	2447
21.19.6	Default	2447
21.20	angle_style mm3 command	2447
21.20.1	Syntax	2447
21.20.2	Examples	2447
21.20.3	Description	2448
21.20.4	Restrictions	2448
21.20.5	Related commands	2448
21.20.6	Default	2448
21.21	angle_style none command	2448
21.21.1	Syntax	2448
21.21.2	Examples	2448
21.21.3	Description	2449
21.21.4	Restrictions	2449
21.21.5	Related commands	2449
21.21.6	Default	2449
21.22	angle_style quartic command	2449
21.22.1	Syntax	2449
21.22.2	Examples	2449
21.22.3	Description	2449
21.22.4	Restrictions	2450
21.22.5	Related commands	2450
21.22.6	Default	2450
21.23	angle_style spica command	2450
21.23.1	Syntax	2450
21.23.2	Examples	2451
21.23.3	Description	2451
21.23.4	Restrictions	2451
21.23.5	Related commands	2452
21.23.6	Default	2452

21.24	angle_style table command . . . . .	2452
21.24.1	Syntax . . . . .	2452
21.24.2	Examples . . . . .	2452
21.24.3	Description . . . . .	2452
21.24.4	Restart, fix_modify, output, run start/stop, minimize info . . . . .	2454
21.24.5	Restrictions . . . . .	2454
21.24.6	Related commands . . . . .	2454
21.24.7	Default . . . . .	2454
21.25	angle_style zero command . . . . .	2454
21.25.1	Syntax . . . . .	2454
21.25.2	Examples . . . . .	2454
21.25.3	Description . . . . .	2454
21.25.4	Restrictions . . . . .	2455
21.25.5	Related commands . . . . .	2455
21.25.6	Default . . . . .	2455
<b>22</b>	<b>Dihedral Styles . . . . .</b>	<b>2457</b>
22.1	dihedral_style charmm command . . . . .	2457
22.2	dihedral_style charmmfsw command . . . . .	2457
22.2.1	Syntax . . . . .	2457
22.2.2	Examples . . . . .	2457
22.2.3	Description . . . . .	2457
22.2.4	Restrictions . . . . .	2459
22.2.5	Related commands . . . . .	2459
22.2.6	Default . . . . .	2459
22.3	dihedral_style class2 command . . . . .	2459
22.3.1	Syntax . . . . .	2459
22.3.2	Examples . . . . .	2459
22.3.3	Description . . . . .	2460
22.3.4	Restrictions . . . . .	2462
22.3.5	Related commands . . . . .	2462
22.3.6	Default . . . . .	2462
22.4	dihedral_style cosine/shift/exp command . . . . .	2462
22.4.1	Syntax . . . . .	2462
22.4.2	Examples . . . . .	2462
22.4.3	Description . . . . .	2463
22.4.4	Restrictions . . . . .	2463
22.4.5	Related commands . . . . .	2463
22.4.6	Default . . . . .	2464
22.5	dihedral_style fourier command . . . . .	2464
22.5.1	Syntax . . . . .	2464
22.5.2	Examples . . . . .	2464
22.5.3	Description . . . . .	2464
22.5.4	Restrictions . . . . .	2465
22.5.5	Related commands . . . . .	2465
22.5.6	Default . . . . .	2465
22.6	dihedral_style harmonic command . . . . .	2465
22.6.1	Syntax . . . . .	2465
22.6.2	Examples . . . . .	2465
22.6.3	Description . . . . .	2465
22.6.4	Restrictions . . . . .	2466
22.6.5	Related commands . . . . .	2466
22.6.6	Default . . . . .	2466
22.7	dihedral_style helix command . . . . .	2466

22.7.1	Syntax . . . . .	2466
22.7.2	Examples . . . . .	2467
22.7.3	Description . . . . .	2467
22.7.4	Restrictions . . . . .	2467
22.7.5	Related commands . . . . .	2467
22.7.6	Default . . . . .	2468
22.8	dihedral_style hybrid command . . . . .	2468
22.8.1	Syntax . . . . .	2468
22.8.2	Examples . . . . .	2468
22.8.3	Description . . . . .	2468
22.8.4	Restrictions . . . . .	2469
22.8.5	Related commands . . . . .	2469
22.8.6	Default . . . . .	2469
22.9	dihedral_style multi/harmonic command . . . . .	2469
22.9.1	Syntax . . . . .	2469
22.9.2	Examples . . . . .	2469
22.9.3	Description . . . . .	2470
22.9.4	Restrictions . . . . .	2470
22.9.5	Related commands . . . . .	2470
22.9.6	Default . . . . .	2470
22.10	dihedral_style nharmonic command . . . . .	2471
22.10.1	Syntax . . . . .	2471
22.10.2	Examples . . . . .	2471
22.10.3	Description . . . . .	2471
22.10.4	Restrictions . . . . .	2472
22.10.5	Related commands . . . . .	2472
22.10.6	Default . . . . .	2472
22.11	dihedral_style none command . . . . .	2472
22.11.1	Syntax . . . . .	2472
22.11.2	Examples . . . . .	2472
22.11.3	Description . . . . .	2472
22.11.4	Restrictions . . . . .	2472
22.11.5	Related commands . . . . .	2472
22.11.6	Default . . . . .	2473
22.12	dihedral_style opls command . . . . .	2473
22.12.1	Syntax . . . . .	2473
22.12.2	Examples . . . . .	2473
22.12.3	Description . . . . .	2473
22.12.4	Restrictions . . . . .	2474
22.12.5	Related commands . . . . .	2474
22.12.6	Default . . . . .	2474
22.13	dihedral_style quadratic command . . . . .	2474
22.13.1	Syntax . . . . .	2474
22.13.2	Examples . . . . .	2474
22.13.3	Description . . . . .	2474
22.13.4	Restrictions . . . . .	2475
22.13.5	Related commands . . . . .	2475
22.13.6	Default . . . . .	2475
22.14	dihedral_style spherical command . . . . .	2475
22.14.1	Syntax . . . . .	2475
22.14.2	Examples . . . . .	2475
22.14.3	Description . . . . .	2476
22.14.4	Restrictions . . . . .	2477
22.14.5	Related commands . . . . .	2477



22.14.6	Default	2477
22.15	dihedral_style table command	2477
22.16	dihedral_style table/cut command	2477
22.16.1	Syntax	2477
22.16.2	Examples	2478
22.16.3	Description	2478
22.16.4	Restart, fix_modify, output, run start/stop, minimize info	2480
22.16.5	Restrictions	2481
22.16.6	Related commands	2481
22.16.7	Default	2481
22.17	dihedral_style zero command	2481
22.17.1	Syntax	2481
22.17.2	Examples	2481
22.17.3	Description	2481
22.17.4	Restrictions	2482
22.17.5	Related commands	2482
22.17.6	Default	2482
<b>23</b>	<b>Improper Styles</b>	<b>2483</b>
23.1	improper_style harmonic command	2483
23.1.1	Syntax	2483
23.1.2	Examples	2483
23.1.3	Description	2483
23.1.4	Restrictions	2484
23.1.5	Related commands	2484
23.1.6	Default	2484
23.2	improper_style class2 command	2484
23.2.1	Syntax	2484
23.2.2	Examples	2484
23.2.3	Description	2484
23.2.4	Restrictions	2486
23.2.5	Related commands	2486
23.2.6	Default	2486
23.3	improper_style cossq command	2486
23.3.1	Syntax	2486
23.3.2	Examples	2486
23.3.3	Description	2486
23.3.4	Restrictions	2487
23.3.5	Related commands	2487
23.3.6	Default	2487
23.4	improper_style cvff command	2487
23.4.1	Syntax	2487
23.4.2	Examples	2487
23.4.3	Description	2488
23.4.4	Restrictions	2488
23.4.5	Related commands	2488
23.4.6	Default	2489
23.5	improper_style distance command	2489
23.5.1	Syntax	2489
23.5.2	Examples	2489
23.5.3	Description	2489
23.5.4	Restrictions	2490
23.5.5	Related commands	2490
23.5.6	Default	2490

23.6	improper_style distharm command	2490
23.6.1	Syntax	2490
23.6.2	Examples	2490
23.6.3	Description	2490
23.6.4	Restrictions	2491
23.6.5	Related commands	2491
23.6.6	Default	2491
23.7	improper_style fourier command	2491
23.7.1	Syntax	2491
23.7.2	Examples	2491
23.7.3	Description	2491
23.7.4	Restrictions	2493
23.7.5	Related commands	2493
23.7.6	Default	2493
23.8	improper_style harmonic command	2493
23.8.1	Syntax	2493
23.8.2	Examples	2493
23.8.3	Description	2493
23.8.4	Restrictions	2494
23.8.5	Related commands	2494
23.8.6	Default	2494
23.9	improper_style hybrid command	2494
23.9.1	Syntax	2494
23.9.2	Examples	2494
23.9.3	Description	2495
23.9.4	Restrictions	2495
23.9.5	Related commands	2496
23.9.6	Default	2496
23.10	improper_style inversion/harmonic command	2496
23.10.1	Syntax	2496
23.10.2	Examples	2496
23.10.3	Description	2496
23.10.4	Restrictions	2497
23.10.5	Related commands	2497
23.10.6	Default	2497
23.11	improper_style none command	2497
23.11.1	Syntax	2497
23.11.2	Examples	2497
23.11.3	Description	2497
23.11.4	Restrictions	2498
23.11.5	Related commands	2498
23.11.6	Default	2498
23.12	improper_style ring command	2498
23.12.1	Syntax	2498
23.12.2	Examples	2498
23.12.3	Description	2498
23.12.4	Restrictions	2499
23.12.5	Related commands	2499
23.13	improper_style sqdistharm command	2499
23.13.1	Syntax	2499
23.13.2	Examples	2499
23.13.3	Description	2500
23.13.4	Restrictions	2500
23.13.5	Related commands	2500

23.13.6 Default . . . . .	2500
23.14 improper_style umbrella command . . . . .	2500
23.14.1 Syntax . . . . .	2500
23.14.2 Examples . . . . .	2501
23.14.3 Description . . . . .	2501
23.14.4 Restrictions . . . . .	2502
23.14.5 Related commands . . . . .	2502
23.14.6 Default . . . . .	2502
23.15 improper_style zero command . . . . .	2502
23.15.1 Syntax . . . . .	2502
23.15.2 Examples . . . . .	2502
23.15.3 Description . . . . .	2502
23.15.4 Restrictions . . . . .	2503
23.15.5 Related commands . . . . .	2503
23.15.6 Default . . . . .	2503
<b>24 Dump Styles</b>	<b>2505</b>
<b>25 Bibliography</b>	<b>2507</b>
<b>IV Indices and tables</b>	<b>2531</b>
<b>Bibliography</b>	<b>2535</b>
<b>Python Module Index</b>	<b>2537</b>
<b>Index</b>	<b>2539</b>



LAMMPS stands for **L**arge-scale **A**tomic/**M**olecular **M**assively **P**arallel **S**imulator.

LAMMPS is a classical molecular dynamics simulation code with a focus on materials modeling. It was designed to run efficiently on parallel computers. It was developed originally at Sandia National Laboratories, a US Department of Energy facility. The majority of funding for LAMMPS has come from the US Department of Energy (DOE). LAMMPS is an open-source code, distributed freely under the terms of the GNU Public License Version 2 (GPLv2).

The [LAMMPS website](#) has a variety of information about the code. It includes links to an on-line version of this manual, an [online forum](#) where users can post questions and discuss LAMMPS, and a [GitHub site](#) where all LAMMPS development is coordinated.

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The content for this manual is part of the LAMMPS distribution. The online version always corresponds to the latest development version. If needed, you can download or build a local copy of the manual as HTML pages or a PDF file by following the steps on the [Build the LAMMPS documentation](#) page. If you have difficulties viewing the pages please [see this note](#).

---

The manual is organized in three parts: 1) the [User Guide](#) for how to install and use LAMMPS, 2) the [Programmer Guide](#) for how to write programs using the LAMMPS library from different programming languages and how to modify and extend LAMMPS, and 3) the [Command Reference](#) which includes detailed descriptions of all commands included in the LAMMPS code.

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**Part I**

**User Guide**





## INTRODUCTION

These pages provide a brief introduction to LAMMPS.

### 1.1 Overview of LAMMPS

LAMMPS is a classical molecular dynamics (MD) code that models ensembles of particles in a liquid, solid, or gaseous state. It can model atomic, polymeric, biological, solid-state (metals, ceramics, oxides), granular, coarse-grained, or macroscopic systems using a variety of interatomic potentials (force fields) and boundary conditions. It can model 2d or 3d systems with only a few particles up to millions or billions.

LAMMPS can be built and run on a laptop or desktop machine, but is designed for parallel computers. It will run in serial and on any parallel machine that supports the [MPI](#) message-passing library. This includes shared-memory boxes and distributed-memory clusters and supercomputers. Parts of LAMMPS also support [OpenMP multi-threading](#), vectorization and GPU acceleration.

LAMMPS is written in C++ and requires a compiler that is at least compatible with the C++-11 standard. Earlier versions were written in F77, F90, and C++-98. See the [History page](#) of the website for details. All versions can be downloaded as source code from the [LAMMPS website](#).

LAMMPS is designed to be easy to modify or extend with new capabilities, such as new force fields, atom types, boundary conditions, or diagnostics. See the [Modify](#) page for more details.

In the most general sense, LAMMPS integrates Newton's equations of motion for a collection of interacting particles. A single particle can be an atom or molecule or electron, a coarse-grained cluster of atoms, or a mesoscopic or macroscopic clump of material. The interaction models that LAMMPS includes are mostly short-range in nature; some long-range models are included as well.

LAMMPS uses neighbor lists to keep track of nearby particles. The lists are optimized for systems with particles that are repulsive at short distances, so that the local density of particles never becomes too large. This is in contrast to methods used for modeling plasma or gravitational bodies (e.g. galaxy formation).

On parallel machines, LAMMPS uses spatial-decomposition techniques with MPI parallelization to partition the simulation domain into small sub-domains of equal computational cost, one of which is assigned to each processor. Processors communicate and store "ghost" atom information for atoms that border their sub-domain. Multi-threading parallelization and GPU acceleration with with particle-decomposition can be used in addition.

## 1.2 What does a LAMMPS version mean

The LAMMPS “version” is the date when it was released, such as 1 May 2014. LAMMPS is updated continuously and we aim to keep it working correctly and reliably at all times. You can follow its development in a public [git repository on GitHub](#).

Whenever we fix a bug or update or add a feature, it will be merged into the *develop* branch of the git repository. When a sufficient number of changes have accumulated *and* the software passes a set of automated tests, we release it in the next *patch* release, which are made every few weeks. The *release* branch of the git repository is updated with every such release. Info on patch releases are on [this website page](#).

Once or twice a year, we apply only bug fixes and small, non-intrusive changes to the *develop* branch and the code is subjected to more detailed and thorough testing than the default automated testing. The latest patch release after such a period is then also labeled as a *stable* version and the *stable* branch is updated with it. Between stable releases we occasionally release some updates to the stable release containing only bug fixes and updates back-ported from *develop* but no new features and update the *stable* branch accordingly.

Each version of LAMMPS contains all the documented features up to and including its version date.

The version date is printed to the screen and logfile every time you run LAMMPS. It is also in the file `src/version.h` and in the LAMMPS directory name created when you unpack a tarball. And it is on the first page of the [manual](#).

- If you browse the HTML pages on the LAMMPS WWW site, they will by default describe the most current patch release version of LAMMPS. In the navigation bar on the bottom left, there is the option to view instead the documentation for the most recent *stable* version or the latest version from the current development branch.
- If you browse the HTML pages included in your tarball, they describe the version you have, which may be older.

## 1.3 LAMMPS features

LAMMPS is a classical molecular dynamics (MD) code with these general classes of functionality:

1. *General features*
2. *Particle and model types*
3. *Interatomic potentials (force fields)*
4. *Atom creation*
5. *Ensembles, constraints, and boundary conditions*
6. *Integrators*
7. *Diagnostics*
8. *Output*
9. *Multi-replica models*
10. *Pre- and post-processing*
11. *Specialized features (beyond MD itself)*

### 1.3.1 General features

- runs on a single processor or in parallel
- distributed memory message-passing parallelism (MPI)
- shared memory multi-threading parallelism (OpenMP)
- spatial decomposition of simulation domain for MPI parallelism
- particle decomposition inside of spatial decomposition for OpenMP and GPU parallelism
- GPLv2 licensed open-source distribution
- highly portable C++-11
- modular code with most functionality in optional packages
- only depends on MPI library for basic parallel functionality, MPI stub for serial compilation
- other libraries are optional and only required for specific packages
- GPU (CUDA, OpenCL, HIP, SYCL), Intel Xeon Phi, and OpenMP support for many code features
- easy to extend with new features and functionality
- runs from an input script
- syntax for defining and using variables and formulas
- syntax for looping over runs and breaking out of loops
- run one or multiple simulations simultaneously (in parallel) from one script
- build as library, invoke LAMMPS through library interface or provided Python wrapper or SWIG based wrappers
- couple with other codes: LAMMPS calls other code, other code calls LAMMPS, umbrella code calls both

### 1.3.2 Particle and model types

(See *atom style* command)

- atoms
- coarse-grained particles (e.g. bead-spring polymers)
- united-atom polymers or organic molecules
- all-atom polymers, organic molecules, proteins, DNA
- metals
- granular materials
- coarse-grained mesoscale models
- finite-size spherical and ellipsoidal particles
- finite-size line segment (2d) and triangle (3d) particles
- finite-size rounded polygons (2d) and polyhedra (3d) particles
- point dipole particles
- particles with magnetic spin
- rigid collections of n particles

- hybrid combinations of these

### 1.3.3 Interatomic potentials (force fields)

(See *pair style*, *bond style*, *angle style*, *dihedral style*, *improper style*, *kpace style* commands)

- pairwise potentials: Lennard-Jones, Buckingham, Morse, Born-Mayer-Huggins, Yukawa, soft, class 2 (COMPASS), hydrogen bond, tabulated
- charged pairwise potentials: Coulombic, point-dipole
- many-body potentials: EAM, Finnis/Sinclair EAM, modified EAM (MEAM), embedded ion method (EIM), EDIP, ADP, Stillinger-Weber, Tersoff, REBO, AIREBO, ReaxFF, COMB, Streitz-Mintmire, 3-body polymorphic, BOP, Vashishta
- machine learning potentials: SNAP, GAP, ACE, N2P2, RANN, AGNI
- long-range interactions for charge, point-dipoles, and LJ dispersion: Ewald, Wolf, PPPM (similar to particle-mesh Ewald), MSM
- polarization models: *QEq*, *core/shell model*, *Drude dipole model*
- charge equilibration (QEq via dynamic, point, shielded, Slater methods)
- coarse-grained potentials: DPD, GayBerne, REsquared, colloidal, DLVO
- mesoscopic potentials: granular, Peridynamics, SPH, mesoscopic tubular potential (MESONT)
- semi-empirical potentials: multi-ion generalized pseudopotential theory (MGPT), second moment tight binding + QEq (SMTB-Q), density functional tight-binding (LATTE)
- electron force field (eFF, AWPMD)
- bond potentials: harmonic, FENE, Morse, nonlinear, class 2, quartic (breakable), tabulated
- angle potentials: harmonic, CHARMM, cosine, cosine/squared, cosine/periodic, class 2 (COMPASS), tabulated
- dihedral potentials: harmonic, CHARMM, multi-harmonic, helix, class 2 (COMPASS), OPLS, tabulated
- improper potentials: harmonic, cvff, umbrella, class 2 (COMPASS), tabulated
- polymer potentials: all-atom, united-atom, bead-spring, breakable
- water potentials: TIP3P, TIP4P, SPC, SPC/E and variants
- interlayer potentials for graphene and analogues
- metal-organic framework potentials (QuickFF, MO-FF)
- implicit solvent potentials: hydrodynamic lubrication, Debye
- force-field compatibility with common CHARMM, AMBER, DREIDING, OPLS, GROMACS, COMPASS options
- access to the [OpenKIM Repository](#) of potentials via *kim command*
- hybrid potentials: multiple pair, bond, angle, dihedral, improper potentials can be used in one simulation
- overlaid potentials: superposition of multiple pair potentials (including many-body) with optional scale factor

### 1.3.4 Atom creation

(See *read\_data*, *lattice*, *create\_atoms*, *delete\_atoms*, *displace\_atoms*, *replicate* commands)

- read in atom coords from files
- create atoms on one or more lattices (e.g. grain boundaries)
- delete geometric or logical groups of atoms (e.g. voids)
- replicate existing atoms multiple times
- displace atoms

### 1.3.5 Ensembles, constraints, and boundary conditions

(See *fix* command)

- 2d or 3d systems
- orthogonal or non-orthogonal (triclinic symmetry) simulation domains
- constant NVE, NVT, NPT, NPH, Parrinello/Rahman integrators
- thermostatting options for groups and geometric regions of atoms
- pressure control via Nose/Hoover or Berendsen barostatting in 1 to 3 dimensions
- simulation box deformation (tensile and shear)
- harmonic (umbrella) constraint forces
- rigid body constraints
- SHAKE bond and angle constraints
- motion constraints to manifold surfaces
- Monte Carlo bond breaking, formation, swapping, template based reaction modeling
- atom/molecule insertion and deletion
- walls of various kinds, static and moving
- non-equilibrium molecular dynamics (NEMD)
- variety of additional boundary conditions and constraints

### 1.3.6 Integrators

(See *run*, *run\_style*, *minimize* commands)

- velocity-Verlet integrator
- Brownian dynamics
- rigid body integration
- energy minimization via conjugate gradient or steepest descent relaxation
- rRESPA hierarchical timestepping
- rerun command for post-processing of dump files

### 1.3.7 Diagnostics

- see various flavors of the *fix* and *compute* commands
- introspection command for system, simulation, and compile time settings and configurations

### 1.3.8 Output

(*dump*, *restart* commands)

- log file of thermodynamic info
- text dump files of atom coords, velocities, other per-atom quantities
- binary restart files
- parallel I/O of dump and restart files
- per-atom quantities (energy, stress, centro-symmetry parameter, CNA, etc)
- user-defined system-wide (log file) or per-atom (dump file) calculations
- custom partitioning (chunks) for binning, and static or dynamic grouping of atoms for analysis
- spatial, time, and per-chunk averaging of per-atom quantities
- time averaging and histogramming of system-wide quantities
- atom snapshots in native, XYZ, XTC, DCD, CFG formats

### 1.3.9 Multi-replica models

- *nudged elastic band*
- *hyperdynamics*
- *parallel replica dynamics*
- *temperature accelerated dynamics*
- *parallel tempering*
- path-integral MD: *first variant* `<fix_pimd>`, *second variant* `<fix_ipi>`
- multi-walker collective variables with *Colvars* and *Plumed*

### 1.3.10 Pre- and post-processing

- A handful of pre- and post-processing tools are packaged with LAMMPS, some of which can convert input and output files to/from formats used by other codes; see the *Tools* page.
- Our group has also written and released a separate toolkit called *Pizza.py* which provides tools for doing setup, analysis, plotting, and visualization for LAMMPS simulations. *Pizza.py* is written in *Python* and is available for download from the *Pizza.py* [WWW site](#).

### 1.3.11 Specialized features

LAMMPS can be built with optional packages which implement a variety of additional capabilities. See the *Optional Packages* page for details.

These are LAMMPS capabilities which you may not think of as typical classical MD options:

- *static and dynamic load-balancing*, optional with recursive bisectioning decomposition
- *generalized aspherical particles*
- *stochastic rotation dynamics (SRD)*
- *real-time visualization and interactive MD, built-in renderer for images and movies*
- calculate *virtual diffraction patterns*
- calculate *finite temperature phonon dispersion* and the *dynamical matrix of minimized structures*
- *atom-to-continuum coupling* with finite elements
- coupled rigid body integration via the *POEMS* library
- *QM/MM coupling*
- Monte Carlo via *GCMC* and *tfMC* and *atom swapping*
- *path-integral molecular dynamics (PIMD)* and *this as well*
- *Direct Simulation Monte Carlo* for low-density fluids
- *Peridynamics modeling*
- *Lattice Boltzmann fluid*
- *targeted* and *steered* molecular dynamics
- *two-temperature electron model*

## 1.4 LAMMPS non-features

LAMMPS is designed to be a fast, parallel engine for molecular dynamics (MD) simulations. It provides only a modest amount of functionality for setting up simulations and analyzing their output.

Specifically, LAMMPS was not conceived and designed for:

- being run through a GUI
- building molecular systems, or building molecular topologies
- assign force-field coefficients automatically
- perform sophisticated analysis of your MD simulation
- visualize your MD simulation interactively
- plot your output data

Over the years some of these limitations have been reduced or removed, through features added to LAMMPS or external tools that either closely interface with LAMMPS or extend LAMMPS.

Here are suggestions on how to perform these tasks:

- **GUI:** LAMMPS can be built as a library and a Python wrapper that wraps the library interface is provided. Thus, GUI interfaces can be written in Python (or C or C++ if desired) that run LAMMPS and visualize or plot its output. Examples of this are provided in the `python` directory and described on the [Python](#) doc page. Also, there are several external wrappers or GUI front ends.
- **Builder:** Several pre-processing tools are packaged with LAMMPS. Some of them convert input files in formats produced by other MD codes such as CHARMM, AMBER, or Insight into LAMMPS input formats. Some of them are simple programs that will build simple molecular systems, such as linear bead-spring polymer chains. The `moltemplate` program is a true molecular builder that will generate complex molecular models. See the [Tools](#) page for details on tools packaged with LAMMPS. The [Pre/post processing page](#) of the LAMMPS website describes a variety of third party tools for this task. Furthermore, some LAMMPS internal commands allow to reconstruct, or selectively add topology information, as well as provide the option to insert molecule templates instead of atoms for building bulk molecular systems.
- **Force-field assignment:** The conversion tools described in the previous bullet for CHARMM, AMBER, and Insight will also assign force field coefficients in the LAMMPS format, assuming you provide CHARMM, AMBER, or BIOVIA (formerly Accelrys) force field files. The tools [ParmEd](#) and [InterMol](#) are particularly powerful and flexible in converting force field and topology data between various MD simulation programs.
- **Simulation analysis:** If you want to perform analysis on-the-fly as your simulation runs, see the [compute](#) and [fix](#) doc pages, which list commands that can be used in a LAMMPS input script. Also see the [Modify](#) page for info on how to add your own analysis code or algorithms to LAMMPS. For post-processing, LAMMPS output such as [dump file snapshots](#) can be converted into formats used by other MD or post-processing codes. To some degree, that conversion can be done directly inside of LAMMPS by interfacing to the VMD molfile plugins. The [rerun](#) command also allows to do some post-processing of existing trajectories, and through being able to read a variety of file formats, this can also be used for analyzing trajectories from other MD codes. Some post-processing tools packaged with LAMMPS will do these conversions. Scripts provided in the `tools/python` directory can extract and massage data in dump files to make it easier to import into other programs. See the [Tools](#) page for details on these various options.
- **Visualization:** LAMMPS can produce NETPBM, JPG or PNG snapshot images on-the-fly via its [dump image](#) command and pass them to an external program, [FFmpeg](#) to generate movies from them. For high-quality, interactive visualization there are many excellent and free tools available. See the [Visualization Tools](#) page of the LAMMPS website for visualization packages that can process LAMMPS output data.
- **Plotting:** See the next bullet about [Pizza.py](#) as well as the [Python](#) page for examples of plotting LAMMPS output. Scripts provided with the `python` tool in the `tools` directory will extract and massage data in log and dump files to make it easier to analyze and plot. See the [Tools](#) doc page for more discussion of the various tools.
- **Pizza.py:** Our group has also written a separate toolkit called [Pizza.py](#) which can do certain kinds of setup, analysis, plotting, and visualization (via OpenGL) for LAMMPS simulations. It thus provides some functionality for several of the above bullets. [Pizza.py](#) is written in [Python](#) and is available for download from [this page](#).

## 1.5 LAMMPS open-source license

### 1.5.1 GPL version of LAMMPS

LAMMPS is an open-source code, available free-of-charge, and distributed under the terms of the [GNU Public License Version 2 \(GPLv2\)](#), which means you can use or modify the code however you wish for your own purposes, but have to adhere to certain rules when redistributing it - specifically in binary form - or are distributing software derived from it or that includes parts of it.

LAMMPS comes with no warranty of any kind.

As each source file states in its header, it is a copyrighted code, and thus not in the public domain. For more information about open-source software and open-source distribution, see [www.gnu.org](http://www.gnu.org) or [www.opensource.org](http://www.opensource.org). The legal text of



the GPL as it applies to LAMMPS is in the LICENSE file included in the LAMMPS distribution.

Here is a more specific summary of what the GPL means for LAMMPS users:

- (1) Anyone is free to use, copy, modify, or extend LAMMPS in any way they choose, including for commercial purposes.
- (2) If you **distribute** a modified version of LAMMPS, it must remain open-source, meaning you are required to distribute **all** of it under the terms of the GPL. You should clearly annotate such a modified code as a derivative version of LAMMPS.
- (3) If you release any code that includes or uses LAMMPS source code, then it must also be open-sourced, meaning you distribute it under the terms of the GPL. You may write code that interfaces LAMMPS to a differently licensed library. In that case the code that provides the interface must be licensed GPL, but not necessarily that library unless you are distributing binaries that require the library to run.
- (4) If you give LAMMPS files to someone else, the GPL LICENSE file and source file headers (including the copyright and GPL notices) should remain part of the code.

## 1.5.2 LGPL version of LAMMPS

We occasionally make stable LAMMPS releases available under the [GNU Lesser Public License v2.1](#). This is on request only and with non-LGPL compliant files removed. This allows uses linking non-GPL compatible software with the (otherwise unmodified) LAMMPS library or loading it dynamically at runtime. Any **modifications** to the LAMMPS code however, even with the LGPL licensed version, must still be made available under the same open source terms as LAMMPS itself.

## 1.6 Authors of LAMMPS

The primary LAMMPS developers are at Sandia National Labs and Temple University:

- [Steve Plimpton](#), sjplimp at sandia.gov
- Aidan Thompson, athomps at sandia.gov
- Stan Moore, stamoor at sandia.gov
- Axel Kohlmeyer, akohlmey at gmail.com
- Richard Berger, richard.berger at outlook.com

Past developers include Paul Crozier and Mark Stevens, both at Sandia, and Ray Shan, now at Materials Design.

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The [Authors](#) page of the [LAMMPS website](#) has a comprehensive list of all the individuals who have contributed code for a new feature or command or tool to LAMMPS.

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The following folks deserve special recognition. Many of the packages they have written are unique for an MD code and LAMMPS would not be as general-purpose as it is without their expertise and efforts.

- Metin Aktulga (MSU), REAXFF package for C/C++ version of ReaxFF
- Mike Brown (Intel), GPU and INTEL packages
- Colin Denniston (U Western Ontario), LATBOLTZ package
- Georg Ganzenmuller (EMI), MACHDYN and SPH packages
- Andres Jaramillo-Botero (Caltech), EFF package for electron force field

- Reese Jones (Sandia) and colleagues, ATC package for atom/continuum coupling
- Christoph Kloss (DCS Computing), LIGGGHTS code for granular materials, built on top of LAMMPS
- Rudra Mukherjee (JPL), POEMS package for articulated rigid body motion
- Trung Ngyuen (Northwestern U), GPU, RIGID, BODY, and DIELECTRIC packages
- Mike Parks (Sandia), PERI package for Peridynamics
- Roy Pollock (LLNL), Ewald and PPPM solvers
- Julien Tranchida (Sandia), SPIN package
- Christian Trott (Sandia), CUDA and KOKKOS packages
- Ilya Valuev (JIHT), AWPMD package for wave packet MD
- Greg Wagner (Northwestern U), MEAM package for MEAM potential

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As discussed on the [History page](#) of the website, LAMMPS originated as a cooperative project between DOE labs and industrial partners. Folks involved in the design and testing of the original version of LAMMPS were the following:

- John Carpenter (Mayo Clinic, formerly at Cray Research)
- Terry Stouch (Lexicon Pharmaceuticals, formerly at Bristol Myers Squibb)
- Steve Lustig (Dupont)
- Jim Belak and Roy Pollock (LLNL)

## 1.7 Citing LAMMPS

### 1.7.1 Core Algorithms

The paper mentioned below is the best overview of LAMMPS, but there are also publications describing particular models or algorithms implemented in LAMMPS or complementary software that it has interfaces to. Please see below for how to cite contributions to LAMMPS.

The latest canonical publication that describes the basic features, the source code design, the program structure, the spatial decomposition approach, the neighbor finding, basic communications algorithms, and how users and developers have contributed to LAMMPS is:

LAMMPS - A flexible simulation tool for particle-based materials modeling at the atomic, meso, and continuum scales, *Comp. Phys. Comm.* 271, 108171 (2022)

So a project using LAMMPS or a derivative application that uses LAMMPS as a simulation engine should cite this paper. The paper is expected to be published in its final form under the same DOI in the first half of 2022. Please also give the URL of the LAMMPS website in your paper, namely <https://www.lammps.org>.

The original publication describing the parallel algorithms used in the initial versions of LAMMPS is:

S. Plimpton, Fast Parallel Algorithms for Short-Range Molecular Dynamics, *J Comp Phys*, 117, 1-19 (1995).

## 1.7.2 DOI for the LAMMPS source code

The LAMMPS developers use the [Zenodo service at CERN](#) to create digital object identifiers (DOI) for stable releases of the LAMMPS source code. There are two types of DOIs for the LAMMPS source code.

The canonical DOI for **all** versions of LAMMPS, which will always point to the **latest** stable release version is:

- DOI: [10.5281/zenodo.3726416](https://doi.org/10.5281/zenodo.3726416)

In addition there are DOIs for individual stable releases. Currently there are:

- 3 March 2020 version: DOI:[10.5281/zenodo.3726417](https://doi.org/10.5281/zenodo.3726417)
- 29 October 2020 version: DOI:[10.5281/zenodo.4157471](https://doi.org/10.5281/zenodo.4157471)
- 29 September 2021 version: DOI:[10.5281/zenodo.6386596](https://doi.org/10.5281/zenodo.6386596)

## 1.7.3 Home page

The LAMMPS website at <https://www.lammps.org/> is the canonical location for information about LAMMPS and its features.

## 1.7.4 Citing contributions

LAMMPS has many features that use either previously published methods and algorithms or novel features. It also includes potential parameter files for specific models. Where available, a reminder about references for optional features used in a specific run is printed to the screen and log file. Style and output location can be selected with the *-cite command-line switch*. Additional references are given in the documentation of the *corresponding commands* or in the *Howto tutorials*. So please make certain, that you provide the proper acknowledgments and citations in any published works using LAMMPS.

## 1.8 Additional website links

The [LAMMPS website](#) has a variety of additional info about LAMMPS, beyond what is in this manual. Some other useful resources available online are listed below.

- [Brief intro and recently added significant features](#)
- [List of features](#)
- [List of non-features](#)
- [Recent bug fixes and new features](#)
- [Download info](#)
- [GitHub site](#)
- [SourceForge site](#)
- [LAMMPS open-source license](#)
- [Glossary of terms relevant to LAMMPS](#)
- [LAMMPS highlights with images](#)
- [LAMMPS highlights with movies](#)
- [LAMMPS forum](#)

- Workshops
- Tutorials
- Pre- and post-processing tools for LAMMPS
- Other software usable with LAMMPS
- Viz tools usable with LAMMPS
- Benchmark performance
- Publications that have cited LAMMPS
- Authors of LAMMPS
- History of LAMMPS development
- Funding for LAMMPS

## INSTALL LAMMPS

You can download LAMMPS as an executable or as source code.

When downloading the LAMMPS source code, you also have to *build LAMMPS*. But you have more flexibility as to what features to include or exclude in the build. When you download and install pre-compiled LAMMPS executables, you are limited to install which version of LAMMPS is available and which features are included of these builds. If you plan to *modify or extend LAMMPS*, then you **must** build LAMMPS from the source code.

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**Note:** If you have questions about the pre-compiled LAMMPS executables, you need to contact the people preparing those executables. The LAMMPS developers have no control over their choices of how they configure and build their packages and when they update them.

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### 2.1 Download an executable for Linux

Binaries are available for different versions of Linux:

- *Pre-built Ubuntu Linux executables*
- *Pre-built Fedora Linux executables*
- *Pre-built EPEL Linux executables (RHEL, CentOS)*
- *Pre-built OpenSuse Linux executables*
- *Gentoo Linux executable*
- *Arch Linux build-script*

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**Note:** If you have questions about these pre-compiled LAMMPS executables, you need to contact the people preparing those packages. The LAMMPS developers have no control over their choices of how they configure and build their packages and when they update them.

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### 2.1.1 Pre-built Ubuntu Linux executables

A pre-built LAMMPS executable suitable for running on the latest Ubuntu Linux versions, can be downloaded as a Debian package. This allows you to install LAMMPS with a single command, and stay (mostly) up-to-date with the current stable version of LAMMPS by simply updating your operating system.

To install LAMMPS do the following once:

```
$ sudo apt-get install lammeps
```

This downloads an executable named `lmp` to your box and multiple packages with supporting data, examples and libraries as well as any missing dependencies. This executable can then be used in the usual way to run input scripts:

```
$ lmp -in in.lj
```

To update LAMMPS to the latest packaged version, do the following:

```
$ sudo apt-get update
```

which will also update other packages on your system.

The `lmp` binary is built with the *KIM package* included, which results in the above command also installing the `kim-api` binaries when LAMMPS is installed. In order to use potentials from [openkim.org](http://openkim.org), you can also install the `openkim-models` package

```
$ sudo apt-get install openkim-models
```

Or use the KIM-API commands to download and install individual models. To un-install LAMMPS, do the following:

```
$ sudo apt-get remove lammeps
```

Please use `lmp -help` to see which compilation options, packages, and styles are included in the binary.

Thanks to Anton Gladky ([gladky.anton@gmail.com](mailto:gladky.anton@gmail.com)) for setting up this Ubuntu package capability.

---

### 2.1.2 Pre-built Fedora Linux executables

Pre-built LAMMPS packages for stable releases are available in the Fedora Linux distribution as of Fedora version 28. The packages can be installed via the `dnf` package manager. There are 3 basic varieties (`lammeps` = no MPI, `lammeps-mpich` = MPICH MPI library, `lammeps-openmpi` = OpenMPI MPI library) and for each support for linking to the C library interface (`lammeps-devel`, `lammeps-mpich-devel`, `lammeps-openmpi-devel`), the header for compiling programs using the C library interface (`lammeps-headers`), and the LAMMPS python module for Python 3. All packages can be installed at the same time and the name of the LAMMPS executable is `lmp` and `lmp_openmpi` or `lmp_mpich` respectively. By default, `lmp` will refer to the serial executable, unless one of the MPI environment modules is loaded (`module load mpi/mpich-x86_64` or `module load mpi/openmpi-x86_64`). Then the corresponding parallel LAMMPS executable can be used. The same mechanism applies when loading the LAMMPS python module.

To install LAMMPS with OpenMPI and run an input `in.lj` with 2 CPUs do:

```
$ dnf install lammeps-openmpi
$ module load mpi/openmpi-x86_64
$ mpirun -np 2 lmp -in in.lj
```

The `dnf install` command is needed only once. In case of a new LAMMPS stable release, `dnf update` will automatically update to the newer version as soon as the RPM files are built and uploaded to the download mirrors. The `module load` command is needed once per (shell) session or shell terminal instance, unless it is automatically loaded from the shell profile.

The LAMMPS binary is built with the *KIM package* which results in the above command also installing the *kim-api* binaries when LAMMPS is installed. In order to use potentials from [openkim.org](http://openkim.org), you can install the *openkim-models* package

```
$ dnf install openkim-models
```

Please use `lmp -help` to see which compilation options, packages, and styles are included in the binary.

Thanks to Christoph Junghans (LANL) for making LAMMPS available in Fedora.

---

### 2.1.3 Pre-built EPEL Linux executable

Pre-built LAMMPS (and KIM) packages for stable releases are available in the [Extra Packages for Enterprise Linux \(EPEL\) repository](#) for use with Red Hat Enterprise Linux (RHEL) or CentOS version 7.x and compatible Linux distributions. Names of packages, executable, and content are the same as described above for Fedora Linux. But RHEL/CentOS 7.x uses the `yum` package manager instead of `dnf` in Fedora 28.

Please use `lmp -help` to see which compilation options, packages, and styles are included in the binary.

Thanks to Christoph Junghans (LANL) for making LAMMPS available in EPEL.

---

### 2.1.4 Pre-built OpenSuse Linux executable

A pre-built LAMMPS package for stable releases is available in OpenSuse as of Leap 15.0. You can install the package with:

```
$ zypper install lammps
```

This includes support for OpenMPI. The name of the LAMMPS executable is `lmp`. Thus to run an input in parallel on 2 CPUs you would do:

```
$ mpirun -np 2 lmp -in in.lj
```

Please use `lmp -help` to see which compilation options, packages, and styles are included in the binary.

The LAMMPS binary is built with the *KIM package* which results in the above command also installing the *kim-api* binaries when LAMMPS is installed. In order to use potentials from [openkim.org](http://openkim.org), you can install the *openkim-models* package

```
$ zypper install openkim-models
```

Thanks to Christoph Junghans (LANL) for making LAMMPS available in OpenSuse.

---

## 2.1.5 Gentoo Linux executable

LAMMPS is part of Gentoo's main package tree and can be installed by typing:

```
% emerge --ask lammmps
```

Note that in Gentoo the LAMMPS source is downloaded and the package is built on the your machine.

Certain LAMMPS packages can be enable via USE flags, type

```
% equery uses lammmps
```

for details.

Thanks to Nicolas Bock and Christoph Junghans (LANL) for setting up this Gentoo capability.

---

## 2.1.6 Archlinux build-script

LAMMPS is available via Arch's unofficial Arch User repository (AUR). There are three scripts available, named lammmps, lammmps-beta and lammmps-git. They respectively package the stable, patch and git releases.

To install, you will need to have the git package installed. You may use any of the above names in-place of lammmps.

```
$ git clone https://aur.archlinux.org/lammmps.git
$ cd lammmps
$ makepkg -s
$ makepkg -i
```

To update, you may repeat the above, or change into the cloned directory, and execute the following, after which, if there are any changes, you may use makepkg as above.

```
$ git pull
```

Alternatively, you may use an AUR helper to install these packages.

Note that the AUR provides build-scripts that download the source and the build the package on your machine.

---

**Note:** It looks like the Arch Linux AUR repository build scripts for LAMMPS have not been updated since the 29 October 2020 version. You may want to consider installing a more current version of LAMMPS from source directly.

---

## 2.2 Download an executable for Mac

LAMMPS can be downloaded, built, and configured for OS X on a Mac with [Homebrew](#). (Alternatively, see the install instructions for *Download an executable via Conda*.) The following LAMMPS packages are unavailable at this time because of additional needs not yet met: GPU, KOKKOS, LATTE, MSCG, MPIIO, POEMS, VORONOI.

After installing Homebrew, you can install LAMMPS on your system with the following commands:

```
% brew install lammmps
```



This will install the executables “lammmps\_serial” and “lammmps\_mpi”, as well as the LAMMPS “doc”, “potentials”, “tools”, “bench”, and “examples” directories.

Once LAMMPS is installed, you can test the installation with the Lennard-Jones benchmark file:

```
% brew test lammmps -v
```

The LAMMPS binary is built with the *KIM package* which results in Homebrew also installing the *kim-api* binaries when LAMMPS is installed. In order to use potentials from [openkim.org](http://openkim.org), you can install the *openkim-models* package

```
% brew install openkim-models
```

If you have problems with the installation you can post issues to [this link](#).

Thanks to Derek Thomas ([derekt at cello.t.u-tokyo.ac.jp](mailto:derekt@cello.t.u-tokyo.ac.jp)) for setting up the Homebrew capability.

## 2.3 Download an executable for Windows

Pre-compiled Windows installers which install LAMMPS executables on a Windows system can be downloaded from this site:

<http://packages.lammps.org/windows.html>

Note that each installer package has a date in its name, which corresponds to the LAMMPS version of the same date. Installers for current and older versions of LAMMPS are available. 32-bit and 64-bit installers are available, and each installer contains both a serial and parallel executable. The installer website also explains how to install the Windows MPI package (MPICH2 from Argonne National Labs), needed to run in parallel with MPI.

The LAMMPS binaries contain *all optional packages* included in the source distribution except: KIM, KOKKOS, MSCG, PYTHON, ADIOS, H5MD, NETCDF, QMMM, ML-QUIP, and VTK. The serial version also does not include the MPIIO and LATBOLTZ packages. The GPU package is compiled for OpenCL with mixed precision kernels.

The LAMMPS library is compiled as a shared library and the *LAMMPS Python module* is installed, so that it is possible to load LAMMPS into a Python interpreter.

The installer site also has instructions on how to run LAMMPS under Windows, once it is installed, in both serial and parallel.

When you download the installer package, you run it on your Windows machine. It will then prompt you with a dialog, where you can choose the installation directory, unpack and copy several executables, potential files, documentation pdfs, selected example files, etc. It will then update a few system settings (e.g. PATH, LAMMPS\_POTENTIALS) and add an entry into the Start Menu (with references to the documentation, LAMMPS homepage and more). From that menu, there is also a link to an uninstaller that removes the files and undoes the environment manipulations.

Note that to update to a newer version of LAMMPS, you should typically uninstall the version you currently have, download a new installer, and go through the install procedure described above. I.e. the same procedure for installing/updating most Windows programs. You can install multiple versions of LAMMPS (in different directories), but only the executable for the last-installed package will be found automatically, so this should only be done for debugging purposes.

## 2.4 Download an executable for Linux or Mac via Conda

Binaries are available for MacOS or Linux via [Conda](#).

First, one must setup the Conda package manager on your system. Follow the instructions to install [Miniconda](#), then create a conda environment (named *my-lammps-env* or whatever you prefer) for your lammps install:

```
% conda config --add channels conda-forge
% conda create -n my-lammps-env
```

Then, you can install lammps on your system with the following command:

```
% conda activate my-lammps-env
% conda install lammps
```

The LAMMPS binary is built with the *KIM package* which results in Conda also installing the *kim-api* binaries when LAMMPS is installed. In order to use potentials from [openkim.org](http://openkim.org), you can install the *openkim-models* package

```
% conda install openkim-models
```

If you have problems with the installation you can post issues to [this link](#). Thanks to Jan Janssen (Max-Planck-Institut fuer Eisenforschung) for setting up the Conda capability.

---

**Note:** If you have questions about these pre-compiled LAMMPS executables, you need to contact the people preparing those packages. The LAMMPS developers have no control over their choices of how they configure and build their packages and when they update them.

---

## 2.5 Download source and documentation as a tarball

You can download a current LAMMPS tarball from the [download page](#) of the [LAMMPS website](#).

You have two choices of tarballs, either the most recent stable release or the most current patch release. Stable releases occur a few times per year, and undergo more testing before release. Patch releases occur a couple times per month. The new contents in all releases are listed on the [bug and feature page](#) of the website.

Both tarballs include LAMMPS documentation (HTML and PDF files) corresponding to that version. The download page also has an option to download the current-version LAMMPS documentation by itself.

Older versions of LAMMPS can also be downloaded from [this page](#).

Once you have a tarball, unzip and untar it with the following command:

```
$ tar -xzvf lammps*.tar.gz
```

This will create a LAMMPS directory with the version date in its name, e.g. lammps-23Jun18.

---

You can also download a compressed tar or zip archives from the “Assets” sections of the [LAMMPS GitHub releases site](#). The file name will be lammps-<version>.zip which can be unzipped with the following command, to create a lammps-<version> dir:

```
$ unzip lammps*.zip
```

This version corresponds to the selected LAMMPS patch or stable release.

## 2.6 Download the LAMMPS source with git

All LAMMPS development is coordinated through the “LAMMPS GitHub site”. If you clone the LAMMPS repository onto your local machine, it has several advantages:

- You can stay current with changes to LAMMPS with a single git command.
- You can create your own development branches to add code to LAMMPS.
- You can submit your new features back to GitHub for inclusion in LAMMPS. For that you should first create your own *fork on GitHub*.

You must have `git` installed on your system to use the commands explained below to communicate with the git servers on GitHub. For people still using subversion (svn), GitHub also provides [limited support for subversion clients](#).

---

**Note:** As of October 2016, the official home of public LAMMPS development is on GitHub. The previously advertised LAMMPS git repositories on `git.lammps.org` and `bitbucket.org` are now offline or deprecated.

---

You can follow the LAMMPS development on 3 different git branches:

- **stable** : this branch is updated from the *release* branch with every stable release version and also has selected bug fixes and updates back-ported from the *develop* branch
- **release** : this branch is updated with every patch release; updates are always “fast forward” merges from *develop*
- **develop** : this branch follows the ongoing development and is updated with every merge commit of a pull request

To access the git repositories on your box, use the clone command to create a local copy of the LAMMPS repository with a command like:

```
$ git clone -b release https://github.com/lammps/lammps.git mylammps
```

where “mylammps” is the name of the directory you wish to create on your machine and “release” is one of the 3 branches listed above. (Note that you actually download all 3 branches; you can switch between them at any time using “git checkout <branch name>”.)

---

### Saving time and disk space when using git clone

The complete git history of the LAMMPS project is quite large because it contains the entire commit history of the project since fall 2006, which includes the time when LAMMPS was managed with subversion. This includes a few commits that have added and removed some large files (mostly by accident). If you do not need access to the entire commit history (most people don’t), you can speed up the “cloning” process and reduce local disk space requirements by using the `-depth` git command line flag. That will create a “shallow clone” of the repository containing only a subset of the git history. Using a depth of 1000 is usually sufficient to include the head commits of the *develop* and the *release* branches. To include the head commit of the *stable* branch you may need a depth of up to 10000. If you later need more of the git history, you can always convert the shallow clone into a “full clone”.

---

Once the command completes, your directory will contain the same files as if you unpacked a current LAMMPS tarball, with the exception, that the HTML documentation files are not included. They can be fetched from the LAMMPS website by typing `make fetch` in the doc directory. Or they can be generated from the content provided in `doc/src` by typing `make html` from the doc directory.

After initial cloning, as bug fixes and new features are added to LAMMPS you can stay up-to-date by typing the following git commands from within the “mylammps” directory:

```
$ git checkout release      # not needed if you always stay in this branch
$ git checkout stable      # use one of these 3 checkout commands
$ git checkout develop     # to choose the branch to follow
$ git pull
```

Doing a “pull” will not change any files you have added to the LAMMPS directory structure. It will also not change any existing LAMMPS files you have edited, unless those files have changed in the repository. In that case, git will attempt to merge the new repository file with your version of the file and tell you if there are any conflicts. See the git documentation for details.

If you want to access a particular previous release version of LAMMPS, you can instead “check out” any version with a published tag. See the output of `git tag -l` for the list of tags. The git command to do this is as follows.

```
$ git checkout tagID
```

Stable versions and what tagID to use for a particular stable version are discussed on [this page](#). Note that this command will print some warnings, because in order to get back to the latest revision and to be able to update with `git pull` again, you will need to do `git checkout release` (or check out any other desired branch) first.

Once you have updated your local files with a `git pull` (or `git checkout`), you still need to re-build LAMMPS if any source files have changed. How to do this depends on the build system you are using.

---

### CMake build

Change to your build folder and type:

```
cmake . --build
```

CMake should auto-detect whether it needs to re-run the CMake configuration step and otherwise redo the build for all files that have been changed or files that depend on changed files. In case some build options have been changed or renamed, you may have to update those by running:

```
cmake .
```

and then rebuild.

---

### Traditional make

Switch to the src directory and type:

```
$ make purge      # remove any deprecated src files
$ make package-update # sync package files with src files
$ make foo        # re-build for your machine (mpi, serial, etc)
```

to enforce consistency of the source between the src folder and package directories. This is OK to do even if you don’t use any packages. The “make purge” command removes any deprecated src files if they were removed by the patch from a package sub-directory.

**Warning:** If you wish to edit/change a src file that is from a package, you should edit the version of the file inside the package sub-directory with src, then re-install the package. The version in the source directory is merely a copy and will be wiped out if you type “make package-update”.

---

## Git protocols

The servers at github.com support the “https://” access protocol for anonymous, read-only access. If you have a suitably configured GitHub account, you may also use SSH protocol with the URL “git@github.com:lammmps/lammmps.git”.

---

The LAMMPS GitHub project is currently managed by Axel Kohlmeyer (Temple U, akohlmey at gmail.com).

These are the files and sub-directories in the LAMMPS distribution:

README	Short description of the LAMMPS package
LICENSE	GNU General Public License (GPL)
SECURITY.md	Security Policy for the LAMMPS package
bench	benchmark problems
cmake	CMake build files
doc	documentation
examples	simple test problems
fortran	Fortran wrapper for LAMMPS
lib	additional provided or external libraries
potentials	interatomic potential files
python	Python wrappers for LAMMPS
src	source files
tools	pre- and post-processing tools
unittest	sources and inputs for testing LAMMPS

You will have all of these if you download source. You will only have some of them if you download executables, as explained on the pages listed above.



## BUILD LAMMPS

LAMMPS is built as a library and an executable from source code using either traditional makefiles for use with GNU make (which may require manual editing), or using a build environment generated by CMake (Unix Makefiles, Ninja, Xcode, Visual Studio, KDevelop, CodeBlocks and more).

As an alternative you can download a package with pre-built executables or automated build trees as described on the [Install](#) page.

### 3.1 Build LAMMPS with CMake

This page describes how to use CMake in general to build LAMMPS. Details for specific compile time settings and options to enable and configure add-on packages are discussed with those packages. Links to those pages on the [Build overview](#) page.

The following text assumes some familiarity with CMake and focuses on using the command line tool `cmake` and what settings are supported for building LAMMPS. A more detailed tutorial on how to use `cmake` itself, the text mode or graphical user interface, change the generated output files for different build tools and development environments is on a [separate page](#).

---

**Note:** LAMMPS currently requires that CMake version 3.10 or later is available; version 3.12 or later is preferred.

---

**Warning:** You must not mix the *traditional make based* LAMMPS build procedure with using CMake. Thus no packages may be installed or a build been previously attempted in the LAMMPS source directory by using `make <machine>`. CMake will detect if this is the case and generate an error. To remove conflicting files from the `src` you can use the command `make no-all purge` which will un-install all packages and delete all auto-generated files.

#### 3.1.1 Advantages of using CMake

CMake is an alternative to compiling LAMMPS in the traditional way through (*manually customized*) *makefiles* and a recent addition to LAMMPS thanks to the efforts of Christoph Junghans (LANL) and Richard Berger (Temple U). Using CMake has multiple advantages that are specifically helpful for people with limited experience in compiling software or for people that want to modify or extend LAMMPS.

- CMake can detect available hardware, tools, features, and libraries and adapt the LAMMPS default build configuration accordingly.
- CMake can generate files for different build tools and integrated development environments (IDE).

- CMake supports customization of settings with a text mode or graphical user interface. No knowledge of file formats or and complex command line syntax required.
- All enabled components are compiled in a single build operation.
- Automated dependency tracking for all files and configuration options.
- Support for true out-of-source compilation. Multiple configurations and settings with different choices of LAMMPS packages, settings, or compilers can be configured and built concurrently from the same source tree.
- Simplified packaging of LAMMPS for Linux distributions, environment modules, or automated build tools like [Homebrew](#).
- Integration of automated regression testing (the LAMMPS side for that is still under development).

### 3.1.2 Getting started

Building LAMMPS with CMake is a two-step process. First you use CMake to generate a build environment in a new directory. For that purpose you can use either the command-line utility `cmake` (or `cmake3`), the text-mode UI utility `ccmake` (or `ccmake3`) or the graphical utility `cmake-gui`, or use them interchangeably. The second step is then the compilation and linking of all objects, libraries, and executables. Here is a minimal example using the command line version of CMake to build LAMMPS with no add-on packages enabled and no customization:

```
cd lammps           # change to the LAMMPS distribution directory
mkdir build; cd build # create and use a build directory
cmake ../cmake      # configuration reading CMake scripts from ../cmake
cmake --build .     # compilation (or type "make")
```

This will create and change into a folder called `build`, then run the configuration step to generate build files for the default build command and then launch that build command to compile LAMMPS. During the configuration step CMake will try to detect whether support for MPI, OpenMP, FFTW, gzip, JPEG, PNG, and ffmpeg are available and enable the corresponding configuration settings. The progress of this configuration can be followed on the screen and a summary of selected options and settings will be printed at the end. The `cmake --build .` command will launch the compilation, which, if successful, will ultimately produce a library `liblammps.a` and the LAMMPS executable `lmp` inside the `build` folder.

Compilation can take a long time, since LAMMPS is a large project with many features. If your machine has multiple CPU cores (most do these days), you can speed this up by compiling sources in parallel with `make -j N` (with `N` being the maximum number of concurrently executed tasks). Also installation of the `ccache` (= Compiler Cache) software may speed up repeated compilation even more, e.g. during code development.

After the initial build, whenever you edit LAMMPS source files, enable or disable packages, change compiler flags or build options, you must re-compile and relink the LAMMPS executable with `cmake --build .` (or `make`). If the compilation fails for some reason, try running `cmake .` and then compile again. The included dependency tracking should make certain that only the necessary subset of files are re-compiled. You can also delete compiled objects, libraries and executables with `cmake --build . --target clean` (or `make clean`).

After compilation, you may optionally install the LAMMPS executable into your system with:

```
make install # optional, copy compiled files into installation location
```

This will install the LAMMPS executable and library, some tools (if configured) and additional files like LAMMPS API headers, manpages, potential and force field files. The location of the installation tree defaults to `${HOME}/.local`.



### 3.1.3 Configuration and build options

The CMake commands have one mandatory argument: a folder containing a file called `CMakeLists.txt` (for LAMMPS it is located in the `cmake` folder) or a build folder containing a file called `CMakeCache.txt`, which is generated at the end of the CMake configuration step. The cache file contains all current CMake settings.

To modify settings, enable or disable features, you need to set *variables* with either the `-D` command line flag (`-D VARIABLE1_NAME=value`) or change them in the text mode of graphical user interface. The `-D` flag can be used several times in one command.

For your convenience we provide *CMake presets* that combine multiple settings to enable optional LAMMPS packages or use a different compiler tool chain. Those are loaded with the `-C` flag (`-C ../cmake/presets/basic.cmake`). This step would only be needed once, as the settings from the preset files are stored in the `CMakeCache.txt` file. It is also possible to customize the build by adding one or more `-D` flags to the CMake command line.

Generating files for alternate build tools (e.g. Ninja) and project files for IDEs like Eclipse, CodeBlocks, or Kate can be selected using the `-G` command line flag. A list of available generator settings for your specific CMake version is given when running `cmake --help`.

### 3.1.4 Multi-configuration build systems

Throughout this manual it is mostly assumed that LAMMPS is being built on a Unix-like operating system with “make” as the underlying “builder”, since this is the most common case. In this case the build “configuration” is chose using `-D CMAKE_BUILD_TYPE=<configuration>` with `<configuration>` being one of “Release”, “Debug”, “RelWithDebInfo”, or “MinSizeRel”. Some build tools, however, can also use or even require to have a so-called multi-configuration build system setup. For those the built type (or configuration) is chosen at compile time using the same build files. E.g. with:

```
cmake --build build-multi --config Release
```

In that case the resulting binaries are not in the build folder directly but in sub-directories corresponding to the build type (i.e. Release in the example from above). Similarly, for running unit tests the configuration is selected with the `-C` flag:

```
ctest -C Debug
```

The CMake scripts in LAMMPS have basic support for being compiled using a multi-config build system, but not all of it has been ported. This is in particular applicable to compiling packages that require additional libraries that would be downloaded and compiled by CMake. The “windows” preset file tries to keep track of which packages can be compiled natively with the MSVC compilers out-of-the box. Not all of those external libraries are portable to Windows either.

### 3.1.5 Installing CMake

Check if your machine already has CMake installed:

```
which cmake           # do you have it?
which cmake3          # version 3 may have this name
cmake --version       # what specific version you have
```

On clusters or supercomputers which use environment modules to manage software packages, do this:

```
module list           # is a module for cmake already loaded?
module avail          # is a module for cmake available?
module load cmake     # load cmake module with appropriate name
```

Most Linux distributions offer pre-compiled cmake packages through their package management system. If you do not have CMake or a recent enough version (Note: for CentOS 7.x you need to enable the EPEL repository), you can download the latest version from <https://cmake.org/download/>. Instructions on how to install it on various platforms can be found [on this page](#).

## 3.2 Build LAMMPS with make

Building LAMMPS with traditional makefiles requires that you have a `Makefile.<machine>` file appropriate for your system in either the `src/MAKE`, `src/MAKE/MACHINES`, `src/MAKE/OPTIONS`, or `src/MAKE/MINE` directory (see below). It can include various options for customizing your LAMMPS build with a number of global compilation options and features.

### 3.2.1 Requirements

Those makefiles are written for and tested with GNU make and may not be compatible with other make programs. In most cases, if the “make” program is not GNU make, then there will be a GNU make program available under the name “gmake”. If GNU make or a compatible make is not available, you may have to first install it or switch to building with *CMake*. The makefiles of the traditional make based build process and the scripts they are calling expect a few additional tools to be available and functioning.

- a working C/C++ compiler toolchain supporting the C++11 standard; on Linux these are often the GNU compilers. Some older compilers require adding flags like `-std=c++11` to enable the C++11 mode.
- a Bourne shell compatible “Unix” shell program (often this is `bash`)
- a few shell utilities: `ls`, `mv`, `ln`, `rm`, `grep`, `sed`, `tr`, `cat`, `touch`, `diff`, `dirname`
- python (optional, required for `make lib-<pkg>` in the `src` folder). python scripts are currently tested with python 2.7 and 3.6. The procedure for *building the documentation* requires python 3.5 or later.

### 3.2.2 Getting started

To include LAMMPS packages (i.e. optional commands and styles) you must enable (or “install”) them first, as discussed on the *Build package* page. If a packages requires (provided or external) libraries, you must configure and build those libraries **before** building LAMMPS itself and especially **before** enabling such a package with `make yes-<package>`. *Building LAMMPS with CMake* can automate much of this for many types of machines, especially workstations, desktops, and laptops, so we suggest you try it first when building LAMMPS in those cases.

The commands below perform a default LAMMPS build, producing the LAMMPS executable `lmp_serial` and `lmp_mpi` in `lammps/src`:

```
cd lammps/src    # change to main LAMMPS source folder
make serial      # build a serial LAMMPS executable using GNU g++
make mpi         # build a parallel LAMMPS executable with MPI
make             # see a variety of make options
```

Compilation can take a long time, since LAMMPS is a large project with many features. If your machine has multiple CPU cores (most do these days), you can speed this up by compiling sources in parallel with `make -j N` (with `N` being the maximum number of concurrently executed tasks). Also installation of the `ccache` (= Compiler Cache) software may speed up repeated compilation even more, e.g. during code development.

After the initial build, whenever you edit LAMMPS source files, or add or remove new files to the source directory (e.g. by installing or uninstalling packages), you must re-compile and relink the LAMMPS executable with the same `make <machine>` command. The makefile’s dependency tracking should insure that only the necessary subset of files

are re-compiled. If you change settings in the makefile, you have to recompile *everything*. To delete all objects you can use `make clean-<machine>`.

**Note:** Before the actual compilation starts, LAMMPS will perform several steps to collect information from the configuration and setup that is then embedded into the executable. When you build LAMMPS for the first time, it will also compile a tool to quickly assemble a list of dependencies, that are required for the make program to correctly detect which parts need to be recompiled after changes were made to the sources.

### 3.2.3 Customized builds and alternate makefiles

The `src/MAKE` directory tree contains the `Makefile.<machine>` files included in the LAMMPS distribution. Typing `make example` uses `Makefile.example` from one of those folders, if available. Thus the `make serial` and `make mpi` lines above use `src/MAKE/Makefile.serial` and `src/MAKE/Makefile.mpi`, respectively. Other makefiles are in these directories:

```
OPTIONS      # Makefiles which enable specific options
MACHINES     # Makefiles for specific machines
MINE         # customized Makefiles you create (you may need to create this folder)
```

Simply typing `make` lists all the available `Makefile.<machine>` files with a single line description toward the end of the output. A file with the same name can appear in multiple folders (not a good idea). The order the directories are searched is as follows: `src/MAKE/MINE`, `src/MAKE`, `src/MAKE/OPTIONS`, `src/MAKE/MACHINES`. This gives preference to a customized file you put in `src/MAKE/MINE`. If you create your own custom makefile under a new name, please edit the first line with the description and machine name, so you will not confuse yourself, when looking at the machine summary.

Makefiles you may wish to try include these (some require a package first be installed). Many of these include specific compiler flags for optimized performance. Please note, however, that some of these customized machine Makefile are contributed by users. Since both compilers, OS configurations, and LAMMPS itself keep changing, their settings may become outdated:

```
make mac      # build serial LAMMPS on a Mac
make mac_mpi  # build parallel LAMMPS on a Mac
make intel_cpu # build with the INTEL package optimized for CPUs
make knl      # build with the INTEL package optimized for KNLs
make opt      # build with the OPT package optimized for CPUs
make omp      # build with the OPENMP package optimized for OpenMP
make kokkos_omp # build with the KOKKOS package for OpenMP
make kokkos_cuda_mpi # build with the KOKKOS package for GPUs
make kokkos_phi # build with the KOKKOS package for KNLs
```

## 3.3 Link LAMMPS as a library to another code

LAMMPS is designed as a library of C++ objects that can be integrated into other applications including Python scripts. The files `src/library.cpp` and `src/library.h` define a C-style API for using LAMMPS as a library. See the [Library interface to LAMMPS](#) page for a description of the interface and how to use it for your needs.

The [Basic build options](#) page explains how to build LAMMPS as either a shared or static library. This results in a file in the compilation folder called `liblammps.a` or `liblammps_<name>.a` in case of building a static library. In case of a shared library the name is the same only that the suffix is going to be either `.so` or `.dylib` or `.dll` instead of `.a`

depending on the OS. In some cases the `.so` file may be a symbolic link to a file with the suffix `.so.0` (or some other number).

---

**Note:** Care should be taken to use the same MPI library for the calling code and the LAMMPS library unless LAMMPS is to be compiled without (real) MPI support using the include STUBS MPI library.

---

### 3.3.1 Link with LAMMPS as a static library

The calling application can link to LAMMPS as a static library with compilation and link commands as in the examples shown below. These are examples for a code written in C in the file `caller.c`. The benefit of linking to a static library is, that the resulting executable is independent of that library since all required executable code from the library is copied into the calling executable.

---

#### CMake build

This assumes that LAMMPS has been configured without setting a `LAMMPS_MACHINE` name, installed with “make install”, and the `PKG_CONFIG_PATH` environment variable has been updated to include the `liblammps.pc` file installed into the configured destination folder. The commands to compile and link a coupled executable are then:

```
mpicc -c -O $(pkgconf liblammps --cflags) caller.c
mpicxx -o caller caller.o -$(pkgconf liblammps --libs)
```

---

#### Traditional make

This assumes that LAMMPS has been compiled in the folder `/${HOME}/lammps/src` with “make mpi”. The commands to compile and link a coupled executable are then:

```
mpicc -c -O -I${HOME}/lammps/src caller.c
mpicxx -o caller caller.o -L${HOME}/lammps/src -llammps_mpi
```

The `-I` argument is the path to the location of the `library.h` header file containing the interface to the LAMMPS C-style library interface. The `-L` argument is the path to where the `liblammps_mpi.a` file is located. The `-llammps_mpi` argument is shorthand for telling the compiler to link the file `liblammps_mpi.a`. If LAMMPS has been built as a shared library, then the linker will use `liblammps_mpi.so` instead. If both files are available, the linker will usually prefer the shared library. In case of a shared library, you may need to update the `LD_LIBRARY_PATH` environment variable or running the `caller` executable will fail since it cannot find the shared library at runtime.

---

However, it is only as simple as shown above for the case of a plain LAMMPS library without any optional packages that depend on libraries (bundled or external) or when using a shared library. Otherwise, you need to include all flags, libraries, and paths for the coupled executable, that are also required to link the LAMMPS executable.

---

#### CMake build

When using CMake, additional libraries with sources in the `lib` folder are built, but not included in `liblammps.a` and (currently) not installed with `make install` and not included in the `pkgconfig` configuration file. They can be found in the top level build folder, but you have to determine the necessary

link flags manually. It is therefore recommended to either use the traditional make procedure to build and link with a static library or build and link with a shared library instead.

### Traditional make

After you have compiled a static LAMMPS library using the conventional build system for example with “make mode=static serial”. And you also have installed the POEMS package after building its bundled library in lib/poems. Then the commands to build and link the coupled executable change to:

```
gcc -c -O -I${HOME}/lammps/src -caller.c
g++ -o caller caller.o -L${HOME}/lammps/lib/poems \
    -L${HOME}/lammps/src/STUBS -L${HOME}/lammps/src \
    -llammps_serial -lpoems -lmpi_stubs
```

Note, that you need to link with g++ instead of gcc even if you have written your code in C, since LAMMPS itself is C++ code. You can display the currently applied settings for building LAMMPS for the “serial” machine target by using the command:

```
make mode=print serial
```

Which should output something like:

```
# Compiler:
CXX=g++
# Linker:
LD=g++
# Compilation:
CXXFLAGS=-g -O3 -DLAMMPS_GZIP -DLAMMPS_MEMALIGN=64 -I${HOME}/compile/lammps/
↪lib/poems -I${HOME}/compile/lammps/src/STUBS
# Linking:
LDFLAGS=-g -O
# Libraries:
LDLIBS=-L${HOME}/compile/lammps/src -llammps_serial -L${HOME}/compile/lammps/
↪lib/poems -L${HOME}/compile/lammps/src/STUBS -lpoems -lmpi_stubs
```

From this you can gather the necessary paths and flags. With makefiles for other *machine* configurations you need to do the equivalent and replace “serial” with the corresponding “machine” name of the makefile.

## 3.3.2 Link with LAMMPS as a shared library

When linking to LAMMPS built as a shared library, the situation becomes much simpler, as all dependent libraries and objects are either included in the shared library or registered as a dependent library in the shared library file. Thus those libraries need not to be specified when linking the calling executable. Only the *-l* flags are needed. So the example case from above of the serial version static LAMMPS library with the POEMS package installed becomes:

### CMake build

The commands with a shared LAMMPS library compiled with the CMake build process are the same as for the static library.

```
mpicxx -c -O $(pkgconf liblammps --cflags) caller.c
mpicxx -o caller caller.o -$(pkgconf --libs)
```

---

### Traditional make

The commands with a shared LAMMPS library compiled with the traditional make build using `make mode=shared serial` becomes:

```
gcc -c -O -I${HOME}/lammps/src -caller.c
g++ -o caller caller.o -L${HOME}/lammps/src -llammps_serial
```

---

### Locating liblammps.so at runtime

Unlike with a static link, now the `liblammps.so` file is required at runtime and needs to be in a folder, where the shared linker program of the operating system can find it. This would be either a folder like `/usr/local/lib64` or `${HOME}/.local/lib64` or a folder pointed to by the `LD_LIBRARY_PATH` environment variable. You can type

```
printenv LD_LIBRARY_PATH
```

to see what directories are in that list.

Or you can add the LAMMPS `src` directory or the directory you performed a CMake style build in to your `LD_LIBRARY_PATH` environment variable, so that the current version of the shared library is always available to programs that use it.

For the Bourne or Korn shells (`/bin/sh`, `/bin/ksh`, `/bin/bash` etc.), you would add something like this to your `${HOME}/.profile` file:

```
LD_LIBRARY_PATH ${LD_LIBRARY_PATH}:/usr/lib64}:${HOME}/lammps/src
export LD_LIBRARY_PATH
```

For the `csh` or `tsh` shells, you would equivalently add something like this to your `${HOME}/.cshrc` file:

```
setenv LD_LIBRARY_PATH ${LD_LIBRARY_PATH}:${HOME}/lammps/src
```

You can verify whether all required shared libraries are found with the `ldd` tool. Example:

```
$ LD_LIBRARY_PATH=/home/user/lammps/src ldd caller
linux-vdso.so.1 (0x00007ffe729e0000)
liblammps.so => /home/user/lammps/src/liblammps.so (0x00007fc91bb9e000)
libstdc++.so.6 => /lib64/libstdc++.so.6 (0x00007fc91b984000)
libm.so.6 => /lib64/libm.so.6 (0x00007fc91b83e000)
libgcc_s.so.1 => /lib64/libgcc_s.so.1 (0x00007fc91b824000)
libc.so.6 => /lib64/libc.so.6 (0x00007fc91b65b000)
/lib64/ld-linux-x86-64.so.2 (0x00007fc91c094000)
```

If a required library is missing, you would get a ‘not found’ entry:

```
$ ldd caller
linux-vdso.so.1 (0x00007ffd672fe000)
liblammps.so => not found
libstdc++.so.6 => /usr/lib64/libstdc++.so.6 (0x00007fb7c7e86000)
libm.so.6 => /usr/lib64/libm.so.6 (0x00007fb7c7d40000)
libgcc_s.so.1 => /usr/lib64/libgcc_s.so.1 (0x00007fb7c7d26000)
```

(continues on next page)

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```
libc.so.6 => /usr/lib64/libc.so.6 (0x00007fb7c7b5d000)
/lib64/ld-linux-x86-64.so.2 (0x00007fb7c80a2000)
```

## 3.4 Basic build options

The following topics are covered on this page, for building with both CMake and make:

- *Serial vs parallel build*
- *Choice of compiler and compile/link options*
- *Build the LAMMPS executable and library*
- *Including and removing debug support*
- *Install LAMMPS after a build*

### 3.4.1 Serial vs parallel build

LAMMPS is written to use the ubiquitous [MPI \(Message Passing Interface\)](#) library API for distributed memory parallel computation. You need to have such a library installed for building and running LAMMPS in parallel using a domain decomposition parallelization. It is compatible with the MPI standard version 2.x and later. LAMMPS can also be built into a “serial” executable for use with a single processor using the bundled MPI STUBS library.

Independent of the distributed memory MPI parallelization, parts of LAMMPS are also written with support for shared memory parallelization using the [OpenMP](#) threading standard. A more detailed discussion of that is below.

#### CMake build

```
-D BUILD_MPI=value      # yes or no, default is yes if CMake finds MPI, else
↳no
-D BUILD_OMP=value      # yes or no, default is yes if a compatible compiler
↳is detected
-D LAMMPS_MACHINE=name  # name = mpi, serial, mybox, titan, laptop, etc
                        # no default value
```

The executable created by CMake (after running make) is named `lmp` unless the `LAMMPS_MACHINE` option is set. When setting `LAMMPS_MACHINE=name` the executable will be called `lmp_name`. Using `BUILD_MPI=no` will enforce building a serial executable using the MPI STUBS library.

#### Traditional make

The build with traditional makefiles has to be done inside the source folder `src`.

```
make mpi                # parallel build, produces lmp_mpi using Makefile.mpi
make serial             # serial build, produces lmp_serial using Makefile/
↳serial
make mybox              # uses Makefile.mybox to produce lmp_mybox
```

Any `make machine` command will look up the make settings from a file `Makefile.machine` in the folder `src/MAKE` or one of its sub-directories `MINE`, `MACHINES`, or `OPTIONS`, create a folder `Obj_machine` with all objects and generated files and an executable called `lmp_machine`. The standard parallel build with `make mpi` assumes a standard MPI installation with MPI compiler wrappers where all necessary compiler and linker flags to get access and link with the suitable MPI headers and libraries are set by the wrapper programs. For other cases or the serial build, you have to adjust the make file variables `MPI_INC`, `MPI_PATH`, `MPI_LIB` as well as `CC` and `LINK`. To enable OpenMP threading usually a compiler specific flag needs to be added to the compile and link commands. For the GNU compilers, this is `-fopenmp`, which can be added to the `CC` and `LINK` makefile variables.

For the serial build the following make variables are set (see `src/MAKE/Makefile.serial`):

<code>CC =</code>	<code>g++</code>
<code>LINK =</code>	<code>g++</code>
<code>MPI_INC =</code>	<code>-I../STUBS</code>
<code>MPI_PATH =</code>	<code>-L../STUBS</code>
<code>MPI_LIB =</code>	<code>-lmpi_stubs</code>

You also need to build the `STUBS` library for your platform before making LAMMPS itself. A `make serial` build does this for you automatically, otherwise, type `make mpi-stubs` from the `src` directory, or `make` from the `src/STUBS` dir. If the build fails, you may need to edit the `STUBS/Makefile` for your platform. The stubs library does not provide MPI/IO functions required by some LAMMPS packages, e.g. `MPIIO` or `LATBOLTZ`, and thus is not compatible with those packages.

---

**Note:** The file `src/STUBS/mpi.cpp` provides a CPU timer function called `MPI_Wtime()` that calls `gettimeofday()`. If your operating system does not support `gettimeofday()`, you will need to insert code to call another timer. Note that the ANSI-standard function `clock()` rolls over after an hour or so, and is therefore insufficient for timing long LAMMPS simulations.

---

## MPI and OpenMP support in LAMMPS

If you are installing MPI yourself to build a parallel LAMMPS executable, we recommend either `MPICH` or `OpenMPI` which are regularly used and tested with LAMMPS by the LAMMPS developers. `MPICH` can be downloaded from the [MPICH home page](#) and `OpenMPI` can be downloaded correspondingly from the [OpenMPI home page](#). Other MPI packages should also work. No specific vendor provided and standard compliant MPI library is currently known to be incompatible with LAMMPS. If you are running on a large parallel machine, your system admins or the vendor should have already installed a version of MPI, which is likely to be faster than a self-installed `MPICH` or `OpenMPI`, so you should study the provided documentation to find out how to build and link with it.

The majority of OpenMP (threading) support in LAMMPS is provided by the `OPENMP` package; see the [OPENMP package](#) page for details. The `INTEL` package also includes OpenMP threading (it is compatible with `OPENMP` and will usually fall back on styles from that package, if a `INTEL` does not exist) and adds vectorization support when compiled with compatible compilers, in particular the Intel compilers on top of OpenMP. Also, the `KOKKOS` package can be compiled to include OpenMP threading.

In addition, there are a few commands in LAMMPS that have native OpenMP support included as well. These are commands in the `MPIIO`, `ML-SNAP`, `DIFFRACTION`, and `DPD-REACT` packages. In addition some packages support OpenMP threading indirectly through the libraries they interface to: e.g. `LATTE`, `KSPACE`, and `COLVARS`. See the [Packages details](#) page for more info on these packages and the pages for their respective commands for OpenMP threading info.

For CMake, if you use `BUILD_OMP=yes`, you can use these packages and turn on their native OpenMP support and turn on their native OpenMP support at run time, by setting the `OMP_NUM_THREADS` environment variable before you



launch LAMMPS.

For building via conventional make, the CCFLAGS and LINKFLAGS variables in Makefile.machine need to include the compiler flag that enables OpenMP. For GNU compilers it is `-fopenmp`. For (recent) Intel compilers it is `-qopenmp`. If you are using a different compiler, please refer to its documentation.

### OpenMP Compiler compatibility

Some compilers do not fully support the `default(none)` directive and others (e.g. GCC version 9 and beyond, Clang version 10 and later) may implement strict OpenMP 4.0 and later semantics, which are incompatible with the OpenMP 3.1 semantics used in LAMMPS for maximal compatibility with compiler versions in use. If compilation with OpenMP enabled fails because of your compiler requiring strict OpenMP 4.0 semantics, you can change the behavior by adding `-D LAMMPS_OMP_COMPAT=4` to the `LMP_INC` variable in your makefile, or add it to the command line while configuring with CMake. LAMMPS will auto-detect a suitable setting for most GNU, Clang, and Intel compilers.

## 3.4.2 Choice of compiler and compile/link options

The choice of compiler and compiler flags can be important for maximum performance. Vendor provided compilers for a specific hardware can produce faster code than open-source compilers like the GNU compilers. On the most common x86 hardware most popular C++ compilers are quite similar in performance of C/C++ code at high optimization levels. When using the INTEL package, there is a distinct advantage in using the [Intel C++ compiler](#) due to much improved vectorization through SSE and AVX instructions on compatible hardware as the source code includes changes and Intel compiler specific directives to enable high degrees of vectorization. This may change over time as equivalent vectorization directives are included into OpenMP standard revisions and other compilers adopt them.

On parallel clusters or supercomputers which use “environment modules” for their compile/link environments, you can often access different compilers by simply loading the appropriate module before building LAMMPS.

### CMake build

By default CMake will use the compiler it finds according to internal preferences and it will add optimization flags appropriate to that compiler and any *accelerator packages* you have included in the build. CMake will check if the detected or selected compiler is compatible with the C++ support requirements of LAMMPS and stop with an error, if this is not the case.

You can tell CMake to look for a specific compiler with setting CMake variables (listed below) during configuration. For a few common choices, there are also presets in the `cmake/presets` folder. For convenience, there is a `CMAKE_TUNE_FLAGS` variable that can be set to apply global compiler options (applied to compilation only), to be used for adding compiler or host specific optimization flags in addition to the “flags” variables listed below. You may also specify the corresponding `CMAKE_*_FLAGS` variables individually, if you want to experiment with alternate optimization flags. You should specify all 3 compilers, so that the (few) LAMMPS source files written in C or Fortran are built with a compiler consistent with the one used for the C++ files:

```
-D CMAKE_CXX_COMPILER=name           # name of C++ compiler
-D CMAKE_C_COMPILER=name            # name of C compiler
-D CMAKE_Fortran_COMPILER=name       # name of Fortran compiler

-D CMAKE_CXX_FLAGS=string            # flags to use with C++ compiler
-D CMAKE_C_FLAGS=string              # flags to use with C compiler
-D CMAKE_Fortran_FLAGS=string        # flags to use with Fortran compiler
```

A few example command lines are:

```
# Building with GNU Compilers:
cmake ../cmake -DCMAKE_C_COMPILER=gcc -DCMAKE_CXX_COMPILER=g++ -DCMAKE_Fortran_
↪COMPILER=gfortran
# Building with Intel Compilers:
cmake ../cmake -DCMAKE_C_COMPILER=icc -DCMAKE_CXX_COMPILER=icpc -DCMAKE_
↪Fortran_COMPILER=ifort
# Building with Intel oneAPI Compilers:
cmake ../cmake -DCMAKE_C_COMPILER=icx -DCMAKE_CXX_COMPILER=icpx -DCMAKE_
↪Fortran_COMPILER=ifx
# Building with LLVM/Clang Compilers:
cmake ../cmake -DCMAKE_C_COMPILER=clang -DCMAKE_CXX_COMPILER=clang++ -DCMAKE_
↪Fortran_COMPILER=flang
# Building with PGI/Nvidia Compilers:
cmake ../cmake -DCMAKE_C_COMPILER=pgcc -DCMAKE_CXX_COMPILER=pgc++ -DCMAKE_
↪Fortran_COMPILER=pgfortran
```

For compiling with the Clang/LLVM compilers a CMake preset is provided that can be loaded with `-C ../cmake/presets/clang.cmake`. Similarly, `-C ../cmake/presets/intel.cmake` should switch the compiler toolchain to the legacy Intel compilers, `-C ../cmake/presets/oneapi.cmake` will switch to the LLVM based oneAPI Intel compilers, and `-C ../cmake/presets/pgi.cmake` will switch the compiler to the PGI compilers.

In addition you can set `CMAKE_TUNE_FLAGS` to specifically add compiler flags to tune for optimal performance on given hosts. By default this variable is empty.

---

**Note:** When the `cmake` command completes, it prints a summary to the screen which compilers it is using and what flags and settings will be used for the compilation. Note that if the top-level compiler is `mpicxx`, it is simply a wrapper on a real compiler. The underlying compiler info is what CMake will try to determine and report. You should check to confirm you are using the compiler and optimization flags you want.

---

---

### Makefile.machine settings for traditional make

The “compiler/linker settings” section of a `Makefile.machine` lists compiler and linker settings for your C++ compiler, including optimization flags. For a parallel build it is recommended to use `mpicxx` or `mpiCC`, since these compiler wrappers will include a variety of settings appropriate for your MPI installation and thus avoiding the guesswork of finding the right flags.

Parallel build (see `src/MAKE/Makefile.mpi`):

```
CC =          mpicxx
CCFLAGS =     -g -O3
LINK =        mpicxx
LINKFLAGS =   -g -O
```

Serial build with GNU `gcc` (see `src/MAKE/Makefile.serial`):

```
CC =          g++
CCFLAGS =     -g -O3
LINK =        g++
LINKFLAGS =   -g -O
```

**Note:** If compilation stops with a message like the following:

```
g++ -g -O3 -DLAMMPS_GZIP -DLAMMPS_MEMALIGN=64 -I../STUBS -c ../main.cpp
In file included from ../pointers.h:24:0,
      from ../input.h:17,
      from ../main.cpp:16:
../lmptype.h:34:2: error: #error LAMMPS requires a C++11 (or later) compliant
↪ compiler. Enable C++11 compatibility or upgrade the compiler.
```

then you have either an unsupported (old) compiler or you have to turn on C++11 mode. The latter applies to GCC 4.8.x shipped with RHEL 7.x and CentOS 7.x or GCC 5.4.x shipped with Ubuntu16.04. For those compilers, you need to add the `-std=c++11` flag. If there is no compiler that supports this flag (or equivalent), you would have to install a newer compiler that supports C++11; either as a binary package or through compiling from source.

If you build LAMMPS with any *Accelerator packages* included, there may be specific compiler or linker flags that are either required or recommended to enable required features and to achieve optimal performance. You need to include these in the `CCFLAGS` and `LINKFLAGS` settings above. For details, see the documentation for the individual packages listed on the *Accelerator packages* page. Or examine these files in the `src/MAKE/OPTIONS` directory. They correspond to each of the 5 accelerator packages and their hardware variants:

```
Makefile.opt           # OPT package
Makefile.omp           # OPENMP package
Makefile.intel_cpu     # INTEL package for CPUs
Makefile.intel_coprocessor # INTEL package for KNLs
Makefile.gpu           # GPU package
Makefile.kokkos_cuda_mpi # KOKKOS package for GPUs
Makefile.kokkos_omp    # KOKKOS package for CPUs (OpenMP)
Makefile.kokkos_phi    # KOKKOS package for KNLs (OpenMP)
```

### 3.4.3 Build the LAMMPS executable and library

LAMMPS is always built as a library of C++ classes plus an executable. The executable is a simple `main()` function that sets up MPI and then creates a LAMMPS class instance from the LAMMPS library, which will then process commands provided via a file or from the console input. The LAMMPS library can also be called from another application or a scripting language. See the *Howto couple* doc page for more info on coupling LAMMPS to other codes. See the *Python* page for more info on wrapping and running LAMMPS from Python via its library interface.

#### CMake build

For CMake builds, you can select through setting CMake variables between building a shared or a static LAMMPS library and what kind of suffix is added to them (in case you want to concurrently install multiple variants of binaries with different settings). If none are set, defaults are applied.

```
-D BUILD_SHARED_LIBS=value # yes or no (default)
-D LAMMPS_MACHINE=name    # name = mpi, serial, mybox, titan, laptop, etc
                          # no default value
```

The compilation will always produce a LAMMPS library and an executable linked to it. By default this will be a static library named `liblammps.a` and an executable named `lmp`. Setting `BUILD_SHARED_LIBS=yes` will instead produce a shared library called `liblammps.so` (or `liblammps.dylib` or `liblammps.dll` depending on the platform). If `LAMMPS_MACHINE=name` is set in addition, the name of the generated libraries will be changed to either `liblammps_name.a` or `liblammps_name.so`, respectively and the executable will be called `lmp_name`.

---

### Traditional make

With the traditional makefile based build process, the choice of the generated executable or library depends on the “mode” setting. Several options are available and `mode=static` is the default.

```
make machine           # build LAMMPS executable lmp_machine
make mode=static machine # same as "make machine"
make mode=shared machine # build LAMMPS shared lib liblammps_machine.so
↳instead
```

The “static” build will generate a static library called `liblammps_machine.a` and an executable named `lmp_machine`, while the “shared” build will generate a shared library `liblammps_machine.so` instead and `lmp_machine` will be linked to it. The build step will also create generic soft links, named `liblammps.a` and `liblammps.so`, which point to the specific `liblammps_machine.a/so` files.

---

### Additional information

Note that for creating a shared library, all the libraries it depends on must be compiled to be compatible with shared libraries. This should be the case for libraries included with LAMMPS, such as the dummy MPI library in `src/STUBS` or any package libraries in the `lib` directory, since they are always built in a shared library compatible way using the `-fPIC` compiler switch. However, if an auxiliary library (like MPI or FFTW) does not exist as a compatible format, the shared library linking step may generate an error. This means you will need to install a compatible version of the auxiliary library. The build instructions for that library should tell you how to do this.

As an example, here is how to build and install the [MPICH library](#), a popular open-source version of MPI, as a shared library in the default `/usr/local/lib` location:

```
./configure --enable-shared
make
make install
```

You may need to use `sudo make install` in place of the last line if you do not have write privileges for `/usr/local/lib` or use the `--prefix` configuration option to select an installation folder, where you do have write access. The end result should be the file `/usr/local/lib/libmpich.so`. On many Linux installations the folder `/${HOME}/.local` is an alternative to using `/usr/local` and does not require superuser or `sudo` access. In that case the configuration step becomes:

```
./configure --enable-shared --prefix=${HOME}/.local
```

Avoiding to use “`sudo`” for custom software installation (i.e. from source and not through a package manager tool provided by the OS) is generally recommended to ensure the integrity of the system software installation.

---

### 3.4.4 Including or removing debug support

By default the compilation settings will include the `-g` flag which instructs the compiler to include debug information (e.g. which line of source code a particular instruction correspond to). This can be extremely useful in case LAMMPS crashes and can help to provide crucial information in *tracking down the origin of a crash* and help the LAMMPS developers fix bugs in the source code. However, this increases the storage requirements for object files, libraries, and the executable 3-5 fold.

If this is a concern, you can change the compilation settings or remove the debug information from the LAMMPS executable:

- **Traditional make:** edit your Makefile.<machine> to remove the `-g` flag from the `CCFLAGS` and `LINKFLAGS` definitions
- **CMake:** use `-D CMAKE_BUILD_TYPE=Release` or explicitly reset the applicable compiler flags (best done using the text mode or graphical user interface).
- **Remove debug info:** If you are only concerned about the executable being too large, you can use the `strip` tool (e.g. `strip lmp_serial`) to remove the debug information from the executable file. Do not strip libraries or object files, as that will render them unusable.

### 3.4.5 Build LAMMPS tools

Some tools described in *Auxiliary tools* can be built directly using CMake or Make.

#### CMake build

```
-D BUILD_TOOLS=value          # yes or no (default)
-D BUILD_LAMMPS_SHELL=value  # yes or no (default)
```

The generated binaries will also become part of the LAMMPS installation (see below).

#### Traditional make

```
cd lammps/tools
make all           # build all binaries of tools
make binary2txt   # build only binary2txt tool
make chain        # build only chain tool
make micelle2d    # build only micelle2d tool
make thermo_extract # build only thermo_extract tool

cd lammps/tools/lammps-shell
make              # build LAMMPS shell
```

### 3.4.6 Install LAMMPS after a build

After building LAMMPS, you may wish to copy the LAMMPS executable or library, along with other LAMMPS files (library header, doc files) to a globally visible place on your system, for others to access. Note that you may need super-user privileges (e.g. `sudo`) if the directory you want to copy files to is protected.

---

#### CMake build

```
cmake -D CMAKE_INSTALL_PREFIX=path [options ...] ../cmake
make                                     # perform make after CMake command
make install                             # perform the installation into prefix
```

During the installation process CMake will by default remove any runtime path settings for loading shared libraries. Because of this you may have to set or modify the `LD_LIBRARY_PATH` (or `DYLD_LIBRARY_PATH`) environment variable, if you are installing LAMMPS into a non-system location and/or are linking to libraries in a non-system location that depend on such runtime path settings. As an alternative you may set the CMake variable `LAMMPS_INSTALL_RPATH` to `on` and then the runtime paths for any linked shared libraries and the library installation folder for the LAMMPS library will be embedded and thus the requirement to set environment variables is avoided. The `off` setting is usually preferred for packaged binaries or when setting up environment modules, the `on` setting is more convenient for installing software into a non-system or personal folder.

---

---

#### Traditional make

There is no “install” option in the `src/Makefile` for LAMMPS. If you wish to do this you will need to first build LAMMPS, then manually copy the desired LAMMPS files to the appropriate system directories.

---

## 3.5 Optional build settings

LAMMPS can be built with several optional settings. Each sub-section explain how to do this for building both with CMake and make.

- *C++11 standard compliance* when building all of LAMMPS
  - *FFT library* for use with the `kpace_style ppm` command
  - *Size of LAMMPS integer types and size limits*
  - *Read or write compressed files*
  - *Output of JPG, PNG, and movie files* via the `dump image` or `dump movie` commands
  - *Memory allocation alignment*
  - *Workaround for long long integers*
  - *Exception handling when using LAMMPS as a library* to capture errors
  - *Trigger selected floating-point exceptions*
-

### 3.5.1 C++11 standard compliance

A C++11 standard compatible compiler is a requirement for compiling LAMMPS. LAMMPS version 3 March 2020 is the last version compatible with the previous C++98 standard for the core code and most packages. Most currently used C++ compilers are compatible with C++11, but some older ones may need extra flags to enable C++11 compliance. Example for GNU c++ 4.8.x:

```
CCFLAGS = -g -O3 -std=c++11
```

### 3.5.2 FFT library

When the KSPACE package is included in a LAMMPS build, the *kspace\_style ppm* command performs 3d FFTs which require use of an FFT library to compute 1d FFTs. The KISS FFT library is included with LAMMPS but other libraries can be faster. LAMMPS can use them if they are available on your system.

#### CMake build

```
-D FFT=value           # FFTW3 or MKL or KISS, default is FFTW3 if found,
↳ else KISS
-D FFT_SINGLE=value    # yes or no (default), no = double precision
-D FFT_PACK=value      # array (default) or pointer or memcpy
```

**Note:** The values for the FFT variable must be in upper-case. This is an exception to the rule that all CMake variables can be specified with lower-case values.

Usually these settings are all that is needed. If FFTW3 is selected, then CMake will try to detect, if threaded FFTW libraries are available and enable them by default. This setting is independent of whether OpenMP threads are enabled and a packages like KOKKOS or OPENMP is used. If CMake cannot detect the FFT library, you can set these variables to assist:

```
-D FFTW3_INCLUDE_DIR=path # path to FFTW3 include files
-D FFTW3_LIBRARY=path     # path to FFTW3 libraries
-D FFTW3_OMP_LIBRARY=path # path to FFTW3 OpenMP wrapper libraries
-D FFT_FFTW_THREADS=on    # enable using OpenMP threaded FFTW3 libraries
-D MKL_INCLUDE_DIR=path   # ditto for Intel MKL library
-D FFT_MKL_THREADS=on     # enable using threaded FFTs with MKL libraries
-D MKL_LIBRARY=path       # path to MKL libraries
```

#### Traditional make

To change the FFT library to be used and its options, you have to edit your machine Makefile. Below are examples how the makefile variables could be changed.

```
FFT_INC = -DFFT_FFTW3      # -DFFT_FFTW3, -DFFT_FFTW (same as -DFFT_FFTW3),
↳ -DFFT_MKL, or -DFFT_KISS
                                # default is KISS if not specified
FFT_INC = -DFFT_SINGLE    # do not specify for double precision
```

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```

FFT_INC = -DFFT_FFTW_THREADS # enable using threaded FFTW3 libraries
FFT_INC = -DFFT_MKL_THREADS  # enable using threaded FFTs with MKL libraries
FFT_INC = -DFFT_PACK_ARRAY   # or -DFFT_PACK_POINTER or -DFFT_PACK_MEMCPY
                               # default is FFT_PACK_ARRAY if not specified

```

```

FFT_INC = -I/usr/local/include
FFT_PATH = -L/usr/local/lib
FFT_LIB = -lfftw3 # FFTW3 double precision
FFT_LIB = -lfftw3 -lfftw3_omp # FFTW3 double precision with threads
↳ (needs -DFFT_FFTW_THREADS)
FFT_LIB = -lfftw3 -lfftw3f # FFTW3 single precision
FFT_LIB = -lmkl_intel_lp64 -lmkl_sequential -lmkl_core # MKL with
↳ Intel compiler, serial interface
FFT_LIB = -lmkl_gf_lp64 -lmkl_sequential -lmkl_core # MKL with GNU
↳ compiler, serial interface
FFT_LIB = -lmkl_intel_lp64 -lmkl_intel_thread -lmkl_core # MKL with
↳ Intel compiler, threaded interface
FFT_LIB = -lmkl_gf_lp64 -lmkl_gnu_thread -lmkl_core # MKL with GNU
↳ compiler, threaded interface
FFT_LIB = -lmkl_rt # MKL with automatic runtime selection of
↳ interface libs

```

As with CMake, you do not need to set paths in `FFT_INC` or `FFT_PATH`, if the compiler can find the FFT header and library files in its default search path. You must specify `FFT_LIB` with the appropriate FFT libraries to include in the link.

The **KISS FFT library** is included in the LAMMPS distribution. It is portable across all platforms. Depending on the size of the FFTs and the number of processors used, the other libraries listed here can be faster.

However, note that long-range Coulombics are only a portion of the per-timestep CPU cost, FFTs are only a portion of long-range Coulombics, and 1d FFTs are only a portion of the FFT cost (parallel communication can be costly). A breakdown of these timings is printed to the screen at the end of a run when using the `kpace_style ppm` command. The [Screen and logfile output](#) page gives more details. A more detailed (and time consuming) report of the FFT performance is generated with the `kpace_modify fftbench yes` command.

FFTW is a fast, portable FFT library that should also work on any platform and can be faster than the KISS FFT library. You can download it from [www.fftw.org](http://www.fftw.org). LAMMPS requires version 3.X; the legacy version 2.1.X is no longer supported.

Building FFTW for your box should be as simple as `./configure; make; make install`. The `install` command typically requires root privileges (e.g. invoke it via `sudo`), unless you specify a local directory with the “`-prefix`” option of `configure`. Type `./configure --help` to see various options.

The Intel MKL math library is part of the Intel compiler suite. It can be used with the Intel or GNU compiler (see the `FFT_LIB` setting above).

Performing 3d FFTs in parallel can be time consuming due to data access and required communication. This cost can be reduced by performing single-precision FFTs instead of double precision. Single precision means the real and imaginary parts of a complex datum are 4-byte floats. Double precision means they are 8-byte doubles. Note that Fourier transform and related PPM operations are somewhat less sensitive to floating point truncation errors and thus the resulting error is less than the difference in precision. Using the `-DFFT_SINGLE` setting trades off a little accuracy for reduced memory use and parallel communication costs for transposing 3d FFT data.

When using `-DFFT_SINGLE` with FFTW3 you may need to build the FFTW library a second time with support for single-precision.



For FFTW3, do the following, which should produce the additional library `libfftw3f.a` or `libfftw3f.so`.

```
make clean
./configure --enable-single; make; make install
```

Performing 3d FFTs requires communication to transpose the 3d FFT grid. The data packing/unpacking for this can be done in one of 3 modes (ARRAY, POINTER, MEMCPY) as set by the `FFT_PACK` syntax above. Depending on the machine, the size of the FFT grid, the number of processors used, one option may be slightly faster. The default is ARRAY mode.

### 3.5.3 Size of LAMMPS integer types and size limits

LAMMPS has a few integer data types which can be defined as either 4-byte (= 32-bit) or 8-byte (= 64-bit) integers at compile time. This has an impact on the size of a system that can be simulated or how large counters can become before “rolling over”. The default setting of “smallbig” is almost always adequate.

#### CMake build

With CMake the choice of integer types is made via setting a variable during configuration.

```
-D LAMMPS_SIZES=value # smallbig (default) or bigbig or smallsmall
```

If the variable is not set explicitly, “smallbig” is used.

#### Traditional build

If you want a setting different from the default, you need to edit the `LMP_INC` variable setting your machine Makefile.

```
LMP_INC = -DLAMMPS_SMALLBIG # or -DLAMMPS_BIGBIG or -DLAMMPS_SMALLSMALL
```

The default setting is `-DLAMMPS_SMALLBIG` if nothing is specified

### LAMMPS system size restrictions

	smallbig	bigbig	smallsmall
Total atom count	$2^{63}$ atoms (= $9.223 \cdot 10^{18}$ )	$2^{63}$ atoms (= $9.223 \cdot 10^{18}$ )	$2^{31}$ atoms (= $2.147 \cdot 10^9$ )
Total timesteps	$2^{63}$ steps (= $9.223 \cdot 10^{18}$ )	$2^{63}$ steps (= $9.223 \cdot 10^{18}$ )	$2^{31}$ steps (= $2.147 \cdot 10^9$ )
Atom ID values	$1 \leq i \leq 2^{31}$ (= $2.147 \cdot 10^9$ )	$1 \leq i \leq 2^{63}$ (= $9.223 \cdot 10^{18}$ )	$1 \leq i \leq 2^{31}$ (= $2.147 \cdot 10^9$ )
Image flag values	$-512 \leq i \leq 511$	$-1\,048\,576 \leq i \leq 1\,048\,575$	$-512 \leq i \leq 511$

The “bigbig” setting increases the size of image flags and atom IDs over “smallbig” and the “smallsmall” setting is only needed if your machine does not support 64-bit integers or incurs performance penalties when using them.

These are limits for the core of the LAMMPS code, specific features or some styles may impose additional limits. The *ATC* package cannot be compiled with the “bigbig” setting. Also, there are limitations when using the library interface where some functions with known issues have been replaced by dummy calls printing a corresponding error message rather than crashing randomly or corrupting data.

Atom IDs are not required for atomic systems which do not store bond topology information, though IDs are enabled by default. The `atom_modify id no` command will turn them off. Atom IDs are required for molecular systems with bond topology (bonds, angles, dihedrals, etc). Similarly, some force or compute or fix styles require atom IDs. Thus if you model a molecular system or use one of those styles with more than 2 billion atoms, you need the “bigbig” setting.

Regardless of the total system size limits, the maximum number of atoms per MPI rank (local + ghost atoms) is limited to 2 billion for atomic systems and 500 million for systems with bonds (the additional restriction is due to using the 2 upper bits of the local atom index in neighbor lists for storing special bonds info).

Image flags store 3 values per atom in a single integer which count the number of times an atom has moved through the periodic box in each dimension. See the `dump` manual page for a discussion. If an atom moves through the periodic box more than this limit, the value will “roll over”, e.g. from 511 to -512, which can cause diagnostics like the mean-squared displacement, as calculated by the `compute msd` command, to be faulty.

Also note that the GPU package requires its `lib/gpu` library to be compiled with the same size setting, or the link will fail. A CMake build does this automatically. When building with make, the setting in whichever `lib/gpu/Makefile` is used must be the same as above.

---

### 3.5.4 Output of JPG, PNG, and movie files

The `dump image` command has options to output JPEG or PNG image files. Likewise the `dump movie` command outputs movie files in a variety of movie formats. Using these options requires the following settings:

---

#### CMake build

```
-D WITH_JPEG=value      # yes or no
                        # default = yes if CMake finds JPEG files, else no
-D WITH_PNG=value       # yes or no
                        # default = yes if CMake finds PNG and ZLIB files,
↳ else no
-D WITH_FFMPEG=value    # yes or no
                        # default = yes if CMake can find ffmpeg, else no
```

Usually these settings are all that is needed. If CMake cannot find the graphics header, library, executable files, you can set these variables:

```
-D JPEG_INCLUDE_DIR=path # path to jpeglib.h header file
-D JPEG_LIBRARY=path     # path to libjpeg.a (.so) file
-D PNG_INCLUDE_DIR=path  # path to png.h header file
-D PNG_LIBRARY=path     # path to libpng.a (.so) file
-D ZLIB_INCLUDE_DIR=path # path to zlib.h header file
-D ZLIB_LIBRARY=path    # path to libz.a (.so) file
-D FFMPEG_EXECUTABLE=path # path to ffmpeg executable
```

---

#### Traditional make

```
LMP_INC = -DLAMMPS_JPEG -DLAMMPS_PNG -DLAMMPS_FFMPEG <other LMP_INC settings>

JPG_INC = -I/usr/local/include # path to jpeglib.h, png.h, zlib.h header
↳ files if make cannot find them
```

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```
JPG_PATH = -L/usr/lib          # paths to libjpeg.a, libpng.a, libz.a (.so)
↪files if make cannot find them
JPG_LIB = -ljpeg -lpng -lz     # library names
```

As with CMake, you do not need to set `JPG_INC` or `JPG_PATH`, if `make` can find the graphics header and library files in their default system locations. You must specify `JPG_LIB` with a list of graphics libraries to include in the link. You must make certain that the `ffmpeg` executable (or `ffmpeg.exe` on Windows) is in a directory where LAMMPS can find it at runtime; that is usually a directory list in your `PATH` environment variable.

Using `ffmpeg` to output movie files requires that your machine supports the “`popen`” function in the standard runtime library.

**Note:** On some clusters with high-speed networks, using the `fork()` library call (required by `popen()`) can interfere with the fast communication library and lead to simulations using `ffmpeg` to hang or crash.

### 3.5.5 Read or write compressed files

If this option is enabled, large files can be read or written with compression by `gzip` or similar tools by several LAMMPS commands, including `read_data`, `rerun`, and `dump`. Currently supported compression tools are: `gzip`, `bzip2`, `zstd`, and `lzma`.

#### CMake build

```
-D WITH_GZIP=value           # yes or no
                             # default is yes if CMake can find the gzip program,
↪else no
```

#### Traditional make

```
LMP_INC = -DLAMMPS_GZIP <other LMP_INC settings>
```

This option requires that your operating system fully supports the “`popen()`” function in the standard runtime library and that a `gzip` or other executable can be found by LAMMPS in the standard search path during a run.

**Note:** On clusters with high-speed networks, using the “`fork()`” library call (required by “`popen()`”) can interfere with the fast communication library and lead to simulations using compressed output or input to hang or crash. For selected operations, compressed file I/O is also available using a compression library instead, which is what the `COMPRESS` package enables.

### 3.5.6 Memory allocation alignment

This setting enables the use of the “`posix_memalign()`” call instead of “`malloc()`” when LAMMPS allocates large chunks of memory. Vector instructions on CPUs may become more efficient, if dynamically allocated memory is aligned on larger-than-default byte boundaries. On most current operating systems, the “`malloc()`” implementation returns pointers that are aligned to 16-byte boundaries. Using SSE vector instructions efficiently, however, requires memory blocks being aligned on 64-byte boundaries.

---

#### CMake build

```
-D LAMMPS_MEMALIGN=value           # 0, 8, 16, 32, 64 (default)
```

Use a LAMMPS\_MEMALIGN value of 0 to disable using “`posix_memalign()`” and revert to using the “`malloc()`” C-library function instead. When compiling LAMMPS for Windows systems, “`malloc()`” will always be used and this setting is ignored.

---

#### Traditional make

```
LMP_INC = -DLAMMPS_MEMALIGN=value  # 8, 16, 32, 64
```

Do not set `-DLAMMPS_MEMALIGN`, if you want to have memory allocated with the “`malloc()`” function call instead. `-DLAMMPS_MEMALIGN` **cannot** be used on Windows, as Windows different function calls with different semantics for allocating aligned memory, that are not compatible with how LAMMPS manages its dynamical memory.

---

### 3.5.7 Workaround for long long integers

If your system or MPI version does not recognize “long long” data types, the following setting will be needed. It converts “long long” to a “long” data type, which should be the desired 8-byte integer on those systems:

---

#### CMake build

```
-D LAMMPS_LONGLONG_TO_LONG=value    # yes or no (default)
```

---

#### Traditional make

```
LMP_INC = -DLAMMPS_LONGLONG_TO_LONG <other LMP_INC settings>
```

---

### 3.5.8 Exception handling when using LAMMPS as a library

This setting is useful when external codes drive LAMMPS as a library. With this option enabled, LAMMPS errors do not kill the calling code. Instead, the call stack is unwound and control returns to the caller, e.g. to Python. Of course, the calling code has to be set up to *catch* exceptions thrown from within LAMMPS.

---

#### CMake build

```
-D LAMMPS_EXCEPTIONS=value      # yes or no (default)
```

---



---

#### Traditional make

```
LMP_INC = -DLAMMPS_EXCEPTIONS  <other LMP_INC settings>
```

---



---

**Note:** When LAMMPS is running in parallel, it is not always possible to cleanly recover from an exception since not all parallel ranks may throw an exception and thus other MPI ranks may get stuck waiting for messages from the ones with errors.

---

### 3.5.9 Trigger selected floating-point exceptions

Many kinds of CPUs have the capability to detect when a calculation results in an invalid math operation like a division by zero or calling the square root with a negative argument. The default behavior on most operating systems is to continue and have values for NaN (= not a number) or Inf (= infinity). This allows software to detect and recover from such conditions. This behavior can be changed, however, often through use of compiler flags. On Linux systems (or more general on systems using the GNU C library), these so-called floating-point traps can also be selectively enabled through library calls. LAMMPS supports that by setting the `-DLAMMPS_TRAP_FPE` pre-processor define. As it is done in the `main()` function, this applies only to the standalone executable, not the library.

---

#### CMake build

```
-D CMAKE_TUNE_FLAGS--DLAMMPS_TRAP_FPE
```

---



---

#### Traditional make

```
LMP_INC = -DLAMMPS_TRAP_FPE  <other LMP_INC settings>
```

---

After compilation with this flag set, the LAMMPS executable will stop and produce a core dump when a division by zero, overflow, illegal math function argument or other invalid floating point operation is encountered.

## 3.6 Include packages in build

In LAMMPS, a package is a group of files that enable a specific set of features. For example, force fields for molecular systems or rigid-body constraints are in packages. In the `src` directory, each package is a sub-directory with the package name in capital letters.

An overview of packages is given on the [Packages](#) doc page. Brief overviews of each package are on the [Packages details](#) page.

When building LAMMPS, you can choose to include or exclude each package. In general there is no need to include a package if you never plan to use its features.

If you get a run-time error that a LAMMPS command or style is “unknown”, it is often because the command is contained in a package, and your build did not include that package. If the command or style *is* available in a package included in the LAMMPS distribution, the error message will indicate which package would be needed. Running LAMMPS with the `-h` *command-line switch* will print *all* optional commands and packages that were enabled when building that executable.

For the majority of packages, if you follow the single step below to include it, you can then build LAMMPS exactly the same as you would without any packages installed. A few packages may require additional steps, as explained on the [Build extras](#) page.

These links take you to the extra instructions for those select packages:

<a href="#">ADIOS</a>	<a href="#">ATC</a>	<a href="#">AWPMD</a>	<a href="#">COLVARS</a>	<a href="#">COMPRESS</a>	<a href="#">GPU</a>
<a href="#">H5MD</a>	<a href="#">INTEL</a>	<a href="#">KIM</a>	<a href="#">KOKKOS</a>	<a href="#">LATTE</a>	<a href="#">MACHDYN</a>
<a href="#">ML-HDNNP</a>	<a href="#">ML-PACE</a>	<a href="#">ML-QUIP</a>	<a href="#">MOLFILE</a>	<a href="#">MSCG</a>	<a href="#">NETCDF</a>
<a href="#">OPENMP</a>	<a href="#">OPT</a>	<a href="#">PLUMED</a>	<a href="#">POEMS</a>	<a href="#">PYTHON</a>	<a href="#">QMMM</a>
<a href="#">SCAFACOS</a>	<a href="#">VORONOI</a>	<a href="#">VTK</a>			

The mechanism for including packages is simple but different for CMake versus make.

---

### CMake build

```
-D PKG_NAME=value           # yes or no (default)
```

Examples:

```
-D PKG_MANYBODY=yes
-D PKG_INTEL=yes
```

All packages are included the same way. See the shortcut section below for how to install many packages at once with CMake.

---

**Note:** If you switch between building with CMake and make builds, no packages in the `src` directory can be installed when you invoke `cmake`. CMake will give an error if that is not the case, indicating how you can un-install all packages in the `src` dir.

---

---

---

### Traditional make

```
cd lammps/src
make ps           # check which packages are currently installed
make yes-name    # install a package with name
make no-name     # un-install a package with name
make mpi         # build LAMMPS with whatever packages are now
↳ installed
```

Examples:

```
make no-rigid
make yes-intel
```

All packages are included the same way. See the shortcut section below for how to install many packages at once with make.

---

**Note:** You must always re-build LAMMPS (via make) after installing or un-installing a package, for the action to take effect. The included dependency tracking will make certain only files that are required to be rebuilt are recompiled.

---

---

**Note:** You cannot install or un-install packages and build LAMMPS in a single make command with multiple targets, e.g. `make yes-colloid mpi`. This is because the make procedure creates a list of source files that will be out-of-date for the build if the package configuration changes within the same command. You can include or exclude multiple packages in a single make command, e.g. `make yes-colloid no-manybody`.

---

### 3.6.1 Information for both build systems

Almost all packages can be included or excluded in a LAMMPS build, independent of the other packages. However, some packages include files derived from files in other packages. LAMMPS checks for this and does the right thing. Individual files are only included if their dependencies are already included. Likewise, if a package is excluded, other files dependent on that package are also excluded.

---

**Note:** By default no packages are installed. Prior to August 2018, however, if you downloaded a tarball, 3 packages (KSPACE, MANYBODY, MOLECULE) were pre-installed via the traditional make procedure in the `src` directory. That is no longer the case, so that CMake will build as-is without needing to un-install those packages.

---

### 3.6.2 CMake presets for installing many packages

Instead of specifying all the CMake options via the command-line, CMake allows initializing its settings cache using script files. These are regular CMake files which can manipulate and set CMake variables (which represent selected options), and can also contain control flow constructs for more complex operations.

LAMMPS includes several of these files to define configuration “presets”, similar to the options that exist for the Make based system. Using these files you can enable/disable portions of the available packages in LAMMPS. If you need a custom preset you can take one of them as a starting point and customize it to your needs.

```

cmake -C ../cmake/presets/basic.cmake [OPTIONS] ../cmake # enable just a few core_
↳packages
cmake -C ../cmake/presets/most.cmake [OPTIONS] ../cmake # enable most packages
cmake -C ../cmake/presets/download.cmake [OPTIONS] ../cmake # enable packages which_
↳download sources or potential files
cmake -C ../cmake/presets/nolib.cmake [OPTIONS] ../cmake # disable packages that do_
↳require extra libraries or tools
cmake -C ../cmake/presets/clang.cmake [OPTIONS] ../cmake # change settings to use_
↳the Clang compilers by default
cmake -C ../cmake/presets/gcc.cmake [OPTIONS] ../cmake # change settings to use_
↳the GNU compilers by default
cmake -C ../cmake/presets/intel.cmake [OPTIONS] ../cmake # change settings to use_
↳the Intel compilers by default
cmake -C ../cmake/presets/pgi.cmake [OPTIONS] ../cmake # change settings to use_
↳the PGI compilers by default
cmake -C ../cmake/presets/all_on.cmake [OPTIONS] ../cmake # enable all packages
cmake -C ../cmake/presets/all_off.cmake [OPTIONS] ../cmake # disable all packages
mingw64-cmake -C ../cmake/presets/mingw-cross.cmake [OPTIONS] ../cmake # compile with_
↳MinGW cross compilers

```

Presets that have names starting with “windows” are specifically for compiling LAMMPS *natively on Windows* and presets that have names starting with “kokkos” are specifically for selecting configurations for compiling LAMMPS with *KOKKOS*.

**Note:** Running cmake this way manipulates the CMake settings cache in your current build directory. You can combine multiple presets and options in a single cmake run, or change settings incrementally by running cmake with new flags. If you use a preset for selecting a set of compilers, it will reset all settings from previous CMake runs.

#### Example

```

# build LAMMPS with most commonly used packages, but then remove
# those requiring additional library or tools, but still enable
# GPU package and configure it for using CUDA. You can run.
mkdir build
cd build
cmake -C ../cmake/presets/most.cmake -C ../cmake/presets/nolib.cmake -D PKG_GPU=on -D_
↳GPU_API=cuda ../cmake

# to add another package, say BODY to the previous configuration you can run:
cmake -D PKG_BODY=on .

```

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```
# to reset the package selection from above to the default of no packages
# but leaving all other settings untouched. You can run:
cmake -C ../cmake/presets/all_off.cmake .
```

### 3.6.3 Make shortcuts for installing many packages

The following commands are useful for managing package source files and their installation when building LAMMPS via traditional make. Just type make in lammmps/src to see a one-line summary.

These commands install/un-install sets of packages:

```
make yes-all           # install all packages
make no-all           # check for changes and uninstall all packages
make no-installed     # only check and uninstall installed packages
make yes-basic         # install a few commonly used packages'
make no-basic         # remove a few commonly used packages'
make yes-most         # install most packages w/o libs'
make no-most         # remove most packages w/o libs'
make yes-lib          # install packages that require extra libraries
make no-lib          # uninstall packages that require extra libraries
make yes-ext          # install packages that require external libraries
make no-ext          # uninstall packages that require external libraries
```

which install/un-install various sets of packages. Typing make package will list all the these commands.

**Note:** Installing or un-installing a package for the make based build process works by simply copying files back and forth between the main source directory src and the sub-directories with the package name (e.g. src/KSPACE, src/ATC), so that the files are included or excluded when LAMMPS is built. Only source files in the src folder will be compiled.

The following make commands help manage files that exist in both the src directory and in package sub-directories. You do not normally need to use these commands unless you are editing LAMMPS files or are updating LAMMPS via git.

Type make package-status or make ps to show which packages are currently installed. For those that are installed, it will list any files that are different in the src directory and package sub-directory.

Type make package-installed or make pi to show which packages are currently installed, without listing the status of packages that are not installed.

Type make package-update or make pu to overwrite src files with files from the package sub-directories if the package is installed. It should be used after the checkout has been *updated or changed withy git*, this will only update the files in the package sub-directories, but not the copies in the src folder.

Type make package-overwrite to overwrite files in the package sub-directories with src files.

Type make package-diff to list all differences between pairs of files in both the source directory and the package directory.

## 3.7 Packages with extra build options

When building with some packages, additional steps may be required, in addition to

CMake build	Traditional make
<pre>\$ cmake -D PKG_NAME=yes</pre>	<pre>\$ make yes-name</pre>

as described on the [Build\\_package](#) page.

For a CMake build there may be additional optional or required variables to set. For a build with make, a provided library under the lammps/lib directory may need to be built first. Or an external library may need to exist on your system or be downloaded and built. You may need to tell LAMMPS where it is found on your system.

This is the list of packages that may require additional steps.

<i>ADIOS</i>	<i>ATC</i>	<i>AWPMD</i>	<i>COLVARS</i>	<i>COMPRESS</i>	<i>GPU</i>
<i>H5MD</i>	<i>INTEL</i>	<i>KIM</i>	<i>KOKKOS</i>	<i>LATTE</i>	<i>MACHDYN</i>
<i>MDI</i>	<i>MESONT</i>	<i>ML-HDNNP</i>	<i>ML-IAP</i>	<i>ML-PACE</i>	<i>ML-QUIP</i>
<i>MOLFILE</i>	<i>MSCG</i>	<i>NETCDF</i>	<i>OPENMP</i>	<i>OPT</i>	<i>PLUMED</i>
<i>POEMS</i>	<i>PYTHON</i>	<i>QMMM</i>	<i>SCAFACOS</i>	<i>VORONOI</i>	<i>VTK</i>

---

### 3.7.1 COMPRESS package

To build with this package you must have the [zlib compression library](#) available on your system to build dump styles with a ‘gz’ suffix. There are also styles using the [Zstandard library](#) which have a ‘zstd’ suffix. The zstd library version must be at least 1.4. Older versions use an incompatible API and thus LAMMPS will fail to compile.

---

#### CMake build

If CMake cannot find the zlib library or include files, you can set these variables:

```
-D ZLIB_INCLUDE_DIR=path    # path to zlib.h header file
-D ZLIB_LIBRARY=path        # path to libz.a (.so) file
```

Support for Zstandard compression is auto-detected and for that CMake depends on the [pkg-config](#) tool to identify the necessary flags to compile with this library, so the corresponding `libzstandard.pc` file must be in a folder where `pkg-config` can find it, which may require adding it to the `PKG_CONFIG_PATH` environment variable.

---

#### Traditional make

To include support for Zstandard compression, `-DLAMMPS_ZSTD` must be added to the compiler flags. If make cannot find the libraries, you can edit the file `lib/compress/Makefile.lammps` to specify the paths and library names. This must be done **before** the package is installed.

---

### 3.7.2 GPU package

To build with this package, you must choose options for precision and which GPU hardware to build for. The GPU package currently supports three different types of backends: OpenCL, CUDA and HIP.

#### CMake build

```
-D GPU_API=value           # value = opencl (default) or cuda or hip
-D GPU_PREC=value         # precision setting
                           # value = double or mixed (default) or single
-D HIP_PATH               # path to HIP installation. Must be set if GPU_API=HIP
-D GPU_ARCH=value        # primary GPU hardware choice for GPU_API=cuda
                           # value = sm_XX, see below
                           # default is sm_50
-D HIP_ARCH=value        # primary GPU hardware choice for GPU_API=hip
                           # value depends on selected HIP_PLATFORM
                           # default is 'gfx906' for HIP_PLATFORM=amd and 'sm_50' for HIP_
↳ PLATFORM=nvcc
-D HIP_USE_DEVICE_SORT=value # enables GPU sorting
                           # value = yes (default) or no
-D CUDPP_OPT=value       # use GPU binning on with CUDA (should be off for modern
↳ GPUs)
                           # enables CUDA Performance Primitives, must be "no" for
↳ CUDA_MPS_SUPPORT=yes
                           # value = yes or no (default)
-D CUDA_MPS_SUPPORT=value # enables some tweaks required to run with active nvidia-
↳ cuda-mps daemon
                           # value = yes or no (default)
-D USE_STATIC_OPENCL_LOADER=value # downloads/includes OpenCL ICD loader library, no
↳ local OpenCL headers/libs needed
                           # value = yes (default) or no
```

GPU\_ARCH settings for different GPU hardware is as follows:

- sm\_30 for Kepler (supported since CUDA 5 and until CUDA 10.x)
- sm\_35 or sm\_37 for Kepler (supported since CUDA 5 and until CUDA 11.x)
- sm\_50 or sm\_52 for Maxwell (supported since CUDA 6)
- sm\_60 or sm\_61 for Pascal (supported since CUDA 8)
- sm\_70 for Volta (supported since CUDA 9)
- sm\_75 for Turing (supported since CUDA 10)
- sm\_80 for Ampere (supported since CUDA 11)

A more detailed list can be found, for example, at [Wikipedia's CUDA article](#)

CMake can detect which version of the CUDA toolkit is used and thus will try to include support for **all** major GPU architectures supported by this toolkit. Thus the GPU\_ARCH setting is merely an optimization, to have code for the preferred GPU architecture directly included rather than having to wait for the JIT compiler of the CUDA driver to translate it.

When compiling for CUDA or HIP with CUDA, version 8.0 or later of the CUDA toolkit is required and a GPU architecture of Kepler or later, which must *also* be supported by the CUDA toolkit in use **and** the CUDA driver in use.

When compiling for OpenCL, OpenCL version 1.2 or later is required and the GPU must be supported by the GPU driver and OpenCL runtime bundled with the driver.

When building with CMake, you **must NOT** build the GPU library in `lib/gpu` using the traditional build procedure. CMake will detect files generated by that process and will terminate with an error and a suggestion for how to remove them.

If you are compiling for OpenCL, the default setting is to download, build, and link with a static OpenCL ICD loader library and standard OpenCL headers. This way no local OpenCL development headers or library needs to be present and only OpenCL compatible drivers need to be installed to use OpenCL. If this is not desired, you can set `USE_STATIC_OPENCL_LOADER` to `no`.

If you are compiling with HIP, note that before running CMake you will have to set appropriate environment variables. Some variables such as `HCC_AMDGPU_TARGET` (for ROCm <= 4.0) or `CUDA_PATH` are necessary for `hipcc` and the linker to work correctly.

Using CHIP-SPV implementation of HIP is now supported. It allows one to run HIP code on Intel GPUs via the OpenCL or Level Zero backends. To use CHIP-SPV, you must set `-DHIP_USE_DEVICE_SORT=OFF` in your CMake command line as CHIP-SPV does not yet support `hipCUB`. The use of HIP for Intel GPUs is still experimental so you should only use this option in preparations to run on Aurora system at ANL.

```
# AMDGPU target (ROCm <= 4.0)
export HIP_PLATFORM=hcc
export HIP_PATH=/path/to/HIP/install
export HCC_AMDGPU_TARGET=gfx906
cmake -D PKG_GPU=on -D GPU_API=HIP -D HIP_ARCH=gfx906 -D CMAKE_CXX_COMPILER=hipcc ..
make -j 4
```

```
# AMDGPU target (ROCm >= 4.1)
export HIP_PLATFORM=amd
export HIP_PATH=/path/to/HIP/install
cmake -D PKG_GPU=on -D GPU_API=HIP -D HIP_ARCH=gfx906 -D CMAKE_CXX_COMPILER=hipcc ..
make -j 4
```

```
# CUDA target (not recommended, use GPU_ARCH=cuda)
# !!! DO NOT set CMAKE_CXX_COMPILER !!!
export HIP_PLATFORM=nvcc
export HIP_PATH=/path/to/HIP/install
export CUDA_PATH=/usr/local/cuda
cmake -D PKG_GPU=on -D GPU_API=HIP -D HIP_ARCH=sm_70 ..
make -j 4
```

```
# SPIR-V target (Intel GPUs)
export HIP_PLATFORM=spirv
export HIP_PATH=/path/to/HIP/install
export CMAKE_CXX_COMPILER=<hipcc/clang++>
cmake -D PKG_GPU=on -D GPU_API=HIP ..
make -j 4
```

## Traditional make

Before building LAMMPS, you must build the GPU library in `lib/gpu`. You can do this manually if you prefer; follow the instructions in `lib/gpu/README`. Note that the GPU library uses MPI calls, so you must use the same MPI library (or the STUBS library) settings as the main LAMMPS code. This also applies to the `-DLAMMPS_BIGBIG`, `-DLAMMPS_SMALLBIG`, or `-DLAMMPS_SMALLSMALL` settings in whichever Makefile you use.

You can also build the library in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/gpu/Install.py` script with the specified args:

```
$ make lib-gpu                # print help message
$ make lib-gpu args="-b"      # build GPU library with default Makefile.linux
$ make lib-gpu args="-m xk7 -p single -o xk7.single" # create new Makefile.xk7.single,
↳altered for single-precision
$ make lib-gpu args="-m mpi -a sm_60 -p mixed -b" # build GPU library with mixed,
↳precision and P100 using other settings in Makefile.mpi
```

Note that this procedure starts with a `Makefile.machine` in `lib/gpu`, as specified by the “-m” switch. For your convenience, machine makefiles for “mpi” and “serial” are provided, which have the same settings as the corresponding machine makefiles in the main LAMMPS source folder. In addition you can alter 4 important settings in the `Makefile.machine` you start from via the corresponding `-c`, `-a`, `-p`, `-e` switches (as in the examples above), and also save a copy of the new `Makefile` if desired:

- `CUDA_HOME` = where NVIDIA CUDA software is installed on your system
- `CUDA_ARCH` = `sm_XX`, what GPU hardware you have, same as CMake `GPU_ARCH` above
- `CUDA_PRECISION` = precision (double, mixed, single)
- `EXTRAMAKE` = which `Makefile.lammps.*` file to copy to `Makefile.lammps`

The file `Makefile.cuda` is set up to include support for multiple GPU architectures as supported by the CUDA toolkit in use. This is done through using the “-gencode” flag, which can be used multiple times and thus support all GPU architectures supported by your CUDA compiler.

To enable GPU binning via CUDA performance primitives set the Makefile variable `CUDPP_OPT = -DUSE_CUDPP -Icudpp_mini`. This should **not** be used with most modern GPUs.

To support the CUDA multiprocessor server you can set the define `-DCUDA_PROXY`. Please note that in this case you must **not** use the CUDA performance primitives and thus set the variable `CUDPP_OPT` to empty.

If the library build is successful, 3 files should be created: `lib/gpu/libgpu.a`, `lib/gpu/nvc_get_devices`, and `lib/gpu/Makefile.lammps`. The latter has settings that enable LAMMPS to link with CUDA libraries. If the settings in `Makefile.lammps` for your machine are not correct, the LAMMPS build will fail, and `lib/gpu/Makefile.lammps` may need to be edited.

---

**Note:** If you re-build the GPU library in `lib/gpu`, you should always un-install the GPU package in `lammps/src`, then re-install it and re-build LAMMPS. This is because the compilation of files in the GPU package uses the library settings from the `lib/gpu/Makefile.machine` used to build the GPU library.

---

### 3.7.3 KIM package

To build with this package, the KIM library with API v2 must be downloaded and built on your system. It must include the KIM models that you want to use with LAMMPS.

If you would like to use the *kim query* command, you also need to have libcurl installed with the matching development headers and the curl-config tool.

If you would like to use the *kim property* command, you need to build LAMMPS with the PYTHON package installed and linked to Python 3.6 or later. See the *PYTHON package build info* for more details on this. After successfully building LAMMPS with Python, you also need to install the *kim-property* Python package, which can be easily done using *pip* as `pip install kim-property`, or from the *conda-forge* channel as `conda install kim-property` if LAMMPS is built in Conda. More detailed information is available at: [kim-property installation](#).

In addition to installing the KIM API, it is also necessary to install the library of KIM models (interatomic potentials). See [Obtaining KIM Models](#) to learn how to install a pre-build binary of the OpenKIM Repository of Models. See the list of all KIM models here: <https://openkim.org/browse/models>

(Also note that when downloading and installing from source the KIM API library with all its models, may take a long time (tens of minutes to hours) to build. Of course you only need to do that once.)

---

#### CMake build

```
-D DOWNLOAD_KIM=value           # download OpenKIM API v2 for build, value = no
↳no (default) or yes
-D LMP_DEBUG_CURL=value         # set libcurl verbose mode on/off, value = off
↳(default) or on
-D LMP_NO_SSL_CHECK=value       # tell libcurl to not verify the peer, value = no
↳no (default) or yes
-D KIM_EXTRA_UNITTESTS=value    # enables extra unit tests, value = no
↳(default) or yes
```

If `DOWNLOAD_KIM` is set to *yes* (or *on*), the KIM API library will be downloaded and built inside the CMake build directory. If the KIM library is already installed on your system (in a location where CMake cannot find it), you may need to set the `PKG_CONFIG_PATH` environment variable so that `libkim-api` can be found, or run the command `source kim-api-activate`.

Extra unit tests can only be available if they are explicitly requested (`KIM_EXTRA_UNITTESTS` is set to *yes* (or *on*)) and the prerequisites are met. See *KIM Extra unit tests* for more details on this.

---

#### Traditional make

You can download and build the KIM library manually if you prefer; follow the instructions in `lib/kim/README`. You can also do this in one step from the `lammps/src` directory, using a command like these, which simply invoke the `lib/kim/Install.py` script with the specified args.

```
$ make lib-kim                  # print help message
$ make lib-kim args="-b "       # (re-)install KIM API lib with only example models
$ make lib-kim args="-b -a Glue_Ercolessi_Adams_Al__MO_324507536345_001" #_
↳ditto plus one model
$ make lib-kim args="-b -a everything" # install KIM API lib with all_
↳models
$ make lib-kim args="-n -a EAM_Dynamo_Ackland_W__MO_141627196590_002" #_
↳add one model or model driver
```

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```
$ make lib-kim args="-p /usr/local" # use an existing KIM API installation at
↳ the provided location
$ make lib-kim args="-p /usr/local -a EAM_Dynamo_Ackland_W__MO_141627196590_002
↳ " # ditto but add one model or driver
```

When using the “-b” option, the KIM library is built using its native cmake build system. The lib/kim/Install.py script supports a CMAKE environment variable if the cmake executable is named other than cmake on your system. Additional environment variables may be provided on the command line for use by cmake. For example, to use the cmake3 executable and tell it to use the gnu version 11 compilers to build KIM, one could use the following command line.

```
$ CMAKE=cmake3 CXX=g++-11 CC=gcc-11 FC=gfortran-11 make lib-kim args="-b " #
↳ (re-)install KIM API lib using cmake3 and gnu v11 compilers with only
↳ example models
```

Settings for debugging OpenKIM web queries discussed below need to be applied by adding them to the LMP\_INC variable through editing the Makefile.machine you are using. For example:

```
LMP_INC = -DLMP_NO_SSL_CHECK
```

## Debugging OpenKIM web queries in LAMMPS

If LMP\_DEBUG\_CURL is set, the libcurl verbose mode will be turned on, and any libcurl calls within the KIM web query display a lot of information about libcurl operations. You hardly ever want this set in production use, you will almost always want this when you debug or report problems.

The libcurl library performs peer SSL certificate verification by default. This verification is done using a CA certificate store that the SSL library can use to make sure the peer’s server certificate is valid. If SSL reports an error (“certificate verify failed”) during the handshake and thus refuses further communicate with that server, you can set LMP\_NO\_SSL\_CHECK to override that behavior. When LAMMPS is compiled with LMP\_NO\_SSL\_CHECK set, libcurl does not verify the peer and connection attempts will succeed regardless of the names in the certificate. This option is insecure. As an alternative, you can specify your own CA cert path by setting the environment variable CURL\_CA\_BUNDLE to the path of your choice. A call to the KIM web query would get this value from the environment variable.

## KIM Extra unit tests (CMake only)

During development, testing, or debugging, if *unit testing* is enabled in LAMMPS, one can also enable extra tests on *KIM commands* by setting the KIM\_EXTRA\_UNITTESTS to *yes* (or *on*).

Enabling the extra unit tests have some requirements,

- It requires to have internet access.
- It requires to have libcurl installed with the matching development headers and the curl-config tool.
- It requires to build LAMMPS with the PYTHON package installed and linked to Python 3.6 or later. See the *PYTHON package build info* for more details on this.
- It requires to have kim-property Python package installed, which can be easily done using *pip* as `pip install kim-property`, or from the *conda-forge* channel as `conda install kim-property` if LAMMPS is built in Conda. More detailed information is available at: [kim-property installation](#).
- It is also necessary to install `EAM_Dynamo_MendeleevAckland_2007v3_Zr__MO_004835508849_000`, `EAM_Dynamo_ErcolessiAdams_1994_A1__MO_123629422045_005`, and `LennardJones612_UniversalShifted__MO_9592`

KIM models. See [Obtaining KIM Models](#) to learn how to install a pre-built binary of the OpenKIM Repository of Models or see [Installing KIM Models](#) to learn how to install the specific KIM models.

### 3.7.4 KOKKOS package

Using the KOKKOS package requires choosing several settings. You have to select whether you want to compile with parallelization on the host and whether you want to include offloading of calculations to a device (e.g. a GPU). The default setting is to have no host parallelization and no device offloading. In addition, you can select the hardware architecture to select the instruction set. Since most hardware is backward compatible, you may choose settings for an older architecture to have an executable that will run on this and newer architectures.

**Note:** If you run Kokkos on a different GPU architecture than what LAMMPS was compiled with, there will be a delay during device initialization while the just-in-time compiler is recompiling all GPU kernels for the new hardware. This is, however, only supported for GPUs of the **same** major hardware version and different minor hardware versions, e.g. 5.0 and 5.2 but not 5.2 and 6.0. LAMMPS will abort with an error message indicating a mismatch, if that happens.

The settings discussed below have been tested with LAMMPS and are confirmed to work. Kokkos is an active project with ongoing improvements and projects working on including support for additional architectures. More information on Kokkos can be found on the [Kokkos GitHub project](#).

#### Available Architecture settings

These are the possible choices for the Kokkos architecture ID. They must be specified in uppercase.

Arch-ID	HOST or GPU	Description
AMDAVX	HOST	AMD 64-bit x86 CPU (AVX 1)
ZEN	HOST	AMD Zen class CPU (AVX 2)
ZEN2	HOST	AMD Zen2 class CPU (AVX 2)
ZEN3	HOST	AMD Zen3 class CPU (AVX 2)
ARMV80	HOST	ARMv8.0 Compatible CPU
ARMV81	HOST	ARMv8.1 Compatible CPU
ARMV8_THUNDERX	HOST	ARMv8 Cavium ThunderX CPU
ARMV8_THUNDERX2	HOST	ARMv8 Cavium ThunderX2 CPU
A64FX	HOST	ARMv8.2 with SVE Support
WSM	HOST	Intel Westmere CPU (SSE 4.2)
SNB	HOST	Intel Sandy/Ivy Bridge CPU (AVX 1)
HSW	HOST	Intel Haswell CPU (AVX 2)
BDW	HOST	Intel Broadwell Xeon E-class CPU (AVX 2 + transactional mem)
SKX	HOST	Intel Sky Lake Xeon E-class HPC CPU (AVX512 + transactional mem)
KNC	HOST	Intel Knights Corner Xeon Phi
KNL	HOST	Intel Knights Landing Xeon Phi
BGQ	HOST	IBM Blue Gene/Q CPU
POWER7	HOST	IBM POWER7 CPU
POWER8	HOST	IBM POWER8 CPU
POWER9	HOST	IBM POWER9 CPU
KEPLER30	GPU	NVIDIA Kepler generation CC 3.0 GPU
KEPLER32	GPU	NVIDIA Kepler generation CC 3.2 GPU
KEPLER35	GPU	NVIDIA Kepler generation CC 3.5 GPU

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Table 1 – continued from previous page

KEPLER37	GPU	NVIDIA Kepler generation CC 3.7 GPU
MAXWELL50	GPU	NVIDIA Maxwell generation CC 5.0 GPU
MAXWELL52	GPU	NVIDIA Maxwell generation CC 5.2 GPU
MAXWELL53	GPU	NVIDIA Maxwell generation CC 5.3 GPU
PASCAL60	GPU	NVIDIA Pascal generation CC 6.0 GPU
PASCAL61	GPU	NVIDIA Pascal generation CC 6.1 GPU
VOLTA70	GPU	NVIDIA Volta generation CC 7.0 GPU
VOLTA72	GPU	NVIDIA Volta generation CC 7.2 GPU
TURING75	GPU	NVIDIA Turing generation CC 7.5 GPU
AMPERE80	GPU	NVIDIA Ampere generation CC 8.0 GPU
AMPERE86	GPU	NVIDIA Ampere generation CC 8.6 GPU
VEGA900	GPU	AMD GPU MI25 GFX900
VEGA906	GPU	AMD GPU MI50/MI60 GFX906
VEGA908	GPU	AMD GPU MI100 GFX908
VEGA90A	GPU	AMD GPU
INTEL_DG1	GPU	Intel Iris XeMAX GPU
INTEL_GEN9	GPU	Intel GPU Gen9
INTEL_GEN11	GPU	Intel GPU Gen11
INTEL_GEN12LP	GPU	Intel GPU Gen12LP
INTEL_XEHP	GPU	Intel GPUs Xe-HP

This list was last updated for version 3.5.0 of the Kokkos library.

---

#### Basic CMake build settings:

For multicore CPUs using OpenMP, set these 2 variables.

```
-D Kokkos_ARCH_HOSTARCH=yes # HOSTARCH = HOST from list above
-D Kokkos_ENABLE_OPENMP=yes
-D BUILD_OMP=yes
```

Please note that enabling OpenMP for KOKKOS requires that OpenMP is also *enabled for the rest of LAMMPS*.

For Intel KNLs using OpenMP, set these variables:

```
-D Kokkos_ARCH_KNL=yes
-D Kokkos_ENABLE_OPENMP=yes
```

For NVIDIA GPUs using CUDA, set these variables:

```
-D Kokkos_ARCH_HOSTARCH=yes # HOSTARCH = HOST from list above
-D Kokkos_ARCH_GPUARCH=yes # GPUARCH = GPU from list above
-D Kokkos_ENABLE_CUDA=yes
-D Kokkos_ENABLE_OPENMP=yes
-D CMAKE_CXX_COMPILER=wrapper # wrapper = full path to Cuda nvcc wrapper
```

This will also enable executing FFTs on the GPU, either via the internal KISSFFT library, or - by preference - with the cuFFT library bundled with the CUDA toolkit, depending on whether CMake can identify its location. The *wrapper* value for CMAKE\_CXX\_COMPILER variable is the path to the CUDA nvcc compiler wrapper provided in the Kokkos library: `lib/kokkos/bin/nvcc_wrapper`. The setting should include the full path name to the wrapper, e.g.

```
-D CMAKE_CXX_COMPILER=${HOME}/lammps/lib/kokkos/bin/nvcc_wrapper
```

For AMD or NVIDIA GPUs using HIP, set these variables:

```
-D Kokkos_ARCH_HOSTARCH=yes # HOSTARCH = HOST from list above
-D Kokkos_ARCH_GPUARCH=yes # GPUARCH = GPU from list above
-D Kokkos_ENABLE_HIP=yes
-D Kokkos_ENABLE_OPENMP=yes
```

This will enable FFTs on the GPU, either by the internal KISSFFT library or with the hipFFT wrapper library, which will call out to the platform-appropriate vendor library: rocFFT on AMD GPUs or cuFFT on NVIDIA GPUs.

To simplify compilation, five preset files are included in the `cmake/presets` folder, `kokkos-serial.cmake`, `kokkos-openmp.cmake`, `kokkos-cuda.cmake`, `kokkos-hip.cmake`, and `kokkos-sycl.cmake`. They will enable the KOKKOS package and enable some hardware choice. So to compile with CUDA device parallelization (for GPUs with CC 5.0 and up) with some common packages enabled, you can do the following:

```
mkdir build-kokkos-cuda
cd build-kokkos-cuda
cmake -C ../cmake/presets/basic.cmake -C ../cmake/presets/kokkos-cuda.cmake ../
↪cmake
cmake --build .
```

---

#### Basic traditional make settings:

Choose which hardware to support in `Makefile.machine` via `KOKKOS_DEVICES` and `KOKKOS_ARCH` settings. See the `src/MAKE/OPTIONS/Makefile.kokkos*` files for examples.

For multicore CPUs using OpenMP:

```
KOKKOS_DEVICES = OpenMP
KOKKOS_ARCH = HOSTARCH # HOSTARCH = HOST from list above
```

For Intel KNLs using OpenMP:

```
KOKKOS_DEVICES = OpenMP
KOKKOS_ARCH = KNL
```

For NVIDIA GPUs using CUDA:

```
KOKKOS_DEVICES = Cuda
KOKKOS_ARCH = HOSTARCH,GPUARCH # HOSTARCH = HOST from list above that is,
↪hosting the GPU
KOKKOS_CUDA_OPTIONS = "enable_lambda"
# GPUARCH = GPU from list above
FFT_INC = -DFFT_CUFFT # enable use of cuFFT (optional)
FFT_LIB = -lcufft # link to cuFFT library
```

For GPUs, you also need the following lines in your `Makefile.machine` before the `CC` line is defined. They tell `mpicxx` to use an `nvcc` compiler wrapper, which will use `nvcc` for compiling CUDA files and a C++ compiler for non-Kokkos, non-CUDA files.

```
# For OpenMPI
KOKKOS_ABSOLUTE_PATH = $(shell cd $(KOKKOS_PATH); pwd)
export OMPI_CXX = $(KOKKOS_ABSOLUTE_PATH)/config/nvcc_wrapper
CC = mpicxx
```

```
# For MPICH and derivatives
KOKKOS_ABSOLUTE_PATH = $(shell cd $(KOKKOS_PATH); pwd)
CC = mpicxx -cxx=$(KOKKOS_ABSOLUTE_PATH)/config/nvcc_wrapper
```

For AMD or NVIDIA GPUs using HIP:

```
KOKKOS_DEVICES = HIP
KOKKOS_ARCH = HOSTARCH,GPUARCH # HOSTARCH = HOST from list above that is
↪hosting the GPU
                                # GPUARCH = GPU from list above
FFT_INC = -DFFT_HIPFFT          # enable use of hipFFT (optional)
FFT_LIB = -lhipfft              # link to hipFFT library
```

### Advanced KOKKOS compilation settings

There are other allowed options when building with the KOKKOS package that can improve performance or assist in debugging or profiling. Below are some examples that may be useful in combination with LAMMPS. For the full list (which keeps changing as the Kokkos package itself evolves), please consult the Kokkos library documentation.

As alternative to using multi-threading via OpenMP (`-DKokkos_ENABLE_OPENMP=on` or `KOKKOS_DEVICES=OpenMP`) it is also possible to use Posix threads directly (`-DKokkos_ENABLE_PTHREAD=on` or `KOKKOS_DEVICES=Pthread`). While binding of threads to individual or groups of CPU cores is managed in OpenMP with environment variables, you need assistance from either the “hwloc” or “libnuma” library for the Pthread thread parallelization option. To enable use with CMake: `-DKokkos_ENABLE_HWLOC=on` or `-DKokkos_ENABLE_LIBNUMA=on`; and with conventional make: `KOKKOS_USE_TPLS=hwloc` or `KOKKOS_USE_TPLS=libnuma`.

The CMake option `-DKokkos_ENABLE_LIBRT=on` or the makefile setting `KOKKOS_USE_TPLS=librt` enables the use of a more accurate timer mechanism on many Unix-like platforms for internal profiling.

The CMake option `-DKokkos_ENABLE_DEBUG=on` or the makefile setting `KOKKOS_DEBUG=yes` enables printing of runtime debugging information that can be useful. It also enables runtime bounds checking on Kokkos data structures. As to be expected, enabling this option will negatively impact the performance and thus is only recommended when developing a Kokkos-enabled style in LAMMPS.

The CMake option `-DKokkos_ENABLE_CUDA_UVM=on` or the makefile setting `KOKKOS_CUDA_OPTIONS=enable_lambda,force_uvm` enables the use of CUDA “Unified Virtual Memory” (UVM) in Kokkos. UVM allows to transparently use RAM on the host to supplement the memory used on the GPU (with some performance penalty) and thus enables running larger problems that would otherwise not fit into the RAM on the GPU.

Please note, that the LAMMPS KOKKOS package must **always** be compiled with the `enable_lambda` option when using GPUs. The CMake configuration will thus always enable it.

### 3.7.5 LATTE package

To build with this package, you must download and build the LATTE library.

---

#### CMake build

```
-D DOWNLOAD_LATTE=value      # download LATTE for build, value = no (default)
↪ or yes
-D LATTE_LIBRARY=path        # LATTE library file (only needed if a custom
↪ location)
-D USE_INTERNAL_LINALG=value # Use the internal linear algebra library instead
↪ of LAPACK
                               # value = no (default) or yes
```

If `DOWNLOAD_LATTE` is set, the LATTE library will be downloaded and built inside the CMake build directory. If the LATTE library is already on your system (in a location CMake cannot find it), `LATTE_LIBRARY` is the filename (plus path) of the LATTE library file, not the directory the library file is in.

The LATTE library requires LAPACK (and BLAS) and CMake can identify their locations and pass that info to the LATTE build script. But on some systems this triggers a (current) limitation of CMake and the configuration will fail. Try enabling `USE_INTERNAL_LINALG` in those cases to use the bundled linear algebra library and work around the limitation.

---

#### Traditional make

You can download and build the LATTE library manually if you prefer; follow the instructions in `lib/latte/README`. You can also do it in one step from the `lammops/src` dir, using a command like these, which simply invokes the `lib/latte/Install.py` script with the specified args:

```
$ make lib-latte                # print help message
$ make lib-latte args="-b"      # download and build in lib/latte/
↪ LATTE-master
$ make lib-latte args="-p $HOME/latte" # use existing LATTE installation in
↪ $HOME/latte
$ make lib-latte args="-b -m gfortran" # download and build in lib/latte and
↪ copy Makefile.lammops.gfortran to
↪ Makefile.lammops
```

Note that 3 symbolic (soft) links, `includelink` and `liblink` and `filelink.o`, are created in `lib/latte` to point to required folders and files in the LATTE home directory. When LAMMPS itself is built it will use these links. You should also check that the `Makefile.lammops` file you create is appropriate for the compiler you use on your system to build LATTE.

---

### 3.7.6 ML-IAP package

Building the ML-IAP package requires including the *ML-SNAP* package. There will be an error message if this requirement is not satisfied. Using the *mliappy* model also requires enabling Python support, which in turn requires to include the *PYTHON* package **and** requires to have the *cython* software installed and with it a working *cythonize* command. This feature requires compiling LAMMPS with Python version 3.6 or later.

---

#### CMake build

```
-D MLIAP_ENABLE_PYTHON=value # enable mliappy model (default is autodetect)
```

Without this setting, CMake will check whether it can find a suitable Python version and the *cythonize* command and choose the default accordingly. During the build procedure the provided .pyx file(s) will be automatically translated to C++ code and compiled. Please do **not** run *cythonize* manually in the *src/ML-IAP* folder, as that can lead to compilation errors if Python support is not enabled. If you did it by accident, please remove the generated .cpp and .h files.

---

#### Traditional make

The build uses the *lib/python/Makefile.mliap\_python* file in the compile/link process to add a rule to update the files generated by the *cythonize* command in case the corresponding .pyx file(s) were modified. You may need to modify *lib/python/Makefile.lammps* if the LAMMPS build fails.

To enable building the ML-IAP package with Python support enabled, you need to add *-DMLIAP\_PYTHON* to the *LMP\_INC* variable in your machine makefile. You may have to manually run the *cythonize* command on .pyx file(s) in the *src* folder, if this is not automatically done during installing the ML-IAP package. Please do **not** run *cythonize* in the *src/ML-IAP* folder, as that can lead to compilation errors if Python support is not enabled. If you did this by accident, please remove the generated .cpp and .h files.

---

### 3.7.7 MSCG package

To build with this package, you must download and build the MS-CG library. Building the MS-CG library requires that the GSL (GNU Scientific Library) headers and libraries are installed on your machine. See the *lib/mscg/README* and *MSCG/Install* files for more details.

---

#### CMake build

```
-D DOWNLOAD_MSCG=value # download MSCG for build, value = no (default) or ↵
↵yes
-D MSCG_LIBRARY=path # MSCG library file (only needed if a custom ↵
↵location)
-D MSCG_INCLUDE_DIR=path # MSCG include directory (only needed if a custom ↵
↵location)
```

If *DOWNLOAD\_MSCG* is set, the MSCG library will be downloaded and built inside the CMake build directory. If the MSCG library is already on your system (in a location CMake cannot find it), *MSCG\_LIBRARY* is the filename (plus path) of the MSCG library file, not the directory the library file is in. *MSCG\_INCLUDE\_DIR* is the directory the MSCG include file is in.

---

---

**Traditional make**

You can download and build the MS-CG library manually if you prefer; follow the instructions in `lib/mscg/README`. You can also do it in one step from the `lammops/src` dir, using a command like these, which simply invoke the `lib/mscg/Install.py` script with the specified args:

```
$ make lib-mscg           # print help message
$ make lib-mscg args="-b -m serial" # download and build in lib/mscg/MSCG-
↪release-master
                               # with the settings compatible with
↪"make serial"
$ make lib-mscg args="-b -m mpi"   # download and build in lib/mscg/MSCG-
↪release-master
                               # with the settings compatible with
↪"make mpi"
$ make lib-mscg args="-p /usr/local/mscg-release" # use the existing MS-CG_
↪installation in /usr/local/mscg-release
```

Note that 2 symbolic (soft) links, `includelink` and `liblink`, will be created in `lib/mscg` to point to the MS-CG `src/installation` dir. When LAMMPS is built in `src` it will use these links. You should not need to edit the `lib/mscg/Makefile.lammops` file.

---

---

### 3.7.8 OPT package

---

**CMake build**

No additional settings are needed besides `-D PKG_OPT=yes`

---

---

**Traditional make**

The compiler flag `-restrict` must be used to build LAMMPS with the OPT package when using Intel compilers. It should be added to the `CCFLAGS` line of your `Makefile.machine`. See `src/MAKE/OPTIONS/Makefile.opt` for an example.

---

---

### 3.7.9 POEMS package

---

**CMake build**

No additional settings are needed besides `-D PKG_OPT=yes`

---

---

**Traditional make**

Before building LAMMPS, you must build the POEMS library in `lib/poems`. You can do this manually if you prefer; follow the instructions in `lib/poems/README`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/poems/Install.py` script with the specified args:

```
$ make lib-poems           # print help message
$ make lib-poems args="-m serial" # build with GNU g++ compiler (settings as
↪with "make serial")
$ make lib-poems args="-m mpi"   # build with default MPI C++ compiler
↪(settings as with "make mpi")
$ make lib-poems args="-m icc"   # build with Intel icc compiler
```

The build should produce two files: `lib/poems/libpoems.a` and `lib/poems/Makefile.lammps`. The latter is copied from an existing `Makefile.lammps.*` and has settings needed to build LAMMPS with the POEMS library (though typically the settings are just blank). If necessary, you can edit/create a new `lib/poems/Makefile.machine` file for your system, which should define an `EXTRAMAKE` variable to specify a corresponding `Makefile.lammps.machine` file.

---

### 3.7.10 PYTHON package

Building with the PYTHON package requires you have a the Python development headers and library available on your system, which needs to be a Python 2.7 version or a Python 3.x version. Since support for Python 2.x has ended, using Python 3.x is strongly recommended. See `lib/python/README` for additional details.

---

#### CMake build

```
-D PYTHON_EXECUTABLE=path # path to Python executable to use
```

Without this setting, CMake will guess the default Python version on your system. To use a different Python version, you can either create a virtualenv, activate it and then run `cmake`. Or you can set the `PYTHON_EXECUTABLE` variable to specify which Python interpreter should be used. Note note that you will also need to have the development headers installed for this version, e.g. `python2-devel`.

---

---

#### Traditional make

The build uses the `lib/python/Makefile.lammps` file in the compile/link process to find Python. You should only need to create a new `Makefile.lammps.*` file (and copy it to `Makefile.lammps`) if the LAMMPS build fails.

---

### 3.7.11 VORONOI package

To build with this package, you must download and build the [Voro++ library](#) or install a binary package provided by your operating system.

---

#### CMake build

```
-D DOWNLOAD_VORO=value      # download Voro++ for build, value = no (default) or ↵  
↵yes  
-D VORO_LIBRARY=path        # Voro++ library file (only needed if at custom ↵  
↵location)  
-D VORO_INCLUDE_DIR=path    # Voro++ include directory (only needed if at custom ↵  
↵location)
```

If `DOWNLOAD_VORO` is set, the Voro++ library will be downloaded and built inside the CMake build directory. If the Voro++ library is already on your system (in a location CMake cannot find it), `VORO_LIBRARY` is the filename (plus path) of the Voro++ library file, not the directory the library file is in. `VORO_INCLUDE_DIR` is the directory the Voro++ include file is in.

---

---

#### Traditional make

You can download and build the Voro++ library manually if you prefer; follow the instructions in `lib/voronoi/README`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/voronoi/Install.py` script with the specified args:

```
$ make lib-voronoi          # print help message  
$ make lib-voronoi args="-b" # download and build the default ↵  
↵version in lib/voronoi/voro++-<version>  
$ make lib-voronoi args="-p $HOME/voro++" # use existing Voro++ installation ↵  
↵in $HOME/voro++  
$ make lib-voronoi args="-b -v voro++0.4.6" # download and build the 0.4.6 ↵  
↵version in lib/voronoi/voro++-0.4.6
```

Note that 2 symbolic (soft) links, `includelink` and `liblink`, are created in `lib/voronoi` to point to the Voro++ source dir. When LAMMPS builds in `src` it will use these links. You should not need to edit the `lib/voronoi/Makefile.lammps` file.

---

### 3.7.12 ADIOS package

The ADIOS package requires the [ADIOS I/O library](#), version 2.3.1 or newer. Make sure that you have ADIOS built either with or without MPI to match if you build LAMMPS with or without MPI. ADIOS compilation settings for LAMMPS are automatically detected, if the `PATH` and `LD_LIBRARY_PATH` environment variables have been updated for the local ADIOS installation and the instructions below are followed for the respective build systems.

---

#### CMake build

```
-D ADIOS2_DIR=path          # path is where ADIOS 2.x is installed  
-D PKG_ADIOS=yes
```



---

**Traditional make**

Turn on the ADIOS package before building LAMMPS. If the ADIOS 2.x software is installed in PATH, there is nothing else to do:

```
$ make yes-adios
```

otherwise, set ADIOS2\_DIR environment variable when turning on the package:

```
$ ADIOS2_DIR=path make yes-adios # path is where ADIOS 2.x is installed
```

---

### 3.7.13 ATC package

The ATC package requires the MANYBODY package also be installed.

---

**CMake build**

No additional settings are needed besides `-D PKG_ATC=yes` and `-D PKG_MANYBODY=yes`.

---

---

**Traditional make**

Before building LAMMPS, you must build the ATC library in `lib/atc`. You can do this manually if you prefer; follow the instructions in `lib/atc/README`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/atc/Install.py` script with the specified args:

```
$ make lib-atc # print help message
$ make lib-atc args="-m serial" # build with GNU g++ compiler and MPI
↪(settings as with "make serial")
$ make lib-atc args="-m mpi" # build with default MPI compiler
↪(settings as with "make mpi")
$ make lib-atc args="-m icc" # build with Intel icc compiler
```

The build should produce two files: `lib/atc/libatc.a` and `lib/atc/Makefile.lammps`. The latter is copied from an existing `Makefile.lammps.*` and has settings needed to build LAMMPS with the ATC library. If necessary, you can edit/create a new `lib/atc/Makefile.machine` file for your system, which should define an `EXTRAMAKE` variable to specify a corresponding `Makefile.lammps.<machine>` file.

Note that the `Makefile.lammps` file has settings for the BLAS and LAPACK linear algebra libraries. As explained in `lib/atc/README` these can either exist on your system, or you can use the files provided in `lib/linalg`. In the latter case you also need to build the library in `lib/linalg` with a command like these:

```
$ make lib-linalg # print help message
$ make lib-linalg args="-m serial" # build with GNU Fortran compiler
↪(settings as with "make serial")
$ make lib-linalg args="-m mpi" # build with default MPI Fortran
```

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```
↪ compiler (settings as with "make mpi")
$ make lib-linalg args="-m gfortran" # build with GNU Fortran compiler
```

---

### 3.7.14 AWPMD package

---

#### CMake build

No additional settings are needed besides `-D PKG_AQPM=yes`.

---

#### Traditional make

Before building LAMMPS, you must build the AWPMD library in `lib/awpmd`. You can do this manually if you prefer; follow the instructions in `lib/awpmd/README`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/awpmd/Install.py` script with the specified args:

```
$ make lib-awpmd # print help message
$ make lib-awpmd args="-m serial" # build with GNU g++ compiler and MPI STUBS
↪ (settings as with "make serial")
$ make lib-awpmd args="-m mpi" # build with default MPI compiler (settings
↪ as with "make mpi")
$ make lib-awpmd args="-m icc" # build with Intel icc compiler
```

The build should produce two files: `lib/awpmd/libawpmd.a` and `lib/awpmd/Makefile.lammps`. The latter is copied from an existing `Makefile.lammps.*` and has settings needed to build LAMMPS with the AWPMD library. If necessary, you can edit/create a new `lib/awpmd/Makefile.machine` file for your system, which should define an `EXTRAMAKE` variable to specify a corresponding `Makefile.lammps.<machine>` file.

Note that the `Makefile.lammps` file has settings for the BLAS and LAPACK linear algebra libraries. As explained in `lib/awpmd/README` these can either exist on your system, or you can use the files provided in `lib/linalg`. In the latter case you also need to build the library in `lib/linalg` with a command like these:

```
$ make lib-linalg # print help message
$ make lib-linalg args="-m serial" # build with GNU Fortran compiler
↪ (settings as with "make serial")
$ make lib-linalg args="-m mpi" # build with default MPI Fortran
↪ compiler (settings as with "make mpi")
$ make lib-linalg args="-m gfortran" # build with GNU Fortran compiler
```

---

### 3.7.15 COLVARS package

This package includes the [Colvars library](#) into the LAMMPS distribution, which can be built for the most part with all major versions of the C++ language.

---

#### CMake build

This is the recommended build procedure for using Colvars in LAMMPS. No additional settings are normally needed besides `-D PKG_COLVARS=yes`.

---

#### Traditional make

Before building LAMMPS, one must build the Colvars library in `lib/colvars`.

This can be done manually in the same folder by using or adapting one of the provided Makefiles: for example, `Makefile.g++` for the GNU C++ compiler. C++11 compatibility may need to be enabled for some older compilers (as is done in the example makefile).

In general, it is safer to use build setting consistent with the rest of LAMMPS. This is best carried out from the LAMMPS `src` directory using a command like these, which simply invoke the `lib/colvars/Install.py` script with the specified args:

```
$ make lib-colvars           # print help message
$ make lib-colvars args="-m serial" # build with GNU g++ compiler
↪(settings as with "make serial")
$ make lib-colvars args="-m mpi"   # build with default MPI compiler
↪(settings as with "make mpi")
$ make lib-colvars args="-m g++-debug" # build with GNU g++ compiler and
↪colvars debugging enabled
```

The “machine” argument of the “-m” flag is used to find a `Makefile.machine` to use as build recipe. If it does not already exist in `lib/colvars`, it will be auto-generated by using compiler flags consistent with those parsed from the core LAMMPS makefiles.

Optional flags may be specified as environment variables:

```
$ COLVARS_DEBUG=yes make lib-colvars args="-m machine" # Build with debug
↪code (much slower)
$ COLVARS_LEPTON=no make lib-colvars args="-m machine" # Build without Lepton
↪(included otherwise)
```

The build should produce two files: the library `lib/colvars/libcolvars.a` (which also includes Lepton objects if enabled) and the specification file `lib/colvars/Makefile.lammps`. The latter is auto-generated, and normally does not need to be edited.

---

### 3.7.16 ELECTRODE package

This package depends on the KSPACE package.

---

#### CMake build

No additional settings are needed besides `-D PKG_KSPACE=yes` and `-D PKG_ELECTRODE=yes`.

---

#### Traditional make

The package is activated with `make yes-KSPACE` and `make yes-ELECTRODE`

Note that the `Makefile.lammps` file has settings for the BLAS and LAPACK linear algebra libraries. As explained in `lib/awpmd/README` these can either exist on your system, or you can use the files provided in `lib/linalg`. In the latter case you also need to build the library in `lib/linalg` with a command like these:

```
$ make lib-linalg           # print help message
$ make lib-linalg args="-m serial" # build with GNU Fortran compiler
↪(settings as with "make serial")
$ make lib-linalg args="-m mpi"   # build with default MPI Fortran
↪compiler (settings as with "make mpi")
$ make lib-linalg args="-m gfortran" # build with GNU Fortran compiler
```

---

### 3.7.17 ML-PACE package

This package requires a library that can be downloaded and built in `lib/pace` or somewhere else, which must be done before building LAMMPS with this package. The code for the library can be found at: <https://github.com/ICAMS/lammps-user-pace/>

---

#### CMake build

By default the library will be downloaded from the git repository and built automatically when the ML-PACE package is enabled with `-D PKG_ML-PACE=yes`. The location for the sources may be customized by setting the variable `PACELIB_URL` when configuring with CMake (e.g. to use a local archive on machines without internet access). Since CMake checks the validity of the archive with `md5sum` you may also need to set `PACELIB_MD5` if you provide a different library version than what is downloaded automatically.

---

#### Traditional make

You can download and build the ML-PACE library in one step from the `lammps/src` dir, using these commands, which invoke the `lib/pace/Install.py` script.

```
$ make lib-pace           # print help message
$ make lib-pace args="-b" # download and build the default
↪version in lib/pace
```

---

You should not need to edit the `lib/pace/Makefile.lammps` file.

---

### 3.7.18 PLUMED package

Before building LAMMPS with this package, you must first build PLUMED. PLUMED can be built as part of the LAMMPS build or installed separately from LAMMPS using the generic [PLUMED installation instructions](#). The PLUMED package has been tested to work with Plumed versions 2.4.x, 2.5.x, and 2.6.x and will error out, when trying to run calculations with a different version of the Plumed kernel.

PLUMED can be linked into MD codes in three different modes: static, shared, and runtime. With the “static” mode, all the code that PLUMED requires is linked statically into LAMMPS. LAMMPS is then fully independent from the PLUMED installation, but you have to rebuild/relink it in order to update the PLUMED code inside it. With the “shared” linkage mode, LAMMPS is linked to a shared library that contains the PLUMED code. This library should preferably be installed in a globally accessible location. When PLUMED is linked in this way the same library can be used by multiple MD packages. Furthermore, the PLUMED library LAMMPS uses can be updated without the need for a recompile of LAMMPS for as long as the shared PLUMED library is ABI-compatible.

The third linkage mode is “runtime” which allows the user to specify which PLUMED kernel should be used at runtime by using the `PLUMED_KERNEL` environment variable. This variable should point to the location of the `libplumed-Kernel.so` dynamical shared object, which is then loaded at runtime. This mode of linking is particularly convenient for doing PLUMED development and comparing multiple PLUMED versions as these sorts of comparisons can be done without recompiling the hosting MD code. All three linkage modes are supported by LAMMPS on selected operating systems (e.g. Linux) and using either CMake or traditional make build. The “static” mode should be the most portable, while the “runtime” mode support in LAMMPS makes the most assumptions about operating system and compiler environment. If one mode does not work, try a different one, switch to a different build system, consider a global PLUMED installation or consider downloading PLUMED during the LAMMPS build.

---

#### CMake build

When the `-D PKG_PLUMED=yes` flag is included in the `cmake` command you must ensure that GSL is installed in locations that are specified in your environment. There are then two additional variables that control the manner in which PLUMED is obtained and linked into LAMMPS.

```
-D DOWNLOAD_PLUMED=value    # download PLUMED for build, value = no (default),  
→ or yes  
-D PLUMED_MODE=value       # Linkage mode for PLUMED, value = static (default),  
→ shared, or runtime
```

If `DOWNLOAD_PLUMED` is set to “yes”, the PLUMED library will be downloaded (the version of PLUMED that will be downloaded is hard-coded to a vetted version of PLUMED, usually a recent stable release version) and built inside the CMake build directory. If `DOWNLOAD_PLUMED` is set to “no” (the default), CMake will try to detect and link to an installed version of PLUMED. For this to work, the PLUMED library has to be installed into a location where the `pkg-config` tool can find it or the `PKG_CONFIG_PATH` environment variable has to be set up accordingly. PLUMED should be installed in such a location if you compile it using the default `make`; `make install` commands.

The `PLUMED_MODE` setting determines the linkage mode for the PLUMED library. The allowed values for this flag are “static” (default), “shared”, or “runtime”. If you want to switch the linkage mode, just re-run CMake with a different setting. For a discussion of PLUMED linkage modes, please see above. When `DOWNLOAD_PLUMED` is enabled the static linkage mode is recommended.

---

#### Traditional make

PLUMED needs to be installed before the PLUMED package is installed so that LAMMPS can find the right settings when compiling and linking the LAMMPS executable. You can either download and build PLUMED inside the LAMMPS plumed library folder or use a previously installed PLUMED library and point LAMMPS to its location. You also have to choose the linkage mode: “static” (default), “shared” or “runtime”. For a discussion of PLUMED linkage modes, please see above.

Download/compilation/configuration of the plumed library can be done from the src folder through the following make args:

```
$ make lib-plumed # print help message
$ make lib-plumed args="-b" # download and build PLUMED in lib/
  →plumed/plumed2
$ make lib-plumed args="-p $HOME/.local" # use existing PLUMED installation
  →in $HOME/.local
$ make lib-plumed args="-p /usr/local -m shared" # use existing PLUMED
  →installation in # /usr/local and use shared
  →linkage mode
```

Note that 2 symbolic (soft) links, `includelink` and `liblink` are created in `lib/plumed` that point to the location of the PLUMED build to use. A new file `lib/plumed/Makefile.lammps` is also created with settings suitable for LAMMPS to compile and link PLUMED using the desired linkage mode. After this step is completed, you can install the PLUMED package and compile LAMMPS in the usual manner:

```
$ make yes-plumed
$ make machine
```

Once this compilation completes you should be able to run LAMMPS in the usual way. For shared linkage mode, `libplumed.so` must be found by the LAMMPS executable, which on many operating systems means, you have to set the `LD_LIBRARY_PATH` environment variable accordingly.

Support for the different linkage modes in LAMMPS varies for different operating systems, using the static linkage is expected to be the most portable, and thus set to be the default.

If you want to change the linkage mode, you have to re-run “make lib-plumed” with the desired settings **and** do a re-install if the PLUMED package with “make yes-plumed” to update the required makefile settings with the changes in the `lib/plumed` folder.

---

### 3.7.19 H5MD package

To build with this package you must have the HDF5 software package installed on your system, which should include the `h5cc` compiler and the HDF5 library.

---

#### CMake build

No additional settings are needed besides `-D PKG_H5MD=yes`.

This should auto-detect the H5MD library on your system. Several advanced CMake H5MD options exist if you need to specify where it is installed. Use the `cmake` (terminal window) or `cmake-gui` (graphical) tools to see these options and set them interactively from their user interfaces.

---

---

#### Traditional make

Before building LAMMPS, you must build the CH5MD library in `lib/h5md`. You can do this manually if you prefer; follow the instructions in `lib/h5md/README`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/h5md/Install.py` script with the specified args:

```
$ make lib-h5md           # print help message
$ make lib-h5md args="-m h5cc" # build with h5cc compiler
```

The build should produce two files: `lib/h5md/libch5md.a` and `lib/h5md/Makefile.lammps`. The latter is copied from an existing `Makefile.lammps.*` and has settings needed to build LAMMPS with the system HDF5 library. If necessary, you can edit/create a new `lib/h5md/Makefile.machine` file for your system, which should define an `EXTRAMAKE` variable to specify a corresponding `Makefile.lammps.<machine>` file.

### 3.7.20 ML-HDNNP package

To build with the ML-HDNNP package it is required to download and build the external `n2p2` library v2.1.4 (or higher). The LAMMPS build process offers an automatic download and compilation of `n2p2` or allows you to choose the installation directory of `n2p2` manually. Please see the boxes below for the CMake and traditional build system for detailed information.

In case of a manual installation of `n2p2` you only need to build the `n2p2` core library `libnnp` and interface library `libnnpif`. When using GCC it should suffice to execute `make libnnpif` in the `n2p2` src directory. For more details please see `lib/hdnnp/README` and the `n2p2` build documentation.

#### CMake build

```
-D DOWNLOAD_N2P2=value    # download n2p2 for build, value = no (default) or
↪yes
-D N2P2_DIR=path          # n2p2 base directory (only needed if a custom
↪location)
```

If `DOWNLOAD_N2P2` is set, the `n2p2` library will be downloaded and built inside the CMake build directory. If the `n2p2` library is already on your system (in a location CMake cannot find it), set the `N2P2_DIR` path where `n2p2` is located. If `n2p2` is located directly in `lib/hdnnp/n2p2` it will be automatically found by CMake.

#### Traditional make

You can download and build the `n2p2` library manually if you prefer; follow the instructions in `lib/hdnnp/README`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/hdnnp/Install.py` script with the specified args:

```
$ make lib-hdnnp          # print help message
$ make lib-hdnnp args="-b" # download and build in lib/hdnnp/n2p2-...
$ make lib-hdnnp args="-b -v 2.1.4" # download and build specific version
$ make lib-hdnnp args="-p /usr/local/n2p2" # use the existing n2p2
↪installation in /usr/local/n2p2
```

Note that 3 symbolic (soft) links, `includelink`, `liblink` and `Makefile.lammps`, will be created in `lib/hdnp` to point to `n2p2/include`, `n2p2/lib` and `n2p2/lib/Makefile.lammps-extra`, respectively. When LAMMPS is built in `src` it will use these links.

---

### 3.7.21 INTEL package

To build with this package, you must choose which hardware you want to build for, either x86 CPUs or Intel KNLs in offload mode. You should also typically *install the OPENMP package*, as it can be used in tandem with the INTEL package to good effect, as explained on the *INTEL package* page.

When using Intel compilers version 16.0 or later is required. You can also use the GNU or Clang compilers and they will provide performance improvements over regular styles and OPENMP styles, but less so than with the Intel compilers. Please also note, that some compilers have been found to apply memory alignment constraints incompletely or incorrectly and thus can cause segmentation faults in otherwise correct code when using features from the INTEL package.

---

#### CMake build

```
-D INTEL_ARCH=value      # value = cpu (default) or knl
-D INTEL_LRT_MODE=value # value = threads, none, or c++11
```

---

---

#### Traditional make

Choose which hardware to compile for in `Makefile.machine` via the following settings. See `src/MAKE/OPTIONS/Makefile.intel_cpu*` and `Makefile.knl` files for examples. and `src/INTEL/README` for additional information.

For CPUs:

```
OPTFLAGS =      -xHost -O2 -fp-model fast=2 -no-prec-div -qoverride-limits -
↳qopt-zmm-usage=high
CCFLAGS =      -g -qopenmp -DLAMMPS_MEMALIGN=64 -no-offload -fno-alias -ansi-
↳alias -restrict $(OPTFLAGS)
LINKFLAGS =    -g -qopenmp $(OPTFLAGS)
LIB =          -ltbbmalloc
```

For KNLs:

```
OPTFLAGS =      -xMIC-AVX512 -O2 -fp-model fast=2 -no-prec-div -qoverride-
↳limits
CCFLAGS =      -g -qopenmp -DLAMMPS_MEMALIGN=64 -no-offload -fno-alias -ansi-
↳alias -restrict $(OPTFLAGS)
LINKFLAGS =    -g -qopenmp $(OPTFLAGS)
LIB =          -ltbbmalloc
```

---

In Long-range thread mode (LRT) a modified verlet style is used, that operates the Kspace calculation in a separate thread concurrently to other calculations. This has to be enabled in the *package intel* command at runtime. With the setting “threads” it used the `pthread` library, while “c++11” will use the built-in thread support of C++11 compilers.



The option “none” skips compilation of this feature. The default is to use “threads” if pthreads is available and otherwise “none”.

Best performance is achieved with Intel hardware, Intel compilers, as well as the Intel TBB and MKL libraries. However, the code also compiles, links, and runs with other compilers / hardware and without TBB and MKL.

### 3.7.22 MDI package

#### CMake build

```
-D DOWNLOAD_MDI=value      # download MDI Library for build, value = no,
↪(default) or yes
```

#### Traditional make

Before building LAMMPS, you must build the MDI Library in `lib/mdi`. You can do this by executing a command like one of the following from the `lib/mdi` directory:

```
$ python Install.py -m gcc      # build using gcc compiler
$ python Install.py -m icc      # build using icc compiler
```

The build should produce two files: `lib/mdi/includelink/mdi.h` and `lib/mdi/liblink/libmdi.so`.

### 3.7.23 MESONT package

This package includes a library written in Fortran 90 in the `lib/mesont` folder, so a working Fortran 90 compiler is required to compile it. Also, the files with the force field data for running the bundled examples are not included in the source distribution. Instead they will be downloaded the first time this package is installed.

#### CMake build

No additional settings are needed besides `-D PKG_MESONT=yes`

#### Traditional make

Before building LAMMPS, you must build the `mesont` library in `lib/mesont`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/mesont/Install.py` script with the specified args:

```
$ make lib-mesont           # print help message
$ make lib-mesont args="-m gfortran" # build with GNU g++ compiler (settings,
↪as with "make serial")
$ make lib-mesont args="-m ifort"   # build with Intel icc compiler
```

The build should produce two files: `lib/mesont/libmesont.a` and `lib/mesont/Makefile.lammps`. The latter is copied from an existing `Makefile.lammps.*` and has settings needed to build LAMMPS with the `mesont` library (though typically the settings contain only the Fortran runtime library). If necessary, you can edit/create a new `lib/mesont/Makefile.machine` file for your system, which should define an `EXTRAMAKE` variable to specify a corresponding `Makefile.lammps.machine` file.

---

### 3.7.24 MOLFILE package

---

#### CMake build

```
-D MOLFILE_INCLUDE_DIR=path # (optional) path where VMD molfile plugin
->headers are installed
-D PKG_MOLFILE=yes
```

Using `-D PKG_MOLFILE=yes` enables the package, and setting `-D MOLFILE_INCLUDE_DIR` allows to provide a custom location for the molfile plugin header files. These should match the ABI of the plugin files used, and thus one typically sets them to include folder of the local VMD installation in use. LAMMPS ships with a couple of default header files that correspond to a popular VMD version, usually the latest release.

---

---

#### Traditional make

The `lib/molfile/Makefile.lammps` file has a setting for a dynamic loading library `libdl.a` that is typically present on all systems. It is required for LAMMPS to link with this package. If the setting is not valid for your system, you will need to edit the `Makefile.lammps` file. See `lib/molfile/README` and `lib/molfile/Makefile.lammps` for details. It is also possible to configure a different folder with the VMD molfile plugin header files. LAMMPS ships with a couple of default headers, but these are not compatible with all VMD versions, so it is often best to change this setting to the location of the same include files of the local VMD installation in use.

---

### 3.7.25 NETCDF package

To build with this package you must have the NetCDF library installed on your system.

---

#### CMake build

No additional settings are needed besides `-D PKG_NETCDF=yes`.

This should auto-detect the NETCDF library if it is installed on your system at standard locations. Several advanced CMake NETCDF options exist if you need to specify where it was installed. Use the `ccmake` (terminal window) or `cmake-gui` (graphical) tools to see these options and set them interactively from their user interfaces.

---

---

#### Traditional make

The `lib/netcdf/Makefile.lammps` file has settings for NetCDF include and library files which LAMMPS needs to build with this package. If the settings are not valid for your system, you will need to edit the `Makefile.lammps` file. See `lib/netcdf/README` for details.

### 3.7.26 OPENMP package

#### CMake build

No additional settings are required besides `-D PKG_OPENMP=yes`. If CMake detects OpenMP compiler support, the OPENMP code will be compiled with multi-threading support enabled, otherwise as optimized serial code.

#### Traditional make

To enable multi-threading support in the OPENMP package (and other styles supporting OpenMP) the following compile and link flags must be added to your `Makefile.machine` file. See `src/MAKE/OPTIONS/Makefile.omp` for an example.

```
CCFLAGS: -fopenmp          # for GNU and Clang Compilers
CCFLAGS: -qopenmp -restrict # for Intel compilers on Linux
LINKFLAGS: -fopenmp       # for GNU and Clang Compilers
LINKFLAGS: -qopenmp       # for Intel compilers on Linux
```

For other platforms and compilers, please consult the documentation about OpenMP support for your compiler.

### 3.7.27 QMMM package

For using LAMMPS to do QM/MM simulations via the QMMM package you need to build LAMMPS as a library. A LAMMPS executable with *fix qmmm* included can be built, but will not be able to do a QM/MM simulation on as such. You must also build a QM code - currently only Quantum ESPRESSO (QE) is supported - and create a new executable which links LAMMPS and the QM code together. Details are given in the `lib/qmmm/README` file. It is also recommended to read the instructions for *linking with LAMMPS as a library* for background information. This requires compatible Quantum Espresso and LAMMPS versions. The current interface and makefiles have last been verified to work in February 2020 with Quantum Espresso versions 6.3 to 6.5.

#### CMake build

When using CMake, building a LAMMPS library is required and it is recommended to build a shared library, since any libraries built from the sources in the `lib` folder (including the essential `libqmmm.a`) are not included in the static LAMMPS library and (currently) not installed, while their code is included in the shared LAMMPS library. Thus a typical command line to configure building LAMMPS for QMMM would be:

```
cmake -C ../cmake/presets/basic.cmake -D PKG_QMMM=yes \
-D BUILD_LIB=yes -DBUILD_SHARED_LIBS=yes ../cmake
```

After completing the LAMMPS build and also configuring and compiling Quantum ESPRESSO with external library support (via “make couple”), go back to the `lib/qmmm` folder and follow the instructions on the README file to build the combined LAMMPS/QE QM/MM executable (`pwqmmm.x`) in the `lib/qmmm` folder.

---

### Traditional make

Before building LAMMPS, you must build the QMMM library in `lib/qmmm`. You can do this manually if you prefer; follow the first two steps explained in `lib/qmmm/README`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/qmmm/Install.py` script with the specified args:

```
$ make lib-qmmm                # print help message
$ make lib-qmmm args="-m serial" # build with GNU Fortran compiler
↪(settings as in "make serial")
$ make lib-qmmm args="-m mpi"   # build with default MPI compiler
↪(settings as in "make mpi")
$ make lib-qmmm args="-m gfortran" # build with GNU Fortran compiler
```

The build should produce two files: `lib/qmmm/libqmmm.a` and `lib/qmmm/Makefile.lammps`. The latter is copied from an existing `Makefile.lammps.*` and has settings needed to build LAMMPS with the QMMM library (though typically the settings are just blank). If necessary, you can edit/create a new `lib/qmmm/Makefile.<machine>` file for your system, which should define an `EXTRAMAKE` variable to specify a corresponding `Makefile.lammps.<machine>` file.

You can then install QMMM package and build LAMMPS in the usual manner. After completing the LAMMPS build and compiling Quantum ESPRESSO with external library support (via “make couple”), go back to the `lib/qmmm` folder and follow the instructions in the README file to build the combined LAMMPS/QE QM/MM executable (`pwqmmm.x`) in the `lib/qmmm` folder.

---

## 3.7.28 ML-QUIP package

To build with this package, you must download and build the QUIP library. It can be obtained from GitHub. For support of GAP potentials, additional files with specific licensing conditions need to be downloaded and configured. The automatic download will from within CMake will download the non-commercial use version.

---

### CMake build

```
-D DOWNLOAD_QUIP=value      # download QUIP library for build, value = no
↪(default) or yes
-D QUIP_LIBRARY=path       # path to libquip.a (only needed if a custom
↪location)
-D USE_INTERNAL_LINALG=value # Use the internal linear algebra library instead
↪of LAPACK
                           # value = no (default) or yes
```

CMake will try to download and build the QUIP library from GitHub, if it is not found on the local machine. This requires to have git installed. It will use the same compilers and flags as used for compiling LAMMPS. Currently this is only supported for the GNU and the Intel compilers. Set the `QUIP_LIBRARY` variable if you want to use a previously compiled and installed QUIP library and CMake cannot find it.

The QUIP library requires LAPACK (and BLAS) and CMake can identify their locations and pass that info to the QUIP build script. But on some systems this triggers a (current) limitation of CMake and the configuration will fail. Try enabling `USE_INTERNAL_LINALG` in those cases to use the bundled linear algebra library and work around the limitation.

---

### Traditional make

The download/build procedure for the QUIP library, described in `lib/quip/README` file requires setting two environment variables, `QUIP_ROOT` and `QUIP_ARCH`. These are accessed by the `lib/quip/Makefile.lammps` file which is used when you compile and link LAMMPS with this package. You should only need to edit `Makefile.lammps` if the LAMMPS build can not use its settings to successfully build on your system.

---

## 3.7.29 SCAFACOS package

To build with this package, you must download and build the [ScaFaCoS Coulomb solver library](#)

---

### CMake build

```
-D DOWNLOAD_SCAFACOS=value      # download ScaFaCoS for build, value = no_
↪(default) or yes
-D SCAFACOS_LIBRARY=path       # ScaFaCos library file (only needed if at_
↪custom location)
-D SCAFACOS_INCLUDE_DIR=path   # ScaFaCoS include directory (only needed if at_
↪custom location)
```

If `DOWNLOAD_SCAFACOS` is set, the ScaFaCoS library will be downloaded and built inside the CMake build directory. If the ScaFaCoS library is already on your system (in a location CMake cannot find it), `SCAFACOS_LIBRARY` is the filename (plus path) of the ScaFaCoS library file, not the directory the library file is in. `SCAFACOS_INCLUDE_DIR` is the directory the ScaFaCoS include file is in.

---

### Traditional make

You can download and build the ScaFaCoS library manually if you prefer; follow the instructions in `lib/scafacos/README`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/scafacos/Install.py` script with the specified args:

```
make lib-scafacos                # print help message
make lib-scafacos args="-b"      # download and build in lib/scafacos/
↪scafacos-<version>
make lib-scafacos args="-p $HOME/scafacos # use existing ScaFaCoS_
↪installation in $HOME/scafacos
```

Note that 2 symbolic (soft) links, `includelink` and `liblink`, are created in `lib/scafacos` to point to the ScaFaCoS src dir. When LAMMPS builds in `src` it will use these links. You should not need to edit the `lib/scafacos/Makefile.lammps` file.

---

### 3.7.30 MACHDYN package

To build with this package, you must download the Eigen3 library. Eigen3 is a template library, so you do not need to build it.

---

#### CMake build

```
-D DOWNLOAD_EIGEN3           # download Eigen3, value = no (default) or yes
-D EIGEN3_INCLUDE_DIR=path   # path to Eigen library (only needed if a custom
↪location)
```

If `DOWNLOAD_EIGEN3` is set, the Eigen3 library will be downloaded and inside the CMake build directory. If the Eigen3 library is already on your system (in a location where CMake cannot find it), set `EIGEN3_INCLUDE_DIR` to the directory the Eigen3 include file is in.

---

---

#### Traditional make

You can download the Eigen3 library manually if you prefer; follow the instructions in `lib/smd/README`. You can also do it in one step from the `lammps/src` dir, using a command like these, which simply invoke the `lib/smd/Install.py` script with the specified args:

```
$ make lib-smd                # print help message
$ make lib-smd args="-b"      # download to lib/smd/eigen3
$ make lib-smd args="-p /usr/include/eigen3" # use existing Eigen
↪installation in /usr/include/eigen3
```

Note that a symbolic (soft) link named `includeLink` is created in `lib/smd` to point to the Eigen dir. When LAMMPS builds it will use this link. You should not need to edit the `lib/smd/Makefile.lammps` file.

---

### 3.7.31 VTK package

To build with this package you must have the VTK library installed on your system.

---

#### CMake build

No additional settings are needed besides `-D PKG_VTK=yes`.

This should auto-detect the VTK library if it is installed on your system at standard locations. Several advanced VTK options exist if you need to specify where it was installed. Use the `ccmake` (terminal window) or `cmake-gui` (graphical) tools to see these options and set them interactively from their user interfaces.

---

---

#### Traditional make

The `lib/vtk/Makefile.lammps` file has settings for accessing VTK files and its library, which LAMMPS needs to build with this package. If the settings are not valid for your system, check if one of the other `lib/vtk/Makefile.lammps.*` files is compatible and copy it to `Makefile.lammps`. If none of the provided files work, you will need to edit the `Makefile.lammps` file. See `lib/vtk/README` for details.

---

## 3.8 Build the LAMMPS documentation

Depending on how you obtained LAMMPS and whether you have built the manual yourself, this directory has a number of sub-directories and files. Here is a list with descriptions:

```

README           # brief info about the documentation
src              # content files for LAMMPS documentation
html            # HTML version of the LAMMPS manual (see html/Manual.html)
utils           # tools and settings for building the documentation
lammmps.1       # man page for the lammmps command
msi2lmp.1       # man page for the msi2lmp command
Manual.pdf      # large PDF version of entire manual
LAMMPS.epub     # Manual in ePUB e-book format
LAMMPS.mobi     # Manual in MOBI e-book format
docenv          # virtualenv folder for processing the manual sources
doctrees        # temporary data from processing the manual
doxygen         # doxygen configuration and output
.gitignore      # list of files and folders to be ignored by git
doxygen-warn.log # logfile with warnings from running doxygen
github-development-workflow.md # notes on the LAMMPS development workflow

```

If you downloaded LAMMPS as a tarball from the [LAMMPS website](https://www.lammps.org/), the html folder and the PDF files should be included.

If you downloaded LAMMPS from the public git repository, then the HTML and PDF files are not included. You can build the HTML or PDF files yourself, by typing `make html` or `make pdf` in the doc folder. This requires various tools and files. Some of them have to be installed (see below). For the rest the build process will attempt to download and install them into a python virtual environment and local folders.

A current version of the manual (latest patch release, that is the state of the *release* branch) is available online at: <https://docs.lammps.org/>. A version of the manual corresponding to the ongoing development (that is the state of the *develop* branch) is available online at: <https://docs.lammps.org/latest/>. A version of the manual corresponding to the latest stable LAMMPS release (that is the state of the *stable* branch) is available online at: <https://docs.lammps.org/stable/>

### 3.8.1 Build using GNU make

The LAMMPS manual is written in `reStructuredText` format which can be translated to different output format using the `Sphinx` document generator tool. It also incorporates programmer documentation extracted from the LAMMPS C++ sources through the `Doxygen` program. Currently the translation to HTML, PDF (via LaTeX), ePUB (for many e-book readers) and MOBI (for Amazon Kindle readers) are supported. For that to work a Python 3 interpreter, the `doxygen` tools and internet access to download additional files and tools are required. This download is usually only required once or after the documentation folder is returned to a pristine state with `make clean-all`.

For the documentation build a python virtual environment is set up in the folder `doc/docenv` and various python packages are installed into that virtual environment via the `pip` tool. For rendering embedded LaTeX code also the `MathJax` JavaScript engine needs to be downloaded. If you need to pass additional options to the pip commands to work (e.g. to use a web proxy or to point to additional SSL certificates) you can set them via the `PIP_OPTIONS` environment variable or uncomment and edit the `PIP_OPTIONS` setting at beginning of the makefile.

The actual translation is then done via `make` commands in the doc folder. The following `make` commands are available:

```

make html       # generate HTML in html dir using Sphinx
make pdf        # generate PDF as Manual.pdf using Sphinx and PDFLaTeX
make epub       # generate LAMMPS.epub in ePUB format using Sphinx

```

(continues on next page)

(continued from previous page)

```
make mobi          # generate LAMMPS.mobi in MOBI format using ebook-convert
make fasthtml     # generate approximate HTML in fasthtml dir using Sphinx
                  # some Sphinx extensions do not work correctly with this
make clean        # remove intermediate RST files created by HTML build
make clean-all   # remove entire build folder and any cached data
make anchor_check # check for duplicate anchor labels
make style_check  # check for complete and consistent style lists
make package_check # check for complete and consistent package lists
make spelling     # spell-check the manual
```

---

### 3.8.2 Build using CMake

It is also possible to create the HTML version (and **only** the HTML version) of the manual within the *CMake build directory*. The reason for this option is to include the installation of the HTML manual pages into the “install” step when installing LAMMPS after the CMake build via `cmake --build . --target install`. The documentation build is included in the default build target, but can also be requested independently with `cmake --build . --target doc`. If you need to pass additional options to the pip commands to work (e.g. to use a web proxy or to point to additional SSL certificates) you can set them via the `PIP_OPTIONS` environment variable.

```
-D BUILD_DOC=value      # yes or no (default)
```

---

### 3.8.3 Prerequisites for HTML

To run the HTML documentation build toolchain, python 3, git, doxygen, and virtualenv have to be installed locally. Here are instructions for common setups:

---

#### Ubuntu

```
sudo apt-get install python-virtualenv git doxygen
```

---

#### RHEL or CentOS (Version 7.x)

```
sudo yum install python3-virtualenv git doxygen
```

---

#### Fedora or RHEL/CentOS (8.x or later)

```
sudo dnf install python3-virtualenv git doxygen
```

---



---

## MacOS X

### *Python 3*

Download the latest Python 3 MacOS X package from <https://www.python.org> and install it. This will install both Python 3 and pip3.

### *virtualenv*

Once Python 3 is installed, open a Terminal and type

```
pip3 install virtualenv
```

This will install virtualenv from the Python Package Index.

---

## 3.8.4 Prerequisites for PDF

In addition to the tools needed for building the HTML format manual, a working LaTeX installation with support for PDFLaTeX and a selection of LaTeX styles/packages are required. To run the PDFLaTeX translation the `latexmk` script needs to be installed as well.

## 3.8.5 Prerequisites for ePUB and MOBI

In addition to the tools needed for building the HTML format manual, a working LaTeX installation with a few add-on LaTeX packages as well as the `dvi.png` tool are required to convert embedded math expressions transparently into embedded images.

For converting the generated ePUB file to a MOBI format file (for e-book readers, like Kindle, that cannot read ePUB), you also need to have the `ebook-convert` tool from the “calibre” software installed. <http://calibre-ebook.com/> Typing `make mobi` will first create the ePUB file and then convert it. On the Kindle readers in particular, you also have support for PDF files, so you could download and view the PDF version as an alternative.

## 3.8.6 Instructions for Developers

When adding new styles or options to the LAMMPS code, corresponding documentation is required and either existing files in the `src` folder need to be updated or new files added. These files are written in `reStructuredText` markup for translation with the Sphinx tool.

Before contributing any documentation, please check that both the HTML and the PDF format documentation can translate without errors. During testing the html translation, you may use the `make fasthtml` command which does an approximate translation (i.e. not all Sphinx features and extensions will work), but runs very fast because it will only translate files that have been changed since the last `make fasthtml` command.

Please also check the output to the console for any warnings or problems. There will be multiple tests run automatically:

- A test for correctness of all anchor labels and their references
- A test that all LAMMPS packages (= folders with sources in `lammeps/src`) are documented and listed. A typical warning shows the name of the folder with the suspected new package code and the documentation files where they need to be listed:

```
Found 88 packages
Package NEWPACKAGE missing in Packages_list.rst
Package NEWPACKAGE missing in Packages_details.rst
```

- A test that only standard, printable ASCII text characters are used. This runs the command `env LC_ALL=C grep -n '[^ -~]' src/*.rst` and thus prints all offending lines with filename and line number prepended to the screen. Special characters like the Angstrom Å should be typeset with embedded math (like this `:math:\`\).`
- A test whether all styles are documented and listed in their respective overview pages. A typical output with warnings looks like this:

```
Parsed style names w/o suffixes from C++ tree in ../src:
  Angle styles:      21   Atom styles:      24
  Body styles:       3   Bond styles:      17
  Command styles:   41   Compute styles:  143
  Dihedral styles:  16   Dump styles:     26
  Fix styles:       223  Improper styles:  13
  Integrate styles:  4   Kspace styles:   15
  Minimize styles:  9   Pair styles:     234
  Reader styles:    4   Region styles:   8
Compute style entry newcomp is missing or incomplete in Commands_compute.rst
Compute style entry newcomp is missing or incomplete in compute.rst
Fix style entry newfix is missing or incomplete in Commands_fix.rst
Fix style entry newfix is missing or incomplete in fix.rst
Pair style entry new is missing or incomplete in Commands_pair.rst
Pair style entry new is missing or incomplete in pair_style.rst
Found 6 issue(s) with style lists
```

In addition, there is the option to run a spellcheck on the entire manual with `make spelling`. This requires a library called `enchant`. To avoid printing out *false positives* (e.g. keywords, names, abbreviations) those can be added to the file `lammps/doc/utils/sphinx-config/false_positives.txt`.

## 3.9 Notes for building LAMMPS on Windows

- *General remarks*
  - *Running Linux on Windows*
  - *Using GNU GCC ported to Windows*
  - *Using Visual Studio*
  - *Using Intel oneAPI compilers and libraries*
  - *Using a cross-compiler*
-

### 3.9.1 General remarks

LAMMPS is developed and tested primarily on Linux machines. The vast majority of HPC clusters and supercomputers today run on Linux as well. While portability to other platforms is desired, it is not always achieved. That is sometimes due to non-portable code in LAMMPS itself, but more often due to portability limitations of external libraries and tools required to build a specific feature or package. The LAMMPS developers are dependent on LAMMPS users giving feedback and providing assistance in resolving portability issues. This is particularly true for compiling LAMMPS on Windows, since this platform has significant differences in some low-level functionality. As of LAMMPS version 14 December 2021, large parts of LAMMPS can be compiled natively with the Microsoft Visual C++ Compilers. As of LAMMPS version 31 May 2022, also the Intel oneAPI compilers can compile large parts of LAMMPS natively on Windows. This is mostly facilitated by using the *Platform abstraction functions* in the `platform` namespace and CMake.

Before trying to build LAMMPS on Windows yourself, please consider the [pre-compiled Windows installer packages](#) and see if they are sufficient for your needs.

### 3.9.2 Running Linux on Windows

If it is necessary for you to compile LAMMPS on a Windows machine (e.g. because it is your main desktop), please also consider using a virtual machine software and compile and run LAMMPS in a Linux virtual machine, or - if you have a sufficiently up-to-date Windows 10 or Windows 11 installation - consider using the Windows subsystem for Linux. This optional Windows feature allows you to run the bash shell of a Linux system (Ubuntu by default) from within Windows and from there on, you can pretty much use that shell like you are running on a regular Ubuntu Linux machine (e.g. installing software via apt-get and more). For more details on that, please see [this tutorial](#).

### 3.9.3 Using a GNU GCC ported to Windows

One option for compiling LAMMPS on Windows natively is to install a Bash shell, Unix shell utilities, Perl, Python, GNU make, and a GNU compiler ported to Windows. The Cygwin package provides a unix/linux interface to low-level Windows functions, so LAMMPS can be compiled on Windows. The necessary (minor) modifications to LAMMPS are included, but may not always be up-to-date for recently added functionality and the corresponding new code. A machine makefile for using cygwin for the old build system is provided. Using CMake for this mode of compilation is untested and not likely to work.

When compiling for Windows do **not** set the `-DLAMMPS_MEMALIGN` define in the `LMP_INC` makefile variable and add `-lwsck32 -lpsapi` to the linker flags in `LIB` makefile variable. Try adding `-static-libgcc` or `-static` or both to the linker flags when your resulting LAMMPS Windows executable complains about missing `.dll` files. The CMake configuration should set this up automatically, but is untested.

In case of problems, you are recommended to contact somebody with experience in using Cygwin. If you do come across portability problems requiring changes to the LAMMPS source code, or figure out corrections yourself, please report them on the [LAMMPS forum at MatSci](#), or file them as an issue or pull request on the LAMMPS GitHub project.

### 3.9.4 Using Microsoft Visual Studio

Following the integration of the *platform namespace* into the LAMMPS code base, portability of LAMMPS for native compilation on Windows using Visual Studio has been significantly improved. This has been tested with Visual Studio 2019 (aka version 16) and Visual Studio 2022 (aka version 17). We strongly recommend using Visual Studio 2022 version 17.1 or later. Not all features and packages in LAMMPS are currently supported out of the box, but a preset `cmake/presets/windows.cmake` is provided that contains the packages that have been compiled successfully so far. You **must** use the CMake based build procedure, since there is no support for GNU make or the Unix shell utilities required for the GNU make build procedure.

It is possible to use both the integrated CMake support of the Visual Studio IDE or use an external CMake installation (e.g. downloaded from [cmake.org](http://cmake.org)) to create build files and compile LAMMPS from the command line.

Compilation via command line and unit tests are checked automatically for the LAMMPS development branch through [GitHub Actions](#).

---

**Note:** Versions of Visual Studio before version 17.1 may scan the entire LAMMPS source tree and likely miss the correct master `CMakeLists.txt` and get confused since there are multiple files of that name in different folders but none in top level folder.

---

Please note, that for either approach CMake will create a so-called “*multi-configuration*” *build environment*, and the command lines for building and testing LAMMPS must be adjusted accordingly.

The LAMMPS `cmake` folder contains a `CMakeSettings.json` file with build configurations for MSVC compilers and the MS provided Clang compiler package in Debug and Release mode.

To support running in parallel you can compile with OpenMP enabled using the OPENMP package or install Microsoft MPI (including the SDK) and compile LAMMPS with MPI enabled.

---

**Note:** This is work in progress and you should contact the LAMMPS developers via [GitHub](#) or the [LAMMPS forum at MatSci](#), if you have questions or LAMMPS specific problems.

---

### 3.9.5 Using Intel oneAPI Compilers and Libraries

New in version 31May2022.

After installing the [Intel oneAPI](#) base toolkit and the HPC toolkit, it is also possible to compile large parts of LAMMPS natively on Windows using Intel compilers. The HPC toolkit provides two sets of C/C++ and Fortran compilers: the so-called “classic” compilers (`icl.exe` and `ifort.exe`) and newer, LLVM based compilers (`icx.exe` and `ifx.exe`). In addition to the compilers and their dependent modules, also the thread building blocks (TBB) and the math kernel library (MKL) need to be installed. Two presets (`cmake/presets/windows-intel-llvm.cmake` and `cmake/presets/windows-intel-classic.cmake`) are provided for selecting the LLVM based or classic compilers, respectively. The preset `cmake/presets/windows.cmake` enables compatible packages that are not dependent on additional features or libraries. You **must** use the CMake based build procedure and use Ninja as build tool. For compiling from the command prompt, thus both `CMake` and `Ninja-build` binaries must be installed. It is also possible to use Visual Studio, if it is started (`devenv.exe`) from a command prompt that has the Intel oneAPI compilers enabled. The Visual Studio settings file in the `cmake` folder contains configurations for both compiler variants in debug and release settings. Those will use the CMake and Ninja binaries bundled with Visual Studio, thus a separate installation is not required.

---

#### Known Limitations

In addition to portability issues with several packages and external libraries, the classic Intel compilers are currently not able to compile the googletest libraries and thus enabling the `-DENABLE_TESTING` option will result in compilation failure. The LLVM based compilers are compatible.

---

---

**Note:** This is work in progress and you should contact the LAMMPS developers via [GitHub](#) or the [LAMMPS forum at MatSci](#), if you have questions or LAMMPS specific problems.

---

### 3.9.6 Using a cross-compiler

If you need to provide custom LAMMPS binaries for Windows, but do not need to do the compilation on Windows, please consider using a Linux to Windows cross-compiler. This is how currently the Windows binary packages are created by the LAMMPS developers. Because of that, this is probably the currently best tested and supported way to build LAMMPS executables for Windows. A CMake preset selecting all packages compatible with this cross-compilation build is provided. The GPU package can only be compiled with OpenCL support. To compile with MPI support, a pre-compiled library and the corresponding header files are required. When building with CMake the matching package will be downloaded automatically, but MPI support has to be explicitly enabled with `-DBUILD_MPI=on`.

Please keep in mind, though, that this only applies to **compiling** LAMMPS. Whether the resulting binaries do work correctly is rarely tested by the LAMMPS developers. We instead rely on the feedback of the users of these pre-compiled LAMMPS packages for Windows. We will try to resolve issues to the best of our abilities if we become aware of them. However this is subject to time constraints and focus on HPC platforms.

## 3.10 Notes for saving disk space when building LAMMPS from source

LAMMPS is a large software project with a large number of source files, extensive documentation, and a large collection of example files. When downloading LAMMPS by cloning the [git repository from GitHub](#) this will by default also download the entire commit history since September 2006. Compiling LAMMPS will add the storage requirements of the compiled object files and libraries to the tally.

In a user account on an HPC cluster with filesystem quotas or in other environments with restricted disk space capacity it may be needed to reduce the storage requirements. Here are some suggestions:

- Create a so-called shallow repository by cloning only the last commit instead of the full project history by using `git clone git@github.com:lammops/lammops --depth=1 --branch=develop`. This reduces the downloaded size to about half. With `--depth=1` it is not possible to check out different versions/branches of LAMMPS, using `--depth=1000` will make multiple recent versions available at little extra storage needs (the entire git history had nearly 30,000 commits in fall 2021).
- Download a tar archive from either the [download section on the LAMMPS homepage](#) or from the [LAMMPS releases page on GitHub](#) these will not contain the git history at all.
- Build LAMMPS without the debug flag (remove `-g` from the machine makefile or use `-DCMAKE_BUILD_TYPE=Release`) or use the `strip` command on the LAMMPS executable when no more debugging would be needed. The `strip` command may also be applied to the LAMMPS shared library. The static library may be deleted entirely.
- Delete compiled object files and libraries after copying the LAMMPS executable to a permanent location. When using the traditional build process, one may use `make clean-<machine>` or `make clean-all` to delete object files in the `src` folder. For CMake based builds, one may use `make clean` or just delete the entire build folder.
- The folders containing the documentation tree (`doc`), the examples (`examples`) are not needed to build and run LAMMPS and can be safely deleted. Some files in the `potentials` folder are large and may be deleted, if not needed. The largest of those files (occupying about 120 MBytes combined) will only be downloaded on demand, when the corresponding package is installed.
- When using the CMake build procedure, the compilation can be done on a (local) scratch storage that will not count toward the quota. A local scratch file system may offer the additional benefit of speeding up creating object files and linking with libraries compared to a networked file system. Also with CMake (and unlike with the traditional `make`) it is possible to compile LAMMPS executables with different settings and packages included from the same source tree since all the configuration information is stored in the build folder. So it is not necessary to have multiple copies of LAMMPS.

## 3.11 Development build options

The build procedures in LAMMPS offers a few extra options which are useful during development, testing or debugging.

---

### 3.11.1 Monitor compilation flags (CMake only)

Sometimes it is necessary to verify the complete sequence of compilation flags generated by the CMake build. To enable a more verbose output during compilation you can use the following option.

```
-D CMAKE_VERBOSE_MAKEFILE=value    # value = no (default) or yes
```

Another way of doing this without reconfiguration is calling make with variable VERBOSE set to 1:

```
make VERBOSE=1
```

---

### 3.11.2 Enable static code analysis with clang-tidy (CMake only)

The `clang-tidy` tool is a static code analysis tool to diagnose (and potentially fix) typical programming errors or coding style violations. It has a modular framework of tests that can be adjusted to help identifying problems before they become bugs and also assist in modernizing large code bases (like LAMMPS). It can be enabled for all C++ code with the following CMake flag

```
-D ENABLE_CLANG_TIDY=value    # value = no (default) or yes
```

With this flag enabled all source files will be processed twice, first to be compiled and then to be analyzed. Please note that the analysis can be significantly more time consuming than the compilation itself.

---

### 3.11.3 Report missing and unneeded '#include' statements (CMake only)

The conventions for how and when to use and order include statements in LAMMPS are documented in *LAMMPS programming style and requirements for contributions*. To assist with following these conventions one can use the `Include What You Use` tool. This tool is still under development and for large and complex projects like LAMMPS there are some false positives, so suggested changes need to be verified manually. It is recommended to use at least version 0.16, which has much fewer incorrect reports than earlier versions. To install the IWYU toolkit, you need to have the clang compiler **and** its development package installed. Download the IWYU version that matches the version of the clang compiler, configure, build, and install it.

The necessary steps to generate the report can be enabled via a CMake variable during CMake configuration.

```
-D ENABLE_IWYU=value    # value = no (default) or yes
```

This will check if the required binary (`include-what-you-use` or `iwyu`) and python script (`iwyu-tool` or `iwyu_tool.py`) can be found in the path. The analysis can then be started with:

```
make iwyu
```

---

This may first run some compilation, as the analysis is dependent on recording all commands required to do the compilation.

### 3.11.4 Address, Undefined Behavior, and Thread Sanitizer Support (CMake only)

Compilers such as GCC and Clang support generating instrumented binaries which use different sanitizer libraries to detect problems in the code during run-time. They can detect issues like:

- memory leaks
- undefined behavior
- data races

Please note that this kind of instrumentation usually comes with a performance hit (but much less than using tools like Valgrind with a more low level approach). To enable these features, additional compiler flags need to be added to the compilation and linking stages. This is done through setting the `ENABLE_SANITIZER` variable during configuration. Examples:

```
-D ENABLE_SANITIZER=none      # no sanitizer active (default)
-D ENABLE_SANITIZER=address  # enable address sanitizer / memory leak checker
-D ENABLE_SANITIZER=leak     # enable memory leak checker (only)
-D ENABLE_SANITIZER=undefined # enable undefined behavior sanitizer
-D ENABLE_SANITIZER=thread   # enable thread sanitizer
```

### 3.11.5 Code Coverage and Unit Testing (CMake only)

The LAMMPS code is subject to multiple levels of automated testing during development: integration testing (i.e. whether the code compiles on various platforms and with a variety of settings), unit testing (i.e. whether certain individual parts of the code produce the expected results for given inputs), run testing (whether selected complete input decks run without crashing for multiple configurations), and regression testing (i.e. whether selected input examples reproduce the same results over a given number of steps and operations within a given error margin). The status of this automated testing can be viewed on <https://ci.lammps.org>.

The scripts and inputs for integration, run, and regression testing are maintained in a [separate repository](#) of the LAMMPS project on GitHub.

The unit testing facility is integrated into the CMake build process of the LAMMPS source code distribution itself. It can be enabled by setting `-D ENABLE_TESTING=on` during the CMake configuration step. It requires the [YAML](#) library and development headers (if those are not found locally a recent version will be downloaded and compiled along with LAMMPS and the test program) to compile and will download and compile a specific recent version of the [Googletest](#) C++ test framework for implementing the tests.

After compilation is complete, the unit testing is started in the build folder using the `ctest` command, which is part of the CMake software. The output of this command will be looking something like this:

```
[...]$ ctest
Test project /home/akohlmeier/compile/lammps/build-testing
   Start  1: MolPairStyle:hybrid-overlay
1/109 Test  #1: MolPairStyle:hybrid-overlay ..... Passed    0.02 sec
   Start  2: MolPairStyle:hybrid
```

(continues on next page)

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```

2/109 Test #2: MolPairStyle:hybrid ..... Passed 0.01 sec
      Start 3: MolPairStyle:lj_class2
      [...]
      Start 107: PotentialFileReader
107/109 Test #107: PotentialFileReader ..... Passed 0.04 sec
      Start 108: EIMPotentialFileReader
108/109 Test #108: EIMPotentialFileReader ..... Passed 0.03 sec
      Start 109: TestSimpleCommands
109/109 Test #109: TestSimpleCommands ..... Passed 0.02 sec

100% tests passed, 0 tests failed out of 26

Total Test time (real) = 25.57 sec

```

The ctest command has many options, the most important ones are:

Option	Function
-V	verbose output: display output of individual test runs
-j <num>	parallel run: run <num> tests in parallel
-R <regex>	run subset of tests matching the regular expression <regex>
-E <regex>	exclude subset of tests matching the regular expression <regex>
-L <regex>	run subset of tests with a label matching the regular expression <regex>
-LE <regex>	exclude subset of tests with a label matching the regular expression <regex>
-N	dry-run: display list of tests without running them
-T memcheck	run tests with valgrind memory checker (if available)

In its full implementation, the unit test framework will consist of multiple kinds of tests implemented in different programming languages (C++, C, Python, Fortran) and testing different aspects of the LAMMPS software and its features. The tests will adapt to the compilation settings of LAMMPS, so that tests will be skipped if prerequisite features are not available in LAMMPS.

---

**Note:** The unit test framework was added in spring 2020 and is under active development. The coverage is not complete and will be expanded over time.

---

Tests for styles of the same kind of style (e.g. pair styles or bond styles) are performed with the same test executable using different input files in YAML format. So to add a test for another style of the same kind it may be sufficient to add a suitable YAML file. *Detailed instructions for adding tests* are provided in the Programmer Guide part of the manual.

### Unit tests for force styles

A large part of LAMMPS are different “styles” for computing non-bonded and bonded interactions selected through the *pair\_style command*, *bond\_style command*, *angle\_style command*, *dihedral\_style command*, *improper\_style command*, and *kpace\_style command*. Since these all share common interfaces, it is possible to write generic test programs that will call those common interfaces for small test systems with less than 100 atoms and compare the results with pre-recorded reference results. A test run is then a collection multiple individual test runs each with many comparisons to reference results based on template input files, individual command settings, relative error margins, and reference data stored in a YAML format file with *.yaml* suffix. Currently the programs *test\_pair\_style*, *test\_bond\_style*, and *test\_angle\_style* are implemented. They will compare forces, energies and (global) stress for all atoms after a run 0 calculation and after a few steps of MD with *fix nve*, each in multiple variants with different settings and also for multiple accelerated styles. If a prerequisite style or package is missing, the individual tests are skipped. All tests



will be executed on a single MPI process, so using the CMake option `-D BUILD_MPI=off` can significantly speed up testing, since this will skip the MPI initialization for each test run. Below is an example command and output:

```
[tests]$ test_pair_style mol-pair-lj_cut.yaml
[=====  
[-----] Global test environment set-up.  
[-----] 6 tests from PairStyle  
[ RUN      ] PairStyle.plain  
[      OK ] PairStyle.plain (24 ms)  
[ RUN      ] PairStyle.omp  
[      OK ] PairStyle.omp (18 ms)  
[ RUN      ] PairStyle.intel  
[      OK ] PairStyle.intel (6 ms)  
[ RUN      ] PairStyle.opt  
[ SKIPPED ] PairStyle.opt (0 ms)  
[ RUN      ] PairStyle.single  
[      OK ] PairStyle.single (7 ms)  
[ RUN      ] PairStyle.extract  
[      OK ] PairStyle.extract (6 ms)  
[-----] 6 tests from PairStyle (62 ms total)  
  
[-----] Global test environment tear-down  
[=====  
[ PASSED ] 5 tests.  
[ SKIPPED ] 1 test, listed below:  
[ SKIPPED ] PairStyle.opt
```

In this particular case, 5 out of 6 sets of tests were conducted, the tests for the `lj/cut/opt` pair style was skipped, since the tests executable did not include it. To learn what individual tests are performed, you (currently) need to read the source code. You can use code coverage recording (see next section) to confirm how well the tests cover the code paths in the individual source files.

The force style test programs have a common set of options:

Option	Function
<code>-g &lt;newfile&gt;</code>	regenerate reference data in new YAML file
<code>-u</code>	update reference data in the original YAML file
<code>-s</code>	print error statistics for each group of comparisons
<code>-v</code>	verbose output: also print the executed LAMMPS commands

The `ctest` tool has no mechanism to directly pass flags to the individual test programs, but a workaround has been implemented where these flags can be set in an environment variable `TEST_ARGS`. Example:

```
env TEST_ARGS=-s ctest -V -R BondStyle
```

To add a test for a style that is not yet covered, it is usually best to copy a YAML file for a similar style to a new file, edit the details of the style (how to call it, how to set its coefficients) and then run test command with either the `-g` and the replace the initial test file with the regenerated one or the `-u` option. The `-u` option will destroy the original file, if the generation run does not complete, so using `-g` is recommended unless the YAML file is fully tested and working.

Some of the force style tests are rather slow to run and some are very sensitive to small differences like CPU architecture, compiler toolchain, compiler optimization. Those tests are flagged with a “slow” and/or “unstable” label, and thus those tests can be selectively excluded with the `-LE` flag or selected with the `-L` flag.

### Recommendations and notes for YAML files

- The reference results should be recorded without any code optimization or related compiler flags enabled.
  - The `epsilon` parameter defines the relative precision with which the reference results must be met. The test geometries often have high and low energy parts and thus a significant impact from floating-point math truncation errors is to be expected. Some functional forms and potentials are more noisy than others, so this parameter needs to be adjusted. Typically a value around  $1.0e-13$  can be used, but it may need to be as large as  $1.0e-8$  in some cases.
  - The tests for pair styles from OPT, OPENMP and INTEL are performed with automatically rescaled epsilon to account for additional loss of precision from code optimizations and different summation orders.
  - When compiling with (aggressive) compiler optimization, some tests are likely to fail. It is recommended to inspect the individual tests in detail to decide, whether the specific error for a specific property is acceptable (it often is), or this may be an indication of mis-compiled code (or an undesired large loss of precision due to significant reordering of operations and thus less error cancellation).
- 

### Unit tests for timestepping related fixes

A substantial subset of *fix styles* are invoked regularly during MD timestepping and manipulate per-atom properties like positions, velocities, and forces. For those fix styles, testing can be done in a very similar fashion as for force fields and thus there is a test program `test_fix_timestep` that shares a lot of code, properties, and command line flags with the force field style testers described in the previous section.

This tester will set up a small molecular system run with verlet run style for 4 MD steps, then write a binary restart and continue for another 4 MD steps. At this point coordinates and velocities are recorded and compared to reference data. Then the system is cleared, restarted and running the second 4 MD steps again and the data is compared to the same reference. That is followed by another restart after which per atom type masses are replaced with per-atom masses and the second 4 MD steps are repeated again and compared to the same reference. Also global scalar and vector data of the fix is recorded and compared. If the fix is a thermostat and thus the internal property `t_target` can be extracted, then this is compared to the reference data. The tests are repeated with the respa run style.

If the fix has a multi-threaded version in the OPENMP package, then the entire set of tests is repeated for that version as well.

For this to work, some additional conditions have to be met by the YAML format test inputs.

- The fix to be tested (and only this fix), should be listed in the `prerequisites:` section
- The fix to be tested must be specified in the `post_commands:` section with the fix-ID test. This section may contain other commands and other fixes (e.g. an instance of `fix nve` for testing a thermostat or force manipulation fix)
- For fixes that can tally contributions to the global virial, the line `fix_modify test virial yes` should be included in the `post_commands:` section of the test input.
- For thermostat fixes the target temperature should be ramped from an arbitrary value (e.g. 50K) to a pre-defined target temperature entered as `${t_target}`.
- For fixes that have thermostatting support included, but do not have it enabled in the input (e.g. `fix rigid` with default settings), the `post_commands:` section should contain the line `variable t_target delete` to disable the target temperature ramp check to avoid false positives.

## Use custom linker for faster link times when ENABLE\_TESTING is active

When compiling LAMMPS with enabled tests, most test executables will need to be linked against the LAMMPS library. Since this can be a very large library with many C++ objects when many packages are enabled, link times can become very long on machines that use the GNU BFD linker (e.g. Linux systems). Alternatives like the lld linker of the LLVM project or the gold linker available with GNU binutils can speed up this step substantially. CMake will by default test if any of the two can be enabled and use it when ENABLE\_TESTING is active. It can also be selected manually through the CMAKE\_CUSTOM\_LINKER CMake variable. Allowed values are lld, gold, bfd, or default. The default option will use the system default linker otherwise, the linker is chosen explicitly. This option is only available for the GNU or Clang C++ compiler.

## Tests for other components and utility functions

Additional tests that validate utility functions or specific components of LAMMPS are implemented as standalone executable which may, or may not require creating a suitable LAMMPS instance. These tests are more specific and do not require YAML format input files. To add a test, either an existing source file needs to be extended or a new file added, which in turn requires additions to the CMakeLists.txt file in the source folder.

## Collect and visualize code coverage metrics

You can also collect code coverage metrics while running LAMMPS or the tests by enabling code coverage support during the CMake configuration:

```
-D ENABLE_COVERAGE=on # enable coverage measurements (off by default)
```

This will instrument all object files to write information about which lines of code were accessed during execution in files next to the corresponding object files. These can be post-processed to visually show the degree of coverage and which code paths are accessed and which are not taken. When working on unit tests (see above), this can be extremely helpful to determine which parts of the code are not executed and thus what kind of tests are still missing. The coverage data is cumulative, i.e. new data is added with each new run.

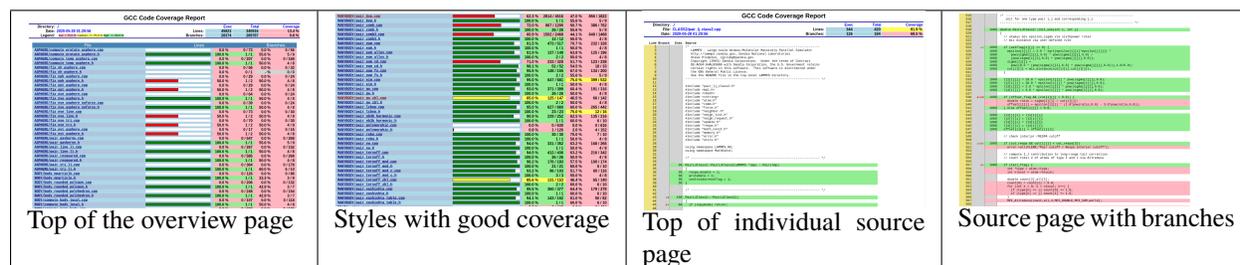
Enabling code coverage will also add the following build targets to generate coverage reports after running the LAMMPS executable or the unit tests:

```
make gen_coverage_html # generate coverage report in HTML format
make gen_coverage_xml # generate coverage report in XML format
make clean_coverage_html # delete folder with HTML format coverage report
make reset_coverage # delete all collected coverage data and HTML output
```

These reports require GCOVR to be installed. The easiest way to do this to install it via pip:

```
pip install git+https://github.com/gcovr/gcovr.git
```

After post-processing with gen\_coverage\_html the results are in a folder coverage\_html and can be viewed with a web browser. The images below illustrate how the data is presented.



### 3.11.6 Coding style utilities

To aid with enforcing some of the coding style conventions in LAMMPS some additional build targets have been added. These require Python 3.5 or later and will only work properly on Unix-like operating and file systems.

The following options are available.

```
make check-whitespace  # search for files with whitespace issues
make fix-whitespace    # correct whitespace issues in files
make check-homepage    # search for files with old LAMMPS homepage URLs
make fix-homepage      # correct LAMMPS homepage URLs in files
make check-errordocs   # search for deprecated error docs in header files
make fix-errordocs     # remove error docs in header files
make check-permissions # search for files with permissions issues
make fix-permissions   # correct permissions issues in files
make check             # run all check targets from above
```

These should help to make source and documentation files conforming to some the coding style preferences of the LAMMPS developers.

### 3.11.7 Clang-format support

For the code in the `unittest` and `src` trees we are transitioning to use the `clang-format` tool to assist with having a consistent source code formatting style. The `clang-format` command bundled with Clang version 8.0 or later is required. The configuration is in files called `.clang-format` in the respective folders. Since the modifications from `clang-format` can be significant and - especially for “legacy style code” - they are not always improving readability, a large number of files currently have a `// clang-format off` at the top, which will disable the processing. As of fall 2021 all files have been either “protected” this way or are enabled for full or partial `clang-format` processing. Over time, the “protected” files will be refactored and updated so that `clang-format` may be applied to them as well.

It is recommended for all newly contributed files to use the `clang-format` processing while writing the code or do the coding style processing (including the scripts mentioned in the previous paragraph)

If `clang-format` is available, files can be updated individually with commands like the following:

```
$ clang-format -i some_file.cpp
```

The following target are available for both, GNU make and CMake:

```
make format-src      # apply clang-format to all files in src and the package folders
make format-tests    # apply clang-format to all files in the unittest tree
```

## RUN LAMMPS

These pages explain how to run LAMMPS once you have *installed an executable* or *downloaded the source code and built an executable*. The *Commands* doc page describes how input scripts are structured and the commands they can contain.

### 4.1 Basics of running LAMMPS

LAMMPS is run from the command line, reading commands from a file via the `-in` command line flag, or from standard input. Using the “`-in in.file`” variant is recommended (see note below). The name of the LAMMPS executable is either `lmp` or `lmp_<machine>` with `<machine>` being the machine string used when compiling LAMMPS. This is required when compiling LAMMPS with the traditional build system (e.g. with `make mpi`), but optional when using CMake to configure and build LAMMPS:

```
$ lmp_serial -in in.file
$ lmp_serial < in.file
$ lmp -in in.file
$ lmp < in.file
$ /path/to/lammps/src/lmp_serial -i in.file
$ mpirun -np 4 lmp_mpi -in in.file
$ mpiexec -np 4 lmp -in in.file
$ mpirun -np 8 /path/to/lammps/src/lmp_mpi -in in.file
$ mpiexec -n 6 /usr/local/bin/lmp -in in.file
```

You normally run the LAMMPS command in the directory where your input script is located. That is also where output files are produced by default, unless you provide specific other paths in your input script or on the command line. As in some of the examples above, the LAMMPS executable itself can be placed elsewhere.

---

**Note:** The redirection operator “`<`” will not always work when running in parallel with `mpirun` or `mpiexec`; for those systems the `-in` form is required.

---

As LAMMPS runs it prints info to the screen and a logfile named `log.lammps`. More info about output is given on the [screen and logfile output](#) page.

If LAMMPS encounters errors in the input script or while running a simulation it will print an ERROR message and stop or a WARNING message and continue. See the [Common Problems](#) page for a discussion of the various kinds of errors LAMMPS can or can’t detect, a list of all ERROR and WARNING messages, and what to do about them.

---

LAMMPS can run the same problem on any number of processors, including a single processor. In theory you should get identical answers on any number of processors and on any machine. In practice, numerical round-off due to using

floating-point math can cause slight differences and an eventual divergence of molecular dynamics trajectories. See the *Errors common* page for discussion of this.

LAMMPS can run as large a problem as will fit in the physical memory of one or more processors. If you run out of memory, you must run on more processors or define a smaller problem. The amount of memory needed and how well it can be distributed across processors may vary based on the models and settings and commands used.

If you run LAMMPS in parallel via `mpirun`, you should be aware of the *processors* command, which controls how MPI tasks are mapped to the simulation box, as well as `mpirun` options that control how MPI tasks are assigned to physical cores of the node(s) of the machine you are running on. These settings can improve performance, though the defaults are often adequate.

For example, it is often important to bind MPI tasks (processes) to physical cores (processor affinity), so that the operating system does not migrate them during a simulation. If this is not the default behavior on your machine, the `mpirun` option “`-bind-to core`” (OpenMPI) or “`-bind-to core`” (MPICH) can be used.

If the LAMMPS command(s) you are using support multi-threading, you can set the number of threads per MPI task via the environment variable `OMP_NUM_THREADS`, before you launch LAMMPS:

```
$ export OMP_NUM_THREADS=2      # bash
$ setenv OMP_NUM_THREADS 2      # csh or tcsh
```

This can also be done via the *package* command or via the *-pk command-line switch* which invokes the package command. See the *package* command or *Speed* doc pages for more details about which accelerator packages and which commands support multi-threading.

---

You can experiment with running LAMMPS using any of the input scripts provided in the examples or bench directory. Input scripts are named `in.*` and sample outputs are named `log.*.P` where P is the number of processors it was run on.

Some of the examples or benchmarks require LAMMPS to be built with optional packages.

## 4.2 Command-line options

At run time, LAMMPS recognizes several optional command-line switches which may be used in any order. Either the full word or a one or two letter abbreviation can be used:

- *-e or -echo*
- *-h or -help*
- *-i or -in*
- *-k or -kokkos*
- *-l or -log*
- *-mdi*
- *-m or -mpicolor*
- *-c or -cite*
- *-nc or -nocite*
- *-nb or -nonbuf*
- *-pk or -package*
- *-p or -partition*

- *-pl or -plog*
- *-ps or -pscreen*
- *-ro or -reorder*
- *-r2data or -restart2data*
- *-r2dump or -restart2dump*
- *-sc or -screen*
- *-sr or skiprun*
- *-sf or -suffix*
- *-v or -var*

For example, the `lmp_mpi` executable might be launched as follows:

```
$ mpirun -np 16 lmp_mpi -v f tmp.out -l my.log -sc none -i in.alloy
$ mpirun -np 16 lmp_mpi -var f tmp.out -log my.log -screen none -in in.alloy
```

---

### **-echo style**

Set the style of command echoing. The style can be *none* or *screen* or *log* or *both*. Depending on the style, each command read from the input script will be echoed to the screen and/or logfile. This can be useful to figure out which line of your script is causing an input error. The default value is *log*. The echo style can also be set by using the *echo* command in the input script itself.

---

### **-help**

Print a brief help summary and a list of options compiled into this executable for each LAMMPS style (*atom\_style*, *fix*, *compute*, *pair\_style*, *bond\_style*, etc). This can tell you if the command you want to use was included via I/O redirection; e.g. `lmp_linux < in.run`. With many MPI implementations I/O redirection also works in parallel, but using the `-in` flag will always work.

---

### **-in file**

Specify a file to use as an input script. This is an optional but recommended switch when running LAMMPS in one-partition mode. If it is not specified, LAMMPS reads its script from standard input, typically from a script via I/O redirection; e.g. `lmp_linux < in.run`. With many MPI implementations I/O redirection also works in parallel, but using the `-in` flag will always work.

Note that this is a required switch when running LAMMPS in multi-partition mode, since multiple processors cannot all read from stdin concurrently. The file name may be “none” for starting multi-partition calculations without reading an initial input file from the library interface.

---

### **-kokkos on/off keyword/value ...**

Explicitly enable or disable KOKKOS support, as provided by the KOKKOS package. Even if LAMMPS is built with this package, as described in the *the KOKKOS package page*, this switch must be set to enable running with KOKKOS-enabled styles the package provides. If the switch is not set (the default), LAMMPS will operate as if the KOKKOS package were not installed; i.e. you can run standard LAMMPS or with the GPU or OPENMP packages, for testing or benchmarking purposes.

Additional optional keyword/value pairs can be specified which determine how Kokkos will use the underlying hardware on your platform. These settings apply to each MPI task you launch via the “`mpirun`” or “`mpiexec`” command. You

may choose to run one or more MPI tasks per physical node. Note that if you are running on a desktop machine, you typically have one physical node. On a cluster or supercomputer there may be dozens or 1000s of physical nodes.

Either the full word or an abbreviation can be used for the keywords. Note that the keywords do not use a leading minus sign. I.e. the keyword is “t”, not “-t”. Also note that each of the keywords has a default setting. Examples of when to use these options and what settings to use on different platforms is given on the *KOKKOS package* doc page.

- d or device
- g or gpus
- t or threads
- n or numa

device Nd

This option is only relevant if you built LAMMPS with CUDA=yes, you have more than one GPU per node, and if you are running with only one MPI task per node. The Nd setting is the ID of the GPU on the node to run on. By default Nd = 0. If you have multiple GPUs per node, they have consecutive IDs numbered as 0,1,2,etc. This setting allows you to launch multiple independent jobs on the node, each with a single MPI task per node, and assign each job to run on a different GPU.

gpus Ng Ns

This option is only relevant if you built LAMMPS with CUDA=yes, you have more than one GPU per node, and you are running with multiple MPI tasks per node (up to one per GPU). The Ng setting is how many GPUs you will use. The Ns setting is optional. If set, it is the ID of a GPU to skip when assigning MPI tasks to GPUs. This may be useful if your desktop system reserves one GPU to drive the screen and the rest are intended for computational work like running LAMMPS. By default Ng = 1 and Ns is not set.

Depending on which flavor of MPI you are running, LAMMPS will look for one of these 4 environment variables

SLURM\_LOCALID (various MPI variants compiled with SLURM support)  
MPT\_LRANK (HPE MPI)  
MV2\_COMM\_WORLD\_LOCAL\_RANK (Mvapich)  
OMPI\_COMM\_WORLD\_LOCAL\_RANK (OpenMPI)

which are initialized by the “srun”, “mpirun” or “mpiexec” commands. The environment variable setting for each MPI rank is used to assign a unique GPU ID to the MPI task.

threads Nt

This option assigns Nt number of threads to each MPI task for performing work when Kokkos is executing in OpenMP or pthreads mode. The default is Nt = 1, which essentially runs in MPI-only mode. If there are Np MPI tasks per physical node, you generally want Np\*Nt = the number of physical cores per node, to use your available hardware optimally. This also sets the number of threads used by the host when LAMMPS is compiled with CUDA=yes.

numa Nm

This option is only relevant when using pthreads with hwloc support. In this case Nm defines the number of NUMA regions (typically sockets) on a node which will be utilized by a single MPI rank. By default Nm = 1. If this option is used the total number of worker-threads per MPI rank is threads\*numa. Currently it is always almost better to assign at least one MPI rank per NUMA region, and leave numa set to its default value of 1. This is because letting a single process span multiple NUMA regions induces a significant amount of cross NUMA data traffic which is slow.

---

**-log file**



Specify a log file for LAMMPS to write status information to. In one-partition mode, if the switch is not used, LAMMPS writes to the file `log.lammps`. If this switch is used, LAMMPS writes to the specified file. In multi-partition mode, if the switch is not used, a `log.lammps` file is created with high-level status information. Each partition also writes to a `log.lammps.N` file where `N` is the partition ID. If the switch is specified in multi-partition mode, the high-level logfile is named “file” and each partition also logs information to a `file.N`. For both one-partition and multi-partition mode, if the specified file is “none”, then no log files are created. Using a *log* command in the input script will override this setting. Option `-plog` will override the name of the partition log files `file.N`.

---

#### **-mdi ‘multiple flags’**

This flag is only recognized and used when LAMMPS has support for the MolSSI Driver Interface (MDI) included as part of the *MDI* package. This flag is specific to the MDI library and controls how LAMMPS interacts with MDI. There are usually multiple flags that have to follow it and those have to be placed in quotation marks. For more information about how to launch LAMMPS in MDI client/server mode please refer to the *MDI Howto*.

---

#### **-mpicolor color**

If used, this must be the first command-line argument after the LAMMPS executable name. It is only used when LAMMPS is launched by an `mpirun` command which also launches another executable(s) at the same time. (The other executable could be LAMMPS as well.) The color is an integer value which should be different for each executable (another application may set this value in a different way). LAMMPS and the other executable(s) perform an `MPI_Comm_split()` with their own colors to shrink the `MPI_COMM_WORLD` communication to be the subset of processors they are actually running on.

---

#### **-cite style or file name**

Select how and where to output a reminder about citing contributions to the LAMMPS code that were used during the run. Available keywords for styles are “both”, “none”, “screen”, or “log”. Any other keyword will be considered a file name to write the detailed citation info to instead of logfile or screen. Default is the “log” style where there is a short summary in the screen output and detailed citations in BibTeX format in the logfile. The option “both” selects the detailed output for both, “none”, the short output for both, and “screen” will write the detailed info to the screen and the short version to the log file. If a dedicated citation info file is requested, the screen and log file output will be in the short format (same as with “none”).

See the *citation page* for more details on how to correctly reference and cite LAMMPS.

---

#### **-nocite**

Disable generating a citation reminder (see above) at all.

---

#### **-nonbuf**

Turn off buffering for screen and logfile output. For performance reasons, output to the screen and logfile is usually buffered, i.e. output is only written to a file if its buffer - typically 4096 bytes - has been filled. When LAMMPS crashes for some reason, however, that can mean that there is important output missing. With this flag the buffering can be turned off (only for screen and logfile output) and any output will be committed immediately. Note that when running in parallel with MPI, the screen output may still be buffered by the MPI library and this cannot be changed by LAMMPS. This flag should only be used for debugging and not for production simulations as the performance impact can be significant, especially for large parallel runs.

---

### **-package style args ....**

Invoke the *package* command with style and args. The syntax is the same as if the command appeared at the top of the input script. For example “-package gpu 2” or “-pk gpu 2” is the same as *package gpu 2* in the input script. The possible styles and args are documented on the *package* doc page. This switch can be used multiple times, e.g. to set options for the INTEL and OPENMP packages which can be used together.

Along with the “-suffix” command-line switch, this is a convenient mechanism for invoking accelerator packages and their options without having to edit an input script.

---

### **-partition 8x2 4 5 ...**

Invoke LAMMPS in multi-partition mode. When LAMMPS is run on P processors and this switch is not used, LAMMPS runs in one partition, i.e. all P processors run a single simulation. If this switch is used, the P processors are split into separate partitions and each partition runs its own simulation. The arguments to the switch specify the number of processors in each partition. Arguments of the form MxN mean M partitions, each with N processors. Arguments of the form N mean a single partition with N processors. The sum of processors in all partitions must equal P. Thus the command “-partition 8x2 4 5” has 10 partitions and runs on a total of 25 processors.

Running with multiple partitions can be useful for running *multi-replica simulations*, where each replica runs on one or a few processors. Note that with MPI installed on a machine (e.g. your desktop), you can run on more (virtual) processors than you have physical processors.

To run multiple independent simulations from one input script, using multiple partitions, see the *Howto multiple* page. World- and universe-style *variables* are useful in this context.

---

### **-plog file**

Specify the base name for the partition log files, so partition N writes log information to file.N. If file is none, then no partition log files are created. This overrides the filename specified in the -log command-line option. This option is useful when working with large numbers of partitions, allowing the partition log files to be suppressed (-plog none) or placed in a sub-directory (-plog replica\_files/log.lammps) If this option is not used the log file for partition N is log.lammps.N or whatever is specified by the -log command-line option.

---

### **-pscreen file**

Specify the base name for the partition screen file, so partition N writes screen information to file.N. If file is “none”, then no partition screen files are created. This overrides the filename specified in the -screen command-line option. This option is useful when working with large numbers of partitions, allowing the partition screen files to be suppressed (-pscreen none) or placed in a sub-directory (-pscreen replica\_files/screen). If this option is not used the screen file for partition N is screen.N or whatever is specified by the -screen command-line option.

---

### **-reorder**

This option has 2 forms:

```
-reorder nth N
-reorder custom filename
```

Reorder the processors in the MPI communicator used to instantiate LAMMPS, in one of several ways. The original MPI communicator ranks all P processors from 0 to P-1. The mapping of these ranks to physical processors is done by MPI before LAMMPS begins. It may be useful in some cases to alter the rank order. E.g. to insure that cores within each node are ranked in a desired order. Or when using the *run\_style verlet/split* command with 2 partitions to insure

---

that a specific Kspace processor (in the second partition) is matched up with a specific set of processors in the first partition. See the *General tips* page for more details.

If the keyword *nth* is used with a setting *N*, then it means every *N*th processor will be moved to the end of the ranking. This is useful when using the *run\_style verlet/split* command with 2 partitions via the *-partition* command-line switch. The first set of processors will be in the first partition, the second set in the second partition. The *-reorder* command-line switch can alter this so that the first *N* procs in the first partition and one proc in the second partition will be ordered consecutively, e.g. as the cores on one physical node. This can boost performance. For example, if you use “*-reorder nth 4*” and “*-partition 9 3*” and you are running on 12 processors, the processors will be reordered from

```
0 1 2 3 4 5 6 7 8 9 10 11
```

to

```
0 1 2 4 5 6 8 9 10 3 7 11
```

so that the processors in each partition will be

```
0 1 2 4 5 6 8 9 10
3 7 11
```

See the “processors” command for how to insure processors from each partition could then be grouped optimally for quad-core nodes.

If the keyword is *custom*, then a file that specifies a permutation of the processor ranks is also specified. The format of the reorder file is as follows. Any number of initial blank or comment lines (starting with a “#” character) can be present. These should be followed by *P* lines of the form:

```
I J
```

where *P* is the number of processors LAMMPS was launched with. Note that if running in multi-partition mode (see the *-partition* switch above) *P* is the total number of processors in all partitions. The *I* and *J* values describe a permutation of the *P* processors. Every *I* and *J* should be values from 0 to *P*-1 inclusive. In the set of *P* *I* values, every proc ID should appear exactly once. Ditto for the set of *P* *J* values. A single *I,J* pairing means that the physical processor with rank *I* in the original MPI communicator will have rank *J* in the reordered communicator.

Note that rank ordering can also be specified by many MPI implementations, either by environment variables that specify how to order physical processors, or by config files that specify what physical processors to assign to each MPI rank. The *-reorder* switch simply gives you a portable way to do this without relying on MPI itself. See the *processors file* command for how to output info on the final assignment of physical processors to the LAMMPS simulation domain.

#### **-restart2data restartfile [remap] datafile keyword value ...**

Convert the restart file into a data file and immediately exit. This is the same operation as if the following 2-line input script were run:

```
read_restart restartfile [remap]
write_data datafile keyword value ...
```

The specified restartfile and/or datafile name may contain the wild-card character “\*”. The restartfile name may also contain the wild-card character “%”. The meaning of these characters is explained on the *read\_restart* and *write\_data* doc pages. The use of “%” means that a parallel restart file can be read. Note that a filename such as file.\* may need to be enclosed in quotes or the “\*” character prefixed with a backslash (“\”) to avoid shell expansion of the “\*” character.

Following restartfile argument, the optional word “remap” may be used. This has the same effect like adding it to a *read\_restart* command, and operates as explained on its doc page. This is useful if reading the restart file triggers an error that atoms have been lost. In that case, use of the remap flag should allow the data file to still be produced.

The syntax following restartfile (or remap), namely

```
datafile keyword value ...
```

is identical to the arguments of the *write\_data* command. See its page for details. This includes its optional keyword/value settings.

---

#### **-restart2dump restartfile [remap] group-ID dumpstyle dumpfile arg1 arg2 ...**

Convert the restart file into a dump file and immediately exit. This is the same operation as if the following 2-line input script were run:

```
read_restart restartfile [remap]
write_dump group-ID dumpstyle dumpfile arg1 arg2 ...
```

Note that the specified restartfile and dumpfile names may contain wild-card characters (“\*” or “%”) as explained on the *read\_restart* and *write\_dump* doc pages. The use of “%” means that a parallel restart file and/or parallel dump file can be read and/or written. Note that a filename such as file.\* may need to be enclosed in quotes or the “\*” character prefixed with a backslash (“\”) to avoid shell expansion of the “\*” character.

Note that following the restartfile argument, the optional word “remap” can be used. This has the effect as adding it to the *read\_restart* command, as explained on its doc page. This is useful if reading the restart file triggers an error that atoms have been lost. In that case, use of the remap flag should allow the dump file to still be produced.

The syntax following restartfile (or remap), namely

```
group-ID dumpstyle dumpfile arg1 arg2 ...
```

is identical to the arguments of the *write\_dump* command. See its page for details. This includes what per-atom fields are written to the dump file and optional dump\_modify settings, including ones that affect how parallel dump files are written, e.g. the *nfile* and *fileper* keywords. See the *dump\_modify* page for details.

---

#### **-screen file**

Specify a file for LAMMPS to write its screen information to. In one-partition mode, if the switch is not used, LAMMPS writes to the screen. If this switch is used, LAMMPS writes to the specified file instead and you will see no screen output. In multi-partition mode, if the switch is not used, high-level status information is written to the screen. Each partition also writes to a screen.N file where N is the partition ID. If the switch is specified in multi-partition mode, the high-level screen dump is named “file” and each partition also writes screen information to a file.N. For both one-partition and multi-partition mode, if the specified file is “none”, then no screen output is performed. Option -pscreen will override the name of the partition screen files file.N.

---

#### **-skiprun**

Insert the command *timer timeout 0 every 1* at the beginning of an input file or after a *clear* command. This has the effect that the entire LAMMPS input script is processed without executing actual *run* or *minimize* and similar commands (their main loops are skipped). This can be helpful and convenient to test input scripts of long running calculations for correctness to avoid having them crash after a long time due to a typo or syntax error in the middle or at the end.

---

#### **-suffix style args**

Use variants of various styles if they exist. The specified style can be *gpu*, *intel*, *kk*, *omp*, *opt*, or *hybrid*. These refer to optional packages that LAMMPS can be built with, as described in *Accelerate performance*. The “gpu” style

---

corresponds to the GPU package, the “intel” style to the INTEL package, the “kk” style to the KOKKOS package, the “opt” style to the OPT package, and the “omp” style to the OPENMP package. The hybrid style is the only style that accepts arguments. It allows for two packages to be specified. The first package specified is the default and will be used if it is available. If no style is available for the first package, the style for the second package will be used if available. For example, “-suffix hybrid intel omp” will use styles from the INTEL package if they are installed and available, but styles for the OPENMP package otherwise.

Along with the “-package” command-line switch, this is a convenient mechanism for invoking accelerator packages and their options without having to edit an input script.

As an example, all of the packages provide a *pair\_style lj/cut* variant, with style names `lj/cut/gpu`, `lj/cut/intel`, `lj/cut/kk`, `lj/cut/omp`, and `lj/cut/opt`. A variant style can be specified explicitly in your input script, e.g. `pair_style lj/cut/gpu`. If the `-suffix` switch is used the specified suffix (`gpu,intel,kk,omp,opt`) is automatically appended whenever your input script command creates a new *atom style*, *pair style*, *fix*, *compute*, or *run style*. If the variant version does not exist, the standard version is created.

For the GPU package, using this command-line switch also invokes the default GPU settings, as if the command “package gpu 1” were used at the top of your input script. These settings can be changed by using the “-package gpu” command-line switch or the *package gpu* command in your script.

For the INTEL package, using this command-line switch also invokes the default INTEL settings, as if the command “package intel 1” were used at the top of your input script. These settings can be changed by using the “-package intel” command-line switch or the *package intel* command in your script. If the OPENMP package is also installed, the hybrid style with “intel omp” arguments can be used to make the omp suffix a second choice, if a requested style is not available in the INTEL package. It will also invoke the default OPENMP settings, as if the command “package omp 0” were used at the top of your input script. These settings can be changed by using the “-package omp” command-line switch or the *package omp* command in your script.

For the KOKKOS package, using this command-line switch also invokes the default KOKKOS settings, as if the command “package kokkos” were used at the top of your input script. These settings can be changed by using the “-package kokkos” command-line switch or the *package kokkos* command in your script.

For the OMP package, using this command-line switch also invokes the default OMP settings, as if the command “package omp 0” were used at the top of your input script. These settings can be changed by using the “-package omp” command-line switch or the *package omp* command in your script.

The *suffix* command can also be used within an input script to set a suffix, or to turn off or back on any suffix setting made via the command line.

---

### **-var name value1 value2 ...**

Specify a variable that will be defined for substitution purposes when the input script is read. This switch can be used multiple times to define multiple variables. “Name” is the variable name which can be a single character (referenced as `$x` in the input script) or a full string (referenced as `${abc}`). An *index-style variable* will be created and populated with the subsequent values, e.g. a set of filenames. Using this command-line option is equivalent to putting the line “variable name index value1 value2 ...” at the beginning of the input script. Defining an index variable as a command-line argument overrides any setting for the same index variable in the input script, since index variables cannot be re-defined.

See the *variable* command for more info on defining index and other kinds of variables and the *Parsing rules* page for more info on using variables in input scripts.

---

**Note:** Currently, the command-line parser looks for arguments that start with “-” to indicate new switches. Thus you cannot specify multiple variable values if any of them start with a “-”, e.g. a negative numeric value. It is OK if the first value1 starts with a “-”, since it is automatically skipped.

---

### 4.3 Screen and logfile output

As LAMMPS reads an input script, it prints information to both the screen and a log file about significant actions it takes to setup a simulation. When the simulation is ready to begin, LAMMPS performs various initializations, and prints info about the run it is about to perform, including the amount of memory (in MBytes per processor) that the simulation requires. It also prints details of the initial thermodynamic state of the system. During the run itself, thermodynamic information is printed periodically, every few timesteps. When the run concludes, LAMMPS prints the final thermodynamic state and a total run time for the simulation. It also appends statistics about the CPU time and storage requirements for the simulation. An example set of statistics is shown here:

```
Loop time of 2.81192 on 4 procs for 300 steps with 2004 atoms

Performance: 18.436 ns/day  1.302 hours/ns  106.689 timesteps/s
97.0% CPU use with 4 MPI tasks x no OpenMP threads
```

MPI task timings breakdown:

Section	min time	avg time	max time	%varavg	%total
Pair	1.9808	2.0134	2.0318	1.4	71.60
Bond	0.0021894	0.0060319	0.010058	4.7	0.21
Kspace	0.3207	0.3366	0.36616	3.1	11.97
Neigh	0.28411	0.28464	0.28516	0.1	10.12
Comm	0.075732	0.077018	0.07883	0.4	2.74
Output	0.00030518	0.00042665	0.00078821	1.0	0.02
Modify	0.086606	0.086631	0.086668	0.0	3.08
Other		0.007178			0.26

```
Nlocal: 501 ave 508 max 490 min
Histogram: 1 0 0 0 0 0 1 1 0 1
Nghost: 6586.25 ave 6628 max 6548 min
Histogram: 1 0 1 0 0 0 1 0 0 1
Neighs: 177007 ave 180562 max 170212 min
Histogram: 1 0 0 0 0 0 0 1 1 1
```

```
Total # of neighbors = 708028
Ave neighs/atom = 353.307
Ave special neighs/atom = 2.34032
Neighbor list builds = 26
Dangerous builds = 0
```

The first section provides a global loop timing summary. The *loop time* is the total wall-clock time for the simulation to run. The *Performance* line is provided for convenience to help predict how long it will take to run a desired physical simulation. The *CPU use* line provides the CPU utilization per MPI task; it should be close to 100% times the number of OpenMP threads (or 1 of not using OpenMP). Lower numbers correspond to delays due to file I/O or insufficient thread utilization.

The *MPI task* section gives the breakdown of the CPU run time (in seconds) into major categories:

- *Pair* = non-bonded force computations
- *Bond* = bonded interactions: bonds, angles, dihedrals, impropers
- *Kspace* = long-range interactions: Ewald, PPPM, MSM

- *Neigh* = neighbor list construction
- *Comm* = inter-processor communication of atoms and their properties
- *Output* = output of thermodynamic info and dump files
- *Modify* = fixes and computes invoked by fixes
- *Other* = all the remaining time

For each category, there is a breakdown of the least, average and most amount of wall time any processor spent on this category of computation. The “%varavg” is the percentage by which the max or min varies from the average. This is an indication of load imbalance. A percentage close to 0 is perfect load balance. A large percentage is imbalance. The final “%total” column is the percentage of the total loop time is spent in this category.

When using the *timer full* setting, an additional column is added that also prints the CPU utilization in percent. In addition, when using *timer full* and the *package omp* command are active, a similar timing summary of time spent in threaded regions to monitor thread utilization and load balance is provided. A new *Thread timings* section is also added, which lists the time spent in reducing the per-thread data elements to the storage for non-threaded computation. These thread timings are measured for the first MPI rank only and thus, because the breakdown for MPI tasks can change from MPI rank to MPI rank, this breakdown can be very different for individual ranks. Here is an example output for this section:

```
Thread timings breakdown (MPI rank 0):
Total threaded time 0.6846 / 90.6%
Section | min time | avg time | max time | %varavg | %total
-----|-----|-----|-----|-----|-----
Pair    | 0.5127   | 0.5147   | 0.5167   | 0.3    | 75.18
Bond    | 0.0043139 | 0.0046779 | 0.0050418 | 0.5    | 0.68
Kspace  | 0.070572  | 0.074541  | 0.07851  | 1.5    | 10.89
Neigh   | 0.084778  | 0.086969  | 0.089161  | 0.7    | 12.70
Reduce  | 0.0036485 | 0.003737  | 0.0038254 | 0.1    | 0.55
```

The third section above lists the number of owned atoms (Nlocal), ghost atoms (Nghost), and pairwise neighbors stored per processor. The max and min values give the spread of these values across processors with a 10-bin histogram showing the distribution. The total number of histogram counts is equal to the number of processors.

The last section gives aggregate statistics (across all processors) for pairwise neighbors and special neighbors that LAMMPS keeps track of (see the *special\_bonds* command). The number of times neighbor lists were rebuilt is tallied, as is the number of potentially *dangerous* rebuilds. If atom movement triggered neighbor list rebuilding (see the *neigh\_modify* command), then dangerous reneighborings are those that were triggered on the first timestep atom movement was checked for. If this count is non-zero you may wish to reduce the delay factor to insure no force interactions are missed by atoms moving beyond the neighbor skin distance before a rebuild takes place.

If an energy minimization was performed via the *minimize* command, additional information is printed, e.g.

```
Minimization stats:
Stopping criterion = linesearch alpha is zero
Energy initial, next-to-last, final =
    -6372.3765206    -8328.46998942    -8328.46998942
Force two-norm initial, final = 1059.36 5.36874
Force max component initial, final = 58.6026 1.46872
Final line search alpha, max atom move = 2.7842e-10 4.0892e-10
Iterations, force evaluations = 701 1516
```

The first line prints the criterion that determined minimization was converged. The next line lists the initial and final energy, as well as the energy on the next-to-last iteration. The next 2 lines give a measure of the gradient of the energy (force on all atoms). The 2-norm is the “length” of this 3N-component force vector; the largest component (x, y, or z) of force (infinity-norm) is also given. Then information is provided about the line search and statistics on how many iterations and force-evaluations the minimizer required. Multiple force evaluations are typically done at each iteration to perform a 1d line minimization in the search direction. See the [minimize](#) page for more details.

---

If a *kspc* long-range Coulombics solver that performs FFTs was used during the run (PPPM, Ewald), then additional information is printed, e.g.

```
FFT time (% of Kspce) = 0.200313 (8.34477)
FFT Gflps 3d 1d-only = 2.31074 9.19989
```

The first line is the time spent doing 3d FFTs (several per timestep) and the fraction it represents of the total KSpace time (listed above). Each 3d FFT requires computation (3 sets of 1d FFTs) and communication (transposes). The total flops performed is  $5N\log_2(N)$ , where N is the number of points in the 3d grid. The FFTs are timed with and without the communication and a Gflop rate is computed. The 3d rate is with communication; the 1d rate is without (just the 1d FFTs). Thus you can estimate what fraction of your FFT time was spent in communication, roughly 75% in the example above.

## 4.4 Running LAMMPS on Windows

To run a serial (non-MPI) executable, follow these steps:

- Get a command prompt by going to Start->Run... , then typing “cmd”.
- Move to the directory where you have your input script, (e.g. by typing: cd “Documents”).
- At the command prompt, type “lmp -in in.file”, where in.file is the name of your LAMMPS input script.

Note that the serial executable includes support for multi-threading parallelization from the styles in the OPENMP packages. To run with 4 threads, you can type this:

```
lmp -in in.lj -pk omp 4 -sf omp
```

---

For the MPI executable, which allows you to run LAMMPS under Windows in parallel, follow these steps.

Download and install a compatible MPI library binary package:

- for 32-bit Windows: [mpich2-1.4.1p1-win-ia32.msi](#)
- for 64-bit Windows: [mpich2-1.4.1p1-win-x86-64.msi](#)

The LAMMPS Windows installer packages will automatically adjust your path for the default location of this MPI package. After the installation of the MPICH2 software, it needs to be integrated into the system. For this you need to start a Command Prompt in *Administrator Mode* (right click on the icon and select it). Change into the MPICH2 installation directory, then into the sub-directory **bin** and execute **smpd.exe -install**. Exit the command window.

- Get a new, regular command prompt by going to Start->Run... , then typing “cmd”.
- Move to the directory where you have your input file (e.g. by typing: cd “Documents”).

Then you can run the executable in serial like in the example above or in parallel using MPI with one of the following commands:



```
mpiexec -localonly 4 lmp -in in.file  
mpiexec -np 4 lmp -in in.file
```

where `in.file` is the name of your LAMMPS input script. For the latter case, you may be prompted to enter the password that you set during installation of the MPI library software.

In this mode, output may not immediately show up on the screen, so if your input script takes a long time to execute, you may need to be patient before the output shows up.

The parallel executable can also run on a single processor by typing something like this:

```
lmp -in in.lj
```

Note that the parallel executable also includes OpenMP multi-threading, which can be combined with MPI using something like:

```
mpiexec -localonly 2 lmp -in in.lj -pk omp 2 -sf omp
```



## COMMANDS

These pages describe how a LAMMPS input script is formatted and the commands in it are used to define a LAMMPS simulation.

### 5.1 LAMMPS input scripts

LAMMPS executes calculations by reading commands from an input script (text file), one line at a time. When the input script ends, LAMMPS exits. This is different from programs that read and process the entire input before starting a calculation.

Each command causes LAMMPS to take some immediate action without regard for any commands that may be processed later. Commands may set an internal variable, read in a file, or run a simulation. These actions can be grouped into three categories:

- a) commands that change a global setting (examples: `timestep`, `newton`, `echo`, `log`, `thermo`, `restart`),
- b) commands that add, modify, remove, or replace “styles” that are executed during a “run” (examples: `pair_style`, `fix`, `compute`, `dump`, `thermo_style`, `pair_modify`), and
- c) commands that execute a “run” or perform some other computation or operation (examples: `print`, `run`, `minimize`, `temper`, `write_dump`, `rerun`, `read_data`, `read_restart`)

Commands in category a) have default settings, which means you only need to use the command if you wish to change the defaults.

In many cases, the ordering of commands in an input script is not important, but can have consequences when the global state is changed between commands in the c) category. The following rules apply:

- (1) LAMMPS does not read your entire input script and then perform a simulation with all the settings. Rather, the input script is read one line at a time and each command takes effect when it is read. Thus this sequence of commands:

```
timestep 0.5
run      100
run      100
```

does something different than this sequence:

```
run      100
timestep 0.5
run      100
```

In the first case, the specified timestep (0.5 fs) is used for two simulations of 100 timesteps each. In the second case, the default timestep (1.0 fs) is used for the first 100 step simulation and a 0.5 fs timestep is used for the second one.

- (2) Some commands are only valid when they follow other commands. For example you cannot set the temperature of a group of atoms until atoms have been defined and a group command is used to define which atoms belong to the group.
- (3) Sometimes command B will use values that can be set by command A. This means command A must precede command B in the input script if it is to have the desired effect. For example, the `read_data` command initializes the system by setting up the simulation box and assigning atoms to processors. If default values are not desired, the `processors` and `boundary` commands need to be used before `read_data` to tell LAMMPS how to map processors to the simulation box.

Many input script errors are detected by LAMMPS and an ERROR or WARNING message is printed. The [Errors](#) page gives more information on what errors mean. The documentation for each command lists restrictions on how the command can be used.

You can use the `-skiprun` command line flag to have LAMMPS skip the execution of any “run”, “minimize”, or similar commands to check the entire input for correct syntax to avoid crashes on typos or syntax errors in long runs.

## 5.2 Parsing rules for input scripts

Each non-blank line in the input script is treated as a command. LAMMPS commands are case sensitive. Command names are lower-case, as are specified command arguments. Upper case letters may be used in file names or user-chosen ID strings.

Here are 6 rules for how each line in the input script is parsed by LAMMPS:

1. If the last printable character on the line is a “&” character, the command is assumed to continue on the next line. The next line is concatenated to the previous line by removing the “&” character and line break. This allows long commands to be continued across two or more lines. See the discussion of triple quotes in [6](#) for how to continue a command across multiple line without using “&” characters.
2. All characters from the first “#” character onward are treated as comment and discarded. The exception to this rule is described in [6](#). Note that a comment after a trailing “&” character will prevent the command from continuing on the next line. Also note that for multi-line commands a single leading “#” will comment out the entire command.

```
# this is a comment
timestep 1.0 # this is also a comment
```

3. The line is searched repeatedly for \$ characters, which indicate variables that are replaced with a text string. The exception to this rule is described in [6](#).

If the \$ is followed by text in curly brackets ‘{}’, then the variable name is the text inside the curly brackets. If no curly brackets follow the \$, then the variable name is the single character immediately following the \$. Thus  `${myTemp}` and  `$x` refer to variables named “myTemp” and “x”, while “\$xx” will be interpreted as a variable named “x” followed by an “x” character.

How the variable is converted to a text string depends on what style of variable it is; see the [variable](#) page for details. It can be a variable that stores multiple text strings, and return one of them. The returned text string can be multiple “words” (space separated) which will then be interpreted as multiple arguments in the input command. The variable can also store a numeric formula which will be evaluated and its numeric result returned as a string.

As a special case, if the \$ is followed by parenthesis “()”, then the text inside the parenthesis is treated as an “immediate” variable and evaluated as an *equal-style variable*. This is a way to use numeric formulas in an input script without having to assign them to variable names. For example, these 3 input script lines:

```
variable X equal (xlo+xhi)/2+sqrt(v_area)
region 1 block $X 2 INF INF EDGE EDGE
variable X delete
```

can be replaced by:

```
region 1 block $((xlo+xhi)/2+sqrt(v_area)) 2 INF INF EDGE EDGE
```

so that you do not have to define (or discard) a temporary variable, “X” in this case.

Additionally, the entire “immediate” variable expression may be followed by a colon, followed by a C-style format string, e.g. “:%f” or “:%.10g”. The format string must be appropriate for a double-precision floating-point value. The format string is used to output the result of the variable expression evaluation. If a format string is not specified, a high-precision “%.20g” is used as the default format.

This can be useful for formatting print output to a desired precision:

```
print "Final energy per atom: $(v_ke_per_atom+v_pe_per_atom:%10.3f) eV/atom"
```

Note that neither the curly-bracket or immediate form of variables can contain nested \$ characters for other variables to substitute for. Thus you may **NOT** do this:

```
variable a equal 2
variable b2 equal 4
print "B2 = ${b$a}"
```

Nor can you specify an expression like “\$(x-1.0)” for an immediate variable, but you could use \$(v\_x-1.0), since the latter is valid syntax for an *equal-style variable*.

See the *variable* command for more details of how strings are assigned to variables and evaluated, and how they can be used in input script commands.

4. The line is broken into “words” separated by white-space (tabs, spaces). Note that words can thus contain letters, digits, underscores, or punctuation characters.
5. The first word is the command name. All successive words in the line are arguments.
6. If you want text with spaces to be treated as a single argument, it can be enclosed in either single (') or double (") or triple (""") quotes. A long single argument enclosed in single or double quotes can span multiple lines if the “&” character is used, as described in 1 above. When the lines are concatenated together by LAMMPS (and the “&” characters and line breaks removed), the combined text will become a single line. If you want multiple lines of an argument to retain their line breaks, the text can be enclosed in triple quotes, in which case “&” characters are not needed and do not function as line continuation character. For example:

```
print "Volume = $v"
print 'Volume = $v'
if "${steps} > 1000" then quit
variable a string "red green blue &
                  purple orange cyan"
print """"
System volume = $v
System temperature = $t
""""
```

In each of these cases, the single, double, or triple quotes are removed and the enclosed text stored internally as a single argument.

See the *dump modify format, print, if, and python* commands for examples.

A “#” or “\$” character that is between quotes will not be treated as a comment indicator in 2 or substituted for as a variable in 3.

---

**Note:** If the argument is itself a command that requires a quoted argument (e.g. using a *print* command as part of an *if* or *run every* command), then single, double, or triple quotes can be nested in the usual manner. See the doc pages for those commands for examples. Only one of level of nesting is allowed, but that should be sufficient for most use cases.

---

---

### ASCII versus UTF-8

LAMMPS expects and processes 7-bit ASCII format text internally. Many modern environments use UTF-8 encoding, which is a superset of the 7-bit ASCII character table and thus mostly compatible. However, there are several non-ASCII characters that can look very similar to their ASCII equivalents or are invisible (so they look like a blank), but are encoded differently. Web browsers, PDF viewers, document editors are known to sometimes replace one with the other for a better looking output. However, that can lead to problems, for instance, when using cut-n-paste of input file examples from web pages, or when using a document editor (not a dedicated plain text editor) for writing LAMMPS inputs. LAMMPS will try to detect this and substitute the non-ASCII characters with their ASCII equivalents where known. There also is going to be a warning printed, if this occurs. It is recommended to avoid such characters altogether in LAMMPS input, data and potential files. The replacement tables are likely incomplete and dependent on users reporting problems processing correctly looking input containing UTF-8 encoded non-ASCII characters.

---

## 5.3 Input script structure

This page describes the structure of a typical LAMMPS input script. The examples directory in the LAMMPS distribution contains many sample input scripts; it is discussed on the *Examples* doc page.

A LAMMPS input script typically has 4 parts:

1. *Initialization*
2. *System definition*
3. *Simulation settings*
4. *Run a simulation*

The last 2 parts can be repeated as many times as desired. I.e. run a simulation, change some settings, run some more, etc. Each of the 4 parts is now described in more detail. Remember that almost all commands need only be used if a non-default value is desired.

### 5.3.1 Initialization

Set parameters that need to be defined before atoms are created or read-in from a file.

The relevant commands are *units*, *dimension*, *newton*, *processors*, *boundary*, *atom\_style*, *atom\_modify*.

If force-field parameters appear in the files that will be read, these commands tell LAMMPS what kinds of force fields are being used: *pair\_style*, *bond\_style*, *angle\_style*, *dihedral\_style*, *improper\_style*.

### 5.3.2 System definition

There are 3 ways to define the simulation cell and reserve space for force field info and fill it with atoms in LAMMPS. Read them in from (1) a data file or (2) a restart file via the *read\_data* or *read\_restart* commands, respectively. These files can also contain molecular topology information. Or (3) create a simulation cell and fill it with atoms on a lattice (with no molecular topology), using these commands: *lattice*, *region*, *create\_box*, *create\_atoms* or *read\_dump*.

The entire set of atoms can be duplicated to make a larger simulation using the *replicate* command.

### 5.3.3 Simulation settings

Once atoms and molecular topology are defined, a variety of settings can be specified: force field coefficients, simulation parameters, output options, and more.

Force field coefficients are set by these commands (they can also be set in the read-in files): *pair\_coeff*, *bond\_coeff*, *angle\_coeff*, *dihedral\_coeff*, *improper\_coeff*, *kspace\_style*, *dielectric*, *special\_bonds*.

Various simulation parameters are set by these commands: *neighbor*, *neigh\_modify*, *group*, *timestep*, *reset\_timestep*, *run\_style*, *min\_style*, *min\_modify*.

Fixes impose a variety of boundary conditions, time integration, and diagnostic options. The *fix* command comes in many flavors.

Various computations can be specified for execution during a simulation using the *compute*, *compute\_modify*, and *variable* commands.

Output options are set by the *thermo*, *dump*, and *restart* commands.

### 5.3.4 Run a simulation

A molecular dynamics simulation is run using the *run* command. Energy minimization (molecular statics) is performed using the *minimize* command. A parallel tempering (replica-exchange) simulation can be run using the *temper* command.

## 5.4 Commands by category

This page lists most of the LAMMPS commands, grouped by category. The *General commands* page lists all general commands alphabetically. Style options for entries like *fix*, *compute*, *pair* etc. have their own pages where they are listed alphabetically.

### 5.4.1 Initialization:

<i>newton</i>	<i>package</i>	<i>processors</i>	<i>suffix</i>	<i>units</i>
---------------	----------------	-------------------	---------------	--------------

### 5.4.2 Setup simulation box:

<i>boundary</i>	<i>box</i>	<i>change_box</i>	<i>create_box</i>
<i>dimension</i>	<i>lattice</i>	<i>region</i>	

### 5.4.3 Setup atoms:

<i>atom_modify</i>	<i>atom_style</i>	<i>balance</i>	<i>create_atoms</i>
<i>create_bonds</i>	<i>delete_atoms</i>	<i>delete_bonds</i>	<i>displace_atoms</i>
<i>group</i>	<i>mass</i>	<i>molecule</i>	<i>read_data</i>
<i>read_dump</i>	<i>read_restart</i>	<i>replicate</i>	<i>set</i>
<i>velocity</i>			

### 5.4.4 Force fields:

<i>angle_coeff</i>	<i>angle_style</i>	<i>bond_coeff</i>	<i>bond_style</i>
<i>bond_write</i>	<i>dielectric</i>	<i>dihedral_coeff</i>	<i>dihedral_style</i>
<i>improper_coeff</i>	<i>improper_style</i>	<i>kspace_modify</i>	<i>kspace_style</i>
<i>pair_coeff</i>	<i>pair_modify</i>	<i>pair_style</i>	<i>pair_write</i>
<i>special_bonds</i>			

### 5.4.5 Settings:

<i>comm_modify</i>	<i>comm_style</i>	<i>info</i>	<i>min_modify</i>
<i>min_style</i>	<i>neigh_modify</i>	<i>neighbor</i>	<i>partition</i>
<i>reset_timestep</i>	<i>run_style</i>	<i>timer</i>	<i>timestep</i>

### 5.4.6 Operations within timestepping (fixes) and diagnostics (computes):

<i>compute</i>	<i>compute_modify</i>	<i>fix</i>	<i>fix_modify</i>
<i>uncompute</i>	<i>unfix</i>		



### 5.4.7 Output:

<i>dump image</i>	<i>dump movie</i>	<i>dump</i>	<i>dump_modify</i>
<i>restart</i>	<i>thermo</i>	<i>thermo_modify</i>	<i>thermo_style</i>
<i>undump</i>	<i>write_coeff</i>	<i>write_data</i>	<i>write_dump</i>
<i>write_restart</i>			

### 5.4.8 Actions:

<i>minimize</i>	<i>neb</i>	<i>neb_spin</i>	<i>prd</i>	<i>rerun</i>	<i>run</i>
<i>tad</i>	<i>temper</i>				

### 5.4.9 Input script control:

<i>clear</i>	<i>echo</i>	<i>if</i>	<i>include</i>	<i>info</i>	<i>jump</i>	<i>label</i>
<i>log</i>	<i>next</i>	<i>print</i>	<i>python</i>	<i>quit</i>	<i>shell</i>	<i>variable</i>

<i>General commands</i>	<i>Fix styles</i>	<i>Compute styles</i>
<i>Pair styles</i>	<i>Bond styles</i>	<i>Angle styles</i>
<i>Dihedral styles</i>	<i>Improper styles</i>	<i>KSpace styles</i>
<i>Dump styles</i>		

## 5.5 General commands

An alphabetic list of general LAMMPS commands. Note that style commands with many variants, can be more easily accessed via the small table above.

<i>angle_coeff</i>	<i>angle_style</i>	<i>atom_modify</i>	<i>atom_style</i>	<i>balance</i>
<i>bond_coeff</i>	<i>bond_style</i>	<i>bond_write</i>	<i>boundary</i>	<i>box</i>
<i>change_box</i>	<i>clear</i>	<i>comm_modify</i>	<i>comm_style</i>	<i>compute</i>
<i>compute_modify</i>	<i>create_atoms</i>	<i>create_bonds</i>	<i>create_box</i>	<i>delete_atoms</i>
<i>delete_bonds</i>	<i>dielectric</i>	<i>dihedral_coeff</i>	<i>dihedral_style</i>	<i>dimension</i>
<i>displace_atoms</i>	<i>dump</i>	<i>dump_modify</i>	<i>echo</i>	<i>fix</i>
<i>fix_modify</i>	<i>group</i>	<i>if</i>	<i>improper_coeff</i>	<i>improper_style</i>
<i>include</i>	<i>info</i>	<i>jump</i>	<i>kspace_modify</i>	<i>kspace_style</i>
<i>label</i>	<i>labelmap</i>	<i>lattice</i>	<i>log</i>	<i>mass</i>
<i>minimize</i>	<i>min_modify</i>	<i>min_style</i>	<i>molecule</i>	<i>neigh_modify</i>
<i>neighbor</i>	<i>newton</i>	<i>next</i>	<i>package</i>	<i>pair_coeff</i>
<i>pair_modify</i>	<i>pair_style</i>	<i>pair_write</i>	<i>partition</i>	<i>print</i>
<i>processors</i>	<i>quit</i>	<i>read_data</i>	<i>read_dump</i>	<i>read_restart</i>
<i>region</i>	<i>replicate</i>	<i>rerun</i>	<i>reset_atom_ids</i>	<i>reset_mol_ids</i>
<i>reset_timestep</i>	<i>restart</i>	<i>run</i>	<i>run_style</i>	<i>set</i>
<i>shell</i>	<i>special_bonds</i>	<i>suffix</i>	<i>thermo</i>	<i>thermo_modify</i>
<i>thermo_style</i>	<i>timer</i>	<i>timestep</i>	<i>uncompute</i>	<i>undump</i>
<i>unfix</i>	<i>units</i>	<i>variable</i>	<i>velocity</i>	<i>write_coeff</i>
<i>write_data</i>	<i>write_dump</i>	<i>write_restart</i>		

Additional general LAMMPS commands provided by packages. A few commands have accelerated versions. This is indicated by an additional letter in parenthesis: k = KOKKOS.

<i>dynamical_matrix (k)</i>	<i>group2ndx</i>	<i>hyper</i>	<i>kim</i>	<i>mdi</i>
<i>ndx2group</i>	<i>neb</i>	<i>neb/spin</i>	<i>plugin</i>	<i>prd</i>
<i>python</i>	<i>tad</i>	<i>temper</i>	<i>temper/grem</i>	<i>temper/npt</i>
<i>third_order (k)</i>				

<i>General commands</i>	<i>Fix styles</i>	<i>Compute styles</i>
<i>Pair styles</i>	<i>Bond styles</i>	<i>Angle styles</i>
<i>Dihedral styles</i>	<i>Improper styles</i>	<i>KSpace styles</i>
<i>Dump styles</i>		

## 5.6 Fix commands

An alphabetic list of all LAMMPS *fix* commands. Some styles have accelerated versions. This is indicated by additional letters in parenthesis: g = GPU, i = INTEL, k = KOKKOS, o = OPENMP, t = OPT.

<i>accelerate/cos</i>	<i>acks2/reaxff (k)</i>	<i>adapt</i>	<i>adapt/fep</i>	<i>addforce</i>
<i>addtorque</i>	<i>amoeba/bitorsion</i>	<i>amoeba/pitorsion</i>	<i>append/atoms</i>	<i>atc</i>
<i>atom/swap</i>	<i>ave/atom</i>	<i>ave/chunk</i>	<i>ave/correlate</i>	<i>ave/correlate/long</i>
<i>ave/histo</i>	<i>ave/histo/weight</i>	<i>ave/time</i>	<i>aveforce</i>	<i>balance</i>
<i>brownian</i>	<i>brownian/asphere</i>	<i>brownian/sphere</i>	<i>bocs</i>	<i>bond/break</i>
<i>bond/create</i>	<i>bond/create/angle</i>	<i>bond/react</i>	<i>bond/swap</i>	<i>box/relax</i>
<i>charge/regulation</i>	<i>cmap</i>	<i>colvars</i>	<i>controller</i>	<i>damping/cundall</i>
<i>deform (k)</i>	<i>deposit</i>	<i>dpd/energy (k)</i>	<i>drag</i>	<i>drude</i>
<i>drude/transform/direct</i>	<i>drude/transform/inverse</i>	<i>dt/reset</i>	<i>edpd/source</i>	<i>efield</i>
<i>ehex</i>	<i>electrode/conp (i)</i>	<i>electrode/conq (i)</i>	<i>electrode/thermo (i)</i>	<i>electron/stopping</i>
<i>electron/stopping/fit</i>	<i>enforce2d (k)</i>	<i>eos/cv</i>	<i>eos/table</i>	<i>eos/table/rx (k)</i>
<i>evaporate</i>	<i>external</i>	<i>ffl</i>	<i>filter/corotate</i>	<i>flow/gauss</i>
<i>freeze (k)</i>	<i>gcmc</i>	<i>gld</i>	<i>gle</i>	<i>gravity (ko)</i>
<i>grem</i>	<i>halt</i>	<i>heat</i>	<i>hyper/global</i>	<i>hyper/local</i>
<i>imd</i>	<i>indent</i>	<i>ipi</i>	<i>langevin (k)</i>	<i>langevin/drude</i>
<i>langevin/eff</i>	<i>langevin/spin</i>	<i>latte</i>	<i>lb/fluid</i>	<i>lb/momentum</i>
<i>lb/viscous</i>	<i>lineforce</i>	<i>manifoldforce</i>	<i>mdi/qm</i>	<i>meso/move</i>
<i>mol/swap</i>	<i>momentum (k)</i>	<i>momentum/chunk</i>	<i>move</i>	<i>mscg</i>
<i>msst</i>	<i>mvv/dpd</i>	<i>mvv/edpd</i>	<i>mvv/tdpd</i>	<i>neb</i>
<i>neb/spin</i>	<i>nph (ko)</i>	<i>nph/asphere (o)</i>	<i>nph/body</i>	<i>nph/eff</i>
<i>nph/sphere (o)</i>	<i>nphug</i>	<i>npt (giko)</i>	<i>npt/asphere (o)</i>	<i>npt/body</i>
<i>npt/cauchy</i>	<i>npt/eff</i>	<i>npt/sphere (o)</i>	<i>npt/uef</i>	<i>numdiff</i>
<i>numdiff/virial</i>	<i>nve (giko)</i>	<i>nve/asphere (gi)</i>	<i>nve/asphere/noforce</i>	<i>nve/awpmd</i>
<i>nve/body</i>	<i>nve/dot</i>	<i>nve/dotc/langevin</i>	<i>nve/eff</i>	<i>nve/limit</i>
<i>nve/line</i>	<i>nve/manifold/rattle</i>	<i>nve/noforce</i>	<i>nve/sphere (ko)</i>	<i>nve/bpm/sphere</i>
<i>nve/spin</i>	<i>nve/tri</i>	<i>nvk</i>	<i>nvt (giko)</i>	<i>nvt/asphere (o)</i>
<i>nvt/body</i>	<i>nvt/eff</i>	<i>nvt/manifold/rattle</i>	<i>nvt/sllod (iko)</i>	<i>nvt/sllod/eff</i>
<i>nvt/sphere (o)</i>	<i>nvt/uef</i>	<i>oneway</i>	<i>orient/bcc</i>	<i>orient/fcc</i>
<i>orient/eco</i>	<i>pafi</i>	<i>pair</i>	<i>phonon</i>	<i>pimd</i>
<i>planeforce</i>	<i>plumed</i>	<i>poems</i>	<i>polarize/bem/gmres</i>	<i>polarize/bem/icc</i>

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Table 1 – continued from previous page

<i>polarize/functional</i>	<i>pour</i>	<i>precession/spin</i>	<i>press/berendsen</i>	<i>print</i>
<i>propel/self</i>	<i>property/atom (k)</i>	<i>python/invoke</i>	<i>python/move</i>	<i>qbmsst</i>
<i>qeq/comb (o)</i>	<i>qeq/dynamic</i>	<i>qeq/fire</i>	<i>qeq/point</i>	<i>qeq/reaxff (ko)</i>
<i>qeq/shielded</i>	<i>qeq/slater</i>	<i>qmmm</i>	<i>qtb</i>	<i>rattle</i>
<i>reaxff/bonds (k)</i>	<i>reaxff/species (k)</i>	<i>recenter</i>	<i>restrain</i>	<i>rhok</i>
<i>rigid (o)</i>	<i>rigid/meso</i>	<i>rigid/nph (o)</i>	<i>rigid/nph/small</i>	<i>rigid/npt (o)</i>
<i>rigid/npt/small</i>	<i>rigid/nve (o)</i>	<i>rigid/nve/small</i>	<i>rigid/nvt (o)</i>	<i>rigid/nvt/small</i>
<i>rigid/small (o)</i>	<i>rx (k)</i>	<i>saed/vtk</i>	<i>setforce (k)</i>	<i>setforce/spin</i>
<i>shake (k)</i>	<i>shardlow (k)</i>	<i>smd</i>	<i>smd/adjust_dt</i>	<i>smd/integrate_tlsph</i>
<i>smd/integrate_ulsph</i>	<i>smd/move_tri_surf</i>	<i>smd/setvel</i>	<i>smd/wall_surface</i>	<i>sph</i>
<i>sph/stationary</i>	<i>spring</i>	<i>spring/chunk</i>	<i>spring/rg</i>	<i>spring/self</i>
<i>srd</i>	<i>store/force</i>	<i>store/state</i>	<i>tdpd/source</i>	<i>temp/berendsen</i>
<i>temp/csld</i>	<i>temp/csvr</i>	<i>temp/rescale</i>	<i>temp/rescale/eff</i>	<i>tfmc</i>
<i>tnpt/drude</i>	<i>tnvt/drude</i>	<i>thermal/conductivity</i>	<i>ti/spring</i>	<i>tmd</i>
<i>ttm</i>	<i>ttm/grid</i>	<i>ttm/mod</i>	<i>tune/kSPACE</i>	<i>vector</i>
<i>viscosity</i>	<i>viscous</i>	<i>viscous/sphere</i>	<i>wall/body/polygon</i>	<i>wall/body/polyhedron</i>
<i>wall/colloid</i>	<i>wall/ees</i>	<i>wall/gran</i>	<i>wall/gran/region</i>	<i>wall/harmonic</i>
<i>wall/lj1043</i>	<i>wall/lj126</i>	<i>wall/lj93 (k)</i>	<i>wall/morse</i>	<i>wall/piston</i>
<i>wall/reflect (k)</i>	<i>wall/reflect/stochastic</i>	<i>wall/region</i>	<i>wall/region/ees</i>	<i>wall/srd</i>
<i>widom</i>				

<i>General commands</i>	<i>Fix styles</i>	<i>Compute styles</i>
<i>Pair styles</i>	<i>Bond styles</i>	<i>Angle styles</i>
<i>Dihedral styles</i>	<i>Improper styles</i>	<i>KSpace styles</i>
<i>Dump styles</i>		

## 5.7 Compute commands

An alphabetic list of all LAMMPS *compute* commands. Some styles have accelerated versions. This is indicated by additional letters in parenthesis: g = GPU, i = INTEL, k = KOKKOS, o = OPENMP, t = OPT.

<i>ackland/atom</i>	<i>adf</i>	<i>aggregate/atom</i>	<i>angle</i>	<i>angle/local</i>
<i>angmom/chunk</i>	<i>ave/sphere/atom (k)</i>	<i>basal/atom</i>	<i>body/local</i>	<i>bond</i>
<i>bond/local</i>	<i>born/matrix</i>	<i>centro/atom</i>	<i>centroid/stress/atom</i>	<i>chunk/atom</i>
<i>chunk/spread/atom</i>	<i>cluster/atom</i>	<i>cna/atom</i>	<i>cnp/atom</i>	<i>com</i>
<i>com/chunk</i>	<i>contact/atom</i>	<i>coord/atom (k)</i>	<i>damage/atom</i>	<i>dihedral</i>
<i>dihedral/local</i>	<i>dilatation/atom</i>	<i>dipole</i>	<i>dipole/chunk</i>	<i>displace/atom</i>
<i>dpd</i>	<i>dpd/atom</i>	<i>edpd/temp/atom</i>	<i>efield/atom</i>	<i>entropy/atom</i>
<i>erotate/asphere</i>	<i>erotate/rigid</i>	<i>erotate/sphere</i>	<i>erotate/sphere/atom</i>	<i>event/displace</i>
<i>fabric</i>	<i>fep</i>	<i>fep/ta</i>	<i>force/tally</i>	<i>fragment/atom</i>
<i>global/atom</i>	<i>group/group</i>	<i>gyration</i>	<i>gyration/chunk</i>	<i>gyration/shape</i>
<i>gyration/shape/chunk</i>	<i>heat/flux</i>	<i>heat/flux/tally</i>	<i>heat/flux/virial/tally</i>	<i>hexorder/atom</i>
<i>hma</i>	<i>improper</i>	<i>improper/local</i>	<i>inertia/chunk</i>	<i>ke</i>
<i>ke/atom</i>	<i>ke/atom/eff</i>	<i>ke/eff</i>	<i>ke/rigid</i>	<i>mesont</i>
<i>mliap</i>	<i>momentum</i>	<i>msd</i>	<i>msd/chunk</i>	<i>msd/nongauss</i>
<i>nbond/atom</i>	<i>omega/chunk</i>	<i>orientorder/atom (k)</i>	<i>pair</i>	<i>pair/local</i>
<i>pe</i>	<i>pe/atom</i>	<i>pe/mol/tally</i>	<i>pe/tally</i>	<i>plasticity/atom</i>
<i>pressure</i>	<i>pressure/uef</i>	<i>property/atom</i>	<i>property/chunk</i>	<i>property/local</i>

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Table 2 – continued from previous page

<i>ptm/atom</i>	<i>rdf</i>	<i>reduce</i>	<i>reduce/chunk</i>	<i>reduce/region</i>
<i>rigid/local</i>	<i>saed</i>	<i>slice</i>	<i>smd/contact/radius</i>	<i>smd/damage</i>
<i>smd/hourglass/error</i>	<i>smd/internal/energy</i>	<i>smd/plastic/strain</i>	<i>smd/plastic/strain/rate</i>	<i>smd/rho</i>
<i>smd/tlsph/defgrad</i>	<i>smd/tlsph/dt</i>	<i>smd/tlsph/num/neighs</i>	<i>smd/tlsph/shape</i>	<i>smd/tlsph/strain</i>
<i>smd/tlsph/strain/rate</i>	<i>smd/tlsph/stress</i>	<i>smd/triangle/vertices</i>	<i>smd/ulsph/effm</i>	<i>smd/ulsph/num/neighs</i>
<i>smd/ulsph/strain</i>	<i>smd/ulsph/strain/rate</i>	<i>smd/ulsph/stress</i>	<i>smd/vol</i>	<i>snap</i>
<i>sna/atom</i>	<i>sna/grid</i>	<i>sna/grid/local</i>	<i>snad/atom</i>	<i>snav/atom</i>
<i>sph/e/atom</i>	<i>sph/rho/atom</i>	<i>sph/t/atom</i>	<i>spin</i>	<i>stress/atom</i>
<i>stress/cartesian</i>	<i>stress/cylinder</i>	<i>stress/mop</i>	<i>stress/mop/profile</i>	<i>stress/spherical</i>
<i>stress/tally</i>	<i>tdpd/cc/atom</i>	<i>temp (k)</i>	<i>temp/asphere</i>	<i>temp/body</i>
<i>temp/chunk</i>	<i>temp/com</i>	<i>temp/cs</i>	<i>temp/deform (k)</i>	<i>temp/deform/eff</i>
<i>temp/drude</i>	<i>temp/eff</i>	<i>temp/partial</i>	<i>temp/profile</i>	<i>temp/ramp</i>
<i>temp/region</i>	<i>temp/region/eff</i>	<i>temp/rotate</i>	<i>temp/sphere</i>	<i>temp/uef</i>
<i>ti</i>	<i>torque/chunk</i>	<i>vacf</i>	<i>vcm/chunk</i>	<i>viscosity/cos</i>
<i>voronoi/atom</i>	<i>xrd</i>			

<i>General commands</i>	<i>Fix styles</i>	<i>Compute styles</i>
<i>Pair styles</i>	<i>Bond styles</i>	<i>Angle styles</i>
<i>Dihedral styles</i>	<i>Improper styles</i>	<i>KSpace styles</i>
<i>Dump styles</i>		

## 5.8 Pair\_style potentials

All LAMMPS *pair\_style* commands. Some styles have accelerated versions. This is indicated by additional letters in parenthesis: g = GPU, i = INTEL, k = KOKKOS, o = OPENMP, t = OPT.

<i>none</i>	<i>zero</i>	<i>hybrid (k)</i>	<i>hybrid/overlay (k)</i>
<i>hybrid/scaled</i>	<i>kim</i>	<i>list</i>	<i>tracker</i>
<i>adp (ko)</i>	<i>agni (o)</i>	<i>airebo (io)</i>	<i>airebo/morse (io)</i>
<i>amoeba</i>	<i>atm</i>	<i>awpmd/cut</i>	<i>beck (go)</i>
<i>body/nparticle</i>	<i>body/rounded/polygon</i>	<i>body/rounded/polyhedron</i>	<i>bop</i>
<i>born (go)</i>	<i>born/coul/dsf</i>	<i>born/coul/dsf/cs</i>	<i>born/coul/long (go)</i>
<i>born/coul/long/cs (g)</i>	<i>born/coul/msm (o)</i>	<i>born/coul/wolf (go)</i>	<i>born/coul/wolf/cs (g)</i>
<i>bpm/spring</i>	<i>brownian (o)</i>	<i>brownian/poly (o)</i>	<i>buck (giko)</i>
<i>buck/coul/cut (giko)</i>	<i>buck/coul/long (giko)</i>	<i>buck/coul/long/cs</i>	<i>buck/coul/msm (o)</i>
<i>buck/long/coul/long (o)</i>	<i>buck/mdf</i>	<i>buck6d/coul/gauss/dsf</i>	<i>buck6d/coul/gauss/long</i>
<i>colloid (go)</i>	<i>comb (o)</i>	<i>comb3</i>	<i>cosine/squared</i>
<i>coul/cut (gko)</i>	<i>coul/cut/dielectric</i>	<i>coul/cut/global (o)</i>	<i>coul/cut/soft (o)</i>
<i>coul/debye (gko)</i>	<i>coul/diel (o)</i>	<i>coul/dsf (gko)</i>	<i>coul/exclude</i>
<i>coul/long (gko)</i>	<i>coul/long/cs (g)</i>	<i>coul/long/dielectric</i>	<i>coul/long/soft (o)</i>
<i>coul/msm (o)</i>	<i>coul/slater/cut</i>	<i>coul/slater/long</i>	<i>coul/shield</i>
<i>coul/streitz</i>	<i>coul/tt</i>	<i>coul/wolf (ko)</i>	<i>coul/wolf/cs</i>
<i>dpd (giko)</i>	<i>dpd/fdt</i>	<i>dpd/ext (k)</i>	<i>dpd/ext/tstat (k)</i>
<i>dpd/fdt/energy (k)</i>	<i>dpd/tstat (gko)</i>	<i>dsmc</i>	<i>e3b</i>
<i>drip</i>	<i>eam (gikot)</i>	<i>eam/alloy (gikot)</i>	<i>eam/cd</i>
<i>eam/cd/old</i>	<i>eam/fs (gikot)</i>	<i>eam/he</i>	<i>edip (o)</i>
<i>edip/multi</i>	<i>edpd</i>	<i>eff/cut</i>	<i>eim (o)</i>

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Table 3 – continued from previous page

<i>exp6/rx (k)</i>	<i>extep</i>	<i>gauss (go)</i>	<i>gauss/cut (o)</i>
<i>gayberne (gio)</i>	<i>gran/hertz/history (o)</i>	<i>gran/hooke (o)</i>	<i>gran/hooke/history (ko)</i>
<i>granular</i>	<i>gw</i>	<i>gw/zbl</i>	<i>harmonic/cut (o)</i>
<i>hbond/dreiding/lj (o)</i>	<i>hbond/dreiding/morse (o)</i>	<i>hdnnp</i>	<i>hippo</i>
<i>ilp/graphene/hbn (t)</i>	<i>ilp/tmd (t)</i>	<i>kolmogorov/crespi/full</i>	<i>kolmogorov/crespi/z</i>
<i>lcbop</i>	<i>lebedeva/z</i>	<i>lennard/mdf</i>	<i>line/lj</i>
<i>lj/charmm/coul/charmm (giko)</i>	<i>lj/charmm/coul/charmm/implicit (ko)</i>	<i>lj/charmm/coul/long (gikot)</i>	<i>lj/charmm/coul/long/soft (o)</i>
<i>lj/charmm/coul/msm (o)</i>	<i>lj/charmmfsw/coul/charmmfsh</i>	<i>lj/charmmfsw/coul/long</i>	<i>lj/class2 (gko)</i>
<i>lj/class2/coul/cut (ko)</i>	<i>lj/class2/coul/cut/soft</i>	<i>lj/class2/coul/long (gko)</i>	<i>lj/class2/coul/long/cs</i>
<i>lj/class2/coul/long/soft</i>	<i>lj/class2/soft</i>	<i>lj/cubic (go)</i>	<i>lj/cut (gikot)</i>
<i>lj/cut/coul/cut (gko)</i>	<i>lj/cut/coul/cut/dielectric (o)</i>	<i>lj/cut/coul/cut/soft (o)</i>	<i>lj/cut/coul/debye (gko)</i>
<i>lj/cut/coul/debye/dielectric (o)</i>	<i>lj/cut/coul/dsf (gko)</i>	<i>lj/cut/coul/long (gikot)</i>	<i>lj/cut/coul/long/cs</i>
<i>lj/cut/coul/long/dielectric (o)</i>	<i>lj/cut/coul/long/soft (o)</i>	<i>lj/cut/coul/msm (go)</i>	<i>lj/cut/coul/msm/dielectric</i>
<i>lj/cut/coul/wolf (o)</i>	<i>lj/cut/dipole/cut (go)</i>	<i>lj/cut/dipole/long (g)</i>	<i>lj/cut/dipole/sf (go)</i>
<i>lj/cut/soft (o)</i>	<i>lj/cut/thole/long (o)</i>	<i>lj/cut/tip4p/cut (o)</i>	<i>lj/cut/tip4p/long (got)</i>
<i>lj/cut/tip4p/long/soft (o)</i>	<i>lj/expand (gko)</i>	<i>lj/expand/coul/long (g)</i>	<i>lj/gromacs (gko)</i>
<i>lj/gromacs/coul/gromacs (ko)</i>	<i>lj/long/coul/long (iot)</i>	<i>lj/long/coul/long/dielectric</i>	<i>lj/long/dipole/long</i>
<i>lj/long/tip4p/long (o)</i>	<i>lj/mdf</i>	<i>lj/relres (o)</i>	<i>lj/spica (gko)</i>
<i>lj/spica/coul/long (go)</i>	<i>lj/spica/coul/msm (o)</i>	<i>lj/sf/dipole/sf (go)</i>	<i>lj/smooth (go)</i>
<i>lj/smooth/linear (o)</i>	<i>lj/switch3/coulgauss/long</i>	<i>lj96/cut (go)</i>	<i>local/density</i>
<i>lubricate (o)</i>	<i>lubricate/poly (o)</i>	<i>lubricateU</i>	<i>lubricateU/poly</i>
<i>mdpd</i>	<i>mdpd/rhosum</i>	<i>meam (k)</i>	<i>meam/spline (o)</i>
<i>meam/sw/spline</i>	<i>mesocnt</i>	<i>mesocnt/viscous</i>	<i>mesont/tpm</i>
<i>mgpt</i>	<i>mie/cut (g)</i>	<i>mliap</i>	<i>mm3/switch3/coulgauss/long</i>
<i>momb</i>	<i>morse (gkot)</i>	<i>morse/smooth/linear (o)</i>	<i>morse/soft</i>
<i>multi/lucy</i>	<i>multi/lucy/rx (k)</i>	<i>nb3b/harmonic</i>	<i>nm/cut (o)</i>
<i>nm/cut/coul/cut (o)</i>	<i>nm/cut/coul/long (o)</i>	<i>nm/cut/split</i>	<i>oxdna/coaxstk</i>
<i>oxdna/excv</i>	<i>oxdna/hbond</i>	<i>oxdna/stk</i>	<i>oxdna/xstk</i>
<i>oxdna2/coaxstk</i>	<i>oxdna2/dh</i>	<i>oxdna2/excv</i>	<i>oxdna2/hbond</i>
<i>oxdna2/stk</i>	<i>oxdna2/xstk</i>	<i>oxrna2/excv</i>	<i>oxrna2/hbond</i>
<i>oxrna2/dh</i>	<i>oxrna2/stk</i>	<i>oxrna2/xstk</i>	<i>oxrna2/coaxstk</i>
<i>pace (k)</i>	<i>peri/eps</i>	<i>peri/lps (o)</i>	<i>peri/pmb (o)</i>
<i>peri/ves</i>	<i>polymorphic</i>	<i>python</i>	<i>quip</i>
<i>rann</i>	<i>reaxff (ko)</i>	<i>rebo (io)</i>	<i>resquared (go)</i>
<i>saip/metal (t)</i>	<i>sdpd/taitwater/isothermal</i>	<i>smatb</i>	<i>smatb/single</i>
<i>smd/hertz</i>	<i>smd/tlsph</i>	<i>smd/tri_surface</i>	<i>smd/ulsph</i>
<i>smtbq</i>	<i>snap (k)</i>	<i>soft (go)</i>	<i>sph/heatconduction</i>
<i>sph/idealgas</i>	<i>sph/lj</i>	<i>sph/rhosum</i>	<i>sph/taitwater</i>
<i>sph/taitwater/morris</i>	<i>spin/dipole/cut</i>	<i>spin/dipole/long</i>	<i>spin/dmi</i>
<i>spin/exchange</i>	<i>spin/exchange/biquadratic</i>	<i>spin/magelec</i>	<i>spin/neel</i>
<i>srp</i>	<i>srp/react</i>	<i>sw (giko)</i>	<i>sw/angle/table</i>
<i>sw/mod (o)</i>	<i>table (gko)</i>	<i>table/rx (k)</i>	<i>tdpd</i>
<i>tersoff (giko)</i>	<i>tersoff/mod (gko)</i>	<i>tersoff/mod/c (o)</i>	<i>tersoff/table (o)</i>
<i>tersoff/zbl (gko)</i>	<i>thole</i>	<i>threebody/table</i>	<i>tip4p/cut (o)</i>
<i>tip4p/long (o)</i>	<i>tip4p/long/soft (o)</i>	<i>tri/lj</i>	<i>ufm (got)</i>
<i>vashishta (gko)</i>	<i>vashishta/table (o)</i>	<i>wf/cut</i>	<i>yukawa (gko)</i>
<i>yukawa/colloid (go)</i>	<i>zbl (gko)</i>		

<i>General commands</i>	<i>Fix styles</i>	<i>Compute styles</i>
<i>Pair styles</i>	<i>Bond styles</i>	<i>Angle styles</i>
<i>Dihedral styles</i>	<i>Improper styles</i>	<i>KSpace styles</i>
<i>Dump styles</i>		

## 5.9 Bond\_style potentials

All LAMMPS *bond\_style* commands. Some styles have accelerated versions. This is indicated by additional letters in parenthesis: g = GPU, i = INTEL, k = KOKKOS, o = OPENMP, t = OPT.

<i>none</i>	<i>zero</i>	<i>hybrid</i>	
<i>bpm/rotational</i>	<i>bpm/spring</i>	<i>class2 (ko)</i>	<i>fene (iko)</i>
<i>fene/expand (o)</i>	<i>fene/nm</i>	<i>gaussian</i>	<i>gromos (o)</i>
<i>harmonic (iko)</i>	<i>harmonic/shift (o)</i>	<i>harmonic/shift/cut (o)</i>	<i>mesocnt</i>
<i>mm3</i>	<i>morse (o)</i>	<i>nonlinear (o)</i>	<i>oxdna/fene</i>
<i>oxdna2/fene</i>	<i>oxrna2/fene</i>	<i>quartic (o)</i>	<i>special</i>
<i>table (o)</i>			

## 5.10 Angle\_style potentials

All LAMMPS *angle\_style* commands. Some styles have accelerated versions. This is indicated by additional letters in parenthesis: g = GPU, i = INTEL, k = KOKKOS, o = OPENMP, t = OPT.

<i>none</i>	<i>zero</i>	<i>hybrid</i>	
<i>amoeba</i>	<i>charmm (iko)</i>	<i>class2 (ko)</i>	<i>class2/p6</i>
<i>cosine (ko)</i>	<i>cosine/buck6d</i>	<i>cosine/delta (o)</i>	<i>cosine/periodic (o)</i>
<i>cosine/shift (o)</i>	<i>cosine/shift/exp (o)</i>	<i>cosine/squared (o)</i>	<i>cross</i>
<i>dipole (o)</i>	<i>fourier (o)</i>	<i>fourier/simple (o)</i>	<i>gaussian</i>
<i>harmonic (iko)</i>	<i>mesocnt</i>	<i>mm3</i>	<i>quartic (o)</i>
<i>spica (o)</i>	<i>table (o)</i>		

## 5.11 Dihedral\_style potentials

All LAMMPS *dihedral\_style* commands. Some styles have accelerated versions. This is indicated by additional letters in parenthesis: g = GPU, i = INTEL, k = KOKKOS, o = OPENMP, t = OPT.

<i>none</i>	<i>zero</i>	<i>hybrid</i>	
<i>charmm (iko)</i>	<i>charmmfsw</i>	<i>class2 (ko)</i>	<i>cosine/shift/exp (o)</i>
<i>fourier (io)</i>	<i>harmonic (iko)</i>	<i>helix (o)</i>	<i>multi/harmonic (o)</i>
<i>nharmonic (o)</i>	<i>opls (iko)</i>	<i>quadratic (o)</i>	<i>spherical</i>
<i>table (o)</i>	<i>table/cut</i>		

## 5.12 Improper\_style potentials

All LAMMPS *improper\_style* commands. Some styles have accelerated versions. This is indicated by additional letters in parenthesis: g = GPU, i = INTEL, k = KOKKOS, o = OPENMP, t = OPT.

<i>none</i>	<i>zero</i>	<i>hybrid</i>	
<i>amoeba</i>	<i>class2 (ko)</i>	<i>cossq (o)</i>	<i>cvff (io)</i>
<i>distance</i>	<i>distharm</i>	<i>fourier (o)</i>	<i>harmonic (iko)</i>
<i>inversion/harmonic</i>	<i>ring (o)</i>	<i>sqdistharm</i>	<i>umbrella (o)</i>

<i>General commands</i>	<i>Fix styles</i>	<i>Compute styles</i>
<i>Pair styles</i>	<i>Bond styles</i>	<i>Angle styles</i>
<i>Dihedral styles</i>	<i>Improper styles</i>	<i>KSpace styles</i>
<i>Dump styles</i>		

## 5.13 KSpace solvers

All LAMMPS *kpace\_style* solvers. Some styles have accelerated versions. This is indicated by additional letters in parenthesis: g = GPU, i = INTEL, k = KOKKOS, o = OPENMP, t = OPT.

<i>ewald (o)</i>	<i>ewald/disp</i>	<i>ewald/disp/dipole</i>	<i>ewald/dipole</i>
<i>ewald/dipole/spin</i>	<i>ewald/electrode</i>	<i>msm (o)</i>	<i>msm/cg (o)</i>
<i>msm/dielectric</i>	<i>pppm (giko)</i>	<i>pppm/cg (o)</i>	<i>pppm/dipole</i>
<i>pppm/dipole/spin</i>	<i>pppm/dielectric</i>	<i>pppm/disp (io)</i>	<i>pppm/disp/tip4p (o)</i>
<i>pppm/disp/dielectric</i>	<i>pppm/stagger</i>	<i>pppm/tip4p (o)</i>	<i>pppm/dielectric</i>
<i>pppm/electrode (i)</i>	<i>scafacos</i>		

<i>General commands</i>	<i>Fix styles</i>	<i>Compute styles</i>
<i>Pair styles</i>	<i>Bond styles</i>	<i>Angle styles</i>
<i>Dihedral styles</i>	<i>Improper styles</i>	<i>KSpace styles</i>
<i>Dump styles</i>		

## 5.14 Dump commands

An alphabetic list of all LAMMPS *dump* commands.

<i>atom</i>	<i>atom/adios</i>	<i>atom/gz</i>	<i>atom/mpiio</i>	<i>atom/zstd</i>
<i>cfg</i>	<i>cfg/gz</i>	<i>cfg/mpiio</i>	<i>cfg/uef</i>	<i>cfg/zstd</i>
<i>custom</i>	<i>custom/adios</i>	<i>custom/gz</i>	<i>custom/mpiio</i>	<i>custom/zstd</i>
<i>dcd</i>	<i>deprecated</i>	<i>h5md</i>	<i>image</i>	<i>local</i>
<i>local/gz</i>	<i>local/zstd</i>	<i>molfile</i>	<i>movie</i>	<i>netcdf</i>
<i>netcdf/mpiio</i>	<i>vtk</i>	<i>xtc</i>	<i>xyz</i>	<i>xyz/gz</i>
<i>xyz/mpiio</i>	<i>xyz/zstd</i>	<i>yaml</i>		

## 5.15 Removed commands and packages

This page lists LAMMPS commands and packages that have been removed from the distribution and provides suggestions for alternatives or replacements. LAMMPS has special dummy styles implemented, that will stop LAMMPS and print a suitable error message in most cases, when a style/command is used that has been removed.

### 5.15.1 Fix ave/spatial and fix ave/spatial/sphere

The fixes `ave/spatial` and `ave/spatial/sphere` have been removed from LAMMPS since they were superseded by the more general and extensible “chunk infrastructure”. Here the system is partitioned in one of many possible ways through the `compute chunk/atom` command and then averaging is done using `fix ave/chunk`. Please refer to the `chunk HOWTO` section for an overview.

### 5.15.2 Reset\_ids command

The `reset_ids` command has been renamed to `reset_atom_ids`.

### 5.15.3 MEAM package

The MEAM package in Fortran has been replaced by a C++ implementation. The code in the `MEAM package` is a translation of the Fortran code of MEAM into C++, which removes several restrictions (e.g. there can be multiple instances in hybrid pair styles) and allows for some optimizations leading to better performance. The pair style `meam` has the exact same syntax.

### 5.15.4 REAX package

The REAX package has been removed since it was superseded by the `REAXFF package`. The REAXFF package has been tested to yield equivalent results to the REAX package, offers better performance, supports OpenMP multi-threading via OPENMP, and GPU and threading parallelization through KOKKOS. The new pair styles are not syntax compatible with the removed `reax` pair style, so input files will have to be adapted.

### 5.15.5 USER-CUDA package

The USER-CUDA package had been removed, since it had been unmaintained for a long time and had known bugs and problems. Significant parts of the design were transferred to the `KOKKOS package`, which has similar performance characteristics on NVIDIA GPUs. Both, the KOKKOS and the `GPU package` are maintained and allow running LAMMPS with GPU acceleration.

### 5.15.6 restart2data tool

The functionality of the `restart2data` tool has been folded into the LAMMPS executable directly instead of having a separate tool. A combination of the commands `read_restart` and `write_data` can be used to the same effect. For added convenience this conversion can also be triggered by *command line flags*



## OPTIONAL PACKAGES

This section gives an overview of the optional packages that extend LAMMPS functionality. Packages are groups of files that enable a specific set of features. For example, force fields for molecular systems or rigid-body constraints are in packages. You can see the list of all packages and “make” commands to manage them by typing “make package” from within the src directory of the LAMMPS distribution. The *Build package* page gives general info on how to install and un-install packages as part of the LAMMPS build process.

### 6.1 Available Packages

This is the list of packages included in LAMMPS. The link for each package name gives more details.

Packages are supported by either the LAMMPS developers or the contributing authors and written in a syntax and style consistent with the rest of LAMMPS.

The “Example” column is a sub-directory in the examples directory of the distribution which has an input script that uses the package. E.g. “peptide” refers to the examples/peptide directory; PACKAGES/atc refers to the examples/PACKAGES/atc directory. The “Library” column indicates whether an extra library is needed to build and use the package:

- no = no library
- sys = system library: you likely have it on your machine
- int = internal library: provided with LAMMPS, but you may need to build it
- ext = external library: you will need to download and install it on your machine

Package	Description	Doc page	Example
<i>ADIOS</i>	dump output via ADIOS	<i>dump adios</i>	PACKAGES,
<i>AMOEB</i> A	AMOEB A and HIPPO force fields	<i>AMOEB A and HIPPO howto</i>	amoeba
<i>ASPH</i> ERE	aspherical particle models	<i>Howto spherical</i>	ellipse
<i>ATC</i>	Atom-to-Continuum coupling	<i>fix atc</i>	PACKAGES,
<i>AWP</i> MD	wave packet MD	<i>pair_style awpmd/cut</i>	PACKAGES,
<i>BOCS</i>	BOCS bottom up coarse graining	<i>fix bocs</i>	PACKAGES,
<i>BOD</i> Y	body-style particles	<i>Howto body</i>	body
<i>BPM</i>	bonded particle models	<i>Howto bpm</i>	bpm
<i>BROW</i> NIAN	Brownian dynamics, self-propelled particles	<i>fix brownian, fix propel/self</i>	PACKAGES,
<i>CG-D</i> NA	coarse-grained DNA force fields	src/CG-DNA/README	PACKAGES,
<i>CG-S</i> PICA	SPICA (SDK) coarse-graining model	<i>pair_style lj/spica</i>	PACKAGES,
<i>CL</i> ASS2	class 2 force fields	<i>pair_style lj/class2</i>	n/a
<i>COL</i> LOID	colloidal particles	<i>atom_style colloid</i>	colloid
<i>COL</i> VARS	collective variables library	<i>fix colvars</i>	PACKAGES,

Table 1 – continued from previous page

Package	Description	Doc page	Example
<i>COMPRESS</i>	I/O compression	<i>dump */gz</i>	n/a
<i>CORESHELL</i>	adiabatic core/shell model	<i>Howto coreshell</i>	coreshell
<i>DIELECTRIC</i>	dielectric boundary solvers and force styles	<i>compute efield/atom</i>	PACKAGES,
<i>DIFFRACTION</i>	virtual x-ray and electron diffraction	<i>compute xrd</i>	PACKAGES,
<i>DIPOLE</i>	point dipole particles	<i>pair_style lj/.../dipole</i>	dipole
<i>DPD-BASIC</i>	basic DPD models	<i>pair_styles dpd dpd/tstat dpd/ext dpd/ext/tstat</i>	PACKAGES,
<i>DPD-MESO</i>	mesoscale DPD models	<i>pair_style edpd</i>	PACKAGES,
<i>DPD-REACT</i>	reactive dissipative particle dynamics	src/DPD-REACT/README	PACKAGES,
<i>DPD-SMOOTH</i>	smoothed dissipative particle dynamics	src/DPD-SMOOTH/README	PACKAGES,
<i>DRUDE</i>	Drude oscillators	<i>Howto drude</i>	PACKAGES,
<i>EFF</i>	electron force field	<i>pair_style eff/cut</i>	PACKAGES,
<i>ELECTRODE</i>	electrode charges to match potential	<i>fix electrode/comp</i>	PACKAGES,
<i>EXTRA-COMPUTE</i>	additional compute styles	<i>compute</i>	n/a
<i>EXTRA-DUMP</i>	additional dump styles	<i>dump</i>	n/a
<i>EXTRA-FIX</i>	additional fix styles	<i>fix</i>	n/a
<i>EXTRA-MOLECULE</i>	additional molecular styles	<i>molecular styles</i>	n/a
<i>EXTRA-PAIR</i>	additional pair styles	<i>pair_style</i>	n/a
<i>FEP</i>	free energy perturbation	<i>compute fep</i>	PACKAGES,
<i>GPU</i>	GPU-enabled styles	<i>Section gpu</i>	Benchmarks
<i>GRANULAR</i>	granular systems	<i>Howto granular</i>	pour
<i>H5MD</i>	dump output via HDF5	<i>dump h5md</i>	n/a
<i>INTEL</i>	optimized Intel CPU and KNL styles	<i>Speed intel</i>	Benchmarks
<i>INTERLAYER</i>	Inter-layer pair potentials	<i>several pair styles</i>	PACKAGES,
<i>KIM</i>	OpenKIM wrapper	<i>pair_style kim</i>	kim
<i>KOKKOS</i>	Kokkos-enabled styles	<i>Speed kokkos</i>	Benchmarks
<i>KSPACE</i>	long-range Coulombic solvers	<i>kpace_style</i>	peptide
<i>LATBOLTZ</i>	Lattice Boltzmann fluid	<i>fix lb/fluid</i>	PACKAGES,
<i>LATTE</i>	quantum DFTB forces via LATTE	<i>fix latte</i>	latte
<i>MACHDYN</i>	smoothed Mach dynamics	SMD User Guide	PACKAGES,
<i>MANIFOLD</i>	motion on 2d surfaces	<i>fix manifoldforce</i>	PACKAGES,
<i>MANYBODY</i>	many-body potentials	<i>pair_style tersoff</i>	shear
<i>MC</i>	Monte Carlo options	<i>fix gcmc</i>	n/a
<i>MDI</i>	client-server code coupling	<i>MDI Howto</i>	PACKAGES,
<i>MEAM</i>	modified EAM potential (C++)	<i>pair_style meam</i>	meam
<i>MESONT</i>	mesoscopic tubular potential model	<i>pair styles mesont/tpm, mesocnt</i>	PACKAGES,
<i>MGPT</i>	fast MGPT multi-ion potentials	<i>pair_style mgpt</i>	PACKAGES,
<i>MISC</i>	miscellaneous single-file commands	n/a	no
<i>ML-HDNNP</i>	High-dimensional neural network potentials	<i>pair_style hdnnp</i>	PACKAGES,
<i>ML-IAP</i>	multiple machine learning potentials	<i>pair_style mliap</i>	mliap
<i>ML-PACE</i>	Atomic Cluster Expansion potential	<i>pair pace</i>	PACKAGES,
<i>ML-QUIP</i>	QUIP/libatoms interface	<i>pair_style quip</i>	PACKAGES,
<i>ML-RANN</i>	Pair style for RANN potentials	<i>pair rann</i>	PACKAGES,
<i>ML-SNAP</i>	quantum-fitted potential	<i>pair_style snap</i>	snap
<i>MOFFF</i>	styles for MOF-FF force field	<i>pair_style buck6d/coul/gauss</i>	PACKAGES,
<i>MOLECULE</i>	molecular system force fields	<i>Howto bioFF</i>	peptide
<i>MOLFILE</i>	VMD molfile plug-ins	<i>dump molfile</i>	n/a
<i>MPIO</i>	MPI parallel I/O dump and restart	<i>dump</i>	n/a
<i>MSCG</i>	multi-scale coarse-graining wrapper	<i>fix mscg</i>	mscg
<i>NETCDF</i>	dump output via NetCDF	<i>dump netcdf</i>	n/a

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Table 1 – continued from previous page

Package	Description	Doc page	Example
<i>OPENMP</i>	OpenMP-enabled styles	<i>Speed omp</i>	Benchmarks
<i>OPT</i>	optimized pair styles	<i>Speed opt</i>	Benchmarks
<i>ORIENT</i>	fixes for orientation depended forces	<i>fix orient/*</i>	PACKAGES.
<i>PERI</i>	Peridynamics models	<i>pair_style peri</i>	peri
<i>PHONON</i>	phonon dynamical matrix	<i>fix phonon</i>	PACKAGES.
<i>PLUGIN</i>	Plugin loader command	<i>plugin</i>	plugins
<i>PLUMED</i>	<i>PLUMED</i> free energy library	<i>fix plumed</i>	PACKAGES.
<i>POEMS</i>	coupled rigid body motion	<i>fix poems</i>	rigid
<i>PTM</i>	Polyhedral Template Matching	<i>compute ptm/atom</i>	n/a
<i>PYTHON</i>	embed Python code in an input script	<i>python</i>	python
<i>QEQ</i>	QEq charge equilibration	<i>fix qeq</i>	qeq
<i>QMMM</i>	QM/MM coupling	<i>fix qmmm</i>	PACKAGES.
<i>QTB</i>	quantum nuclear effects	<i>fix qtb fix qbmsst</i>	qtb
<i>REACTION</i>	chemical reactions in classical MD	<i>fix bond/react</i>	PACKAGES.
<i>REAXFF</i>	ReaxFF potential (C/C++)	<i>pair_style reaxff</i>	reax
<i>REPLICA</i>	multi-replica methods	<i>Howto replica</i>	tad
<i>RIGID</i>	rigid bodies and constraints	<i>fix rigid</i>	rigid
<i>SCAFACOS</i>	wrapper for ScaFaCoS Kspace solver	<i>kpace_style scafacos</i>	PACKAGES.
<i>SHOCK</i>	shock loading methods	<i>fix msst</i>	n/a
<i>SMTBQ</i>	second moment tight binding potentials	<i>pair_style smtbq pair_style smatb</i>	PACKAGES.
<i>SPH</i>	smoothed particle hydrodynamics	SPH User Guide	PACKAGES.
<i>SPIN</i>	magnetic atomic spin dynamics	<i>Howto spins</i>	SPIN
<i>SRD</i>	stochastic rotation dynamics	<i>fix srd</i>	srd
<i>TALLY</i>	pairwise tally computes	<i>compute XXX/tally</i>	PACKAGES.
<i>UEF</i>	extensional flow	<i>fix nvt/uef</i>	PACKAGES.
<i>VORONOI</i>	Voronoi tessellation	<i>compute voronoi/atom</i>	n/a
<i>VTK</i>	dump output via VTK	<i>compute vtk</i>	n/a
<i>YAFF</i>	additional styles implemented in YAFF	<i>angle_style cross</i>	PACKAGES.

## 6.2 Package details

Here is a brief description of all packages in LAMMPS. It lists authors (if applicable) and summarizes the package contents. It has specific instructions on how to install the package, including, if necessary, info on how to download or build any extra library it requires. It also gives links to documentation, example scripts, and pictures/movies (if available) that illustrate use of the package.

The majority of packages can be included in a LAMMPS build with a single setting (`-D PKG_<NAME>=on` for CMake) or command (`make yes-<name>` for make). See the [Build package](#) page for more info. A few packages may require additional steps; this is indicated in the descriptions below. The [Build extras](#) page gives those details.

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**Note:** To see the complete list of commands a package adds to LAMMPS, you can examine the files in its src directory, e.g. “`ls src/GRANULAR`”. Files with names that start with `fix`, `compute`, `atom`, `pair`, `bond`, `angle`, etc correspond to commands with the same style name as contained in the file name.

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<i>ADIOS</i>	<i>AMOEBA</i>	<i>ASPHERE</i>	<i>ATC</i>	<i>AWPMD</i>	<i>BOCS</i>
<i>BODY</i>	<i>BPM</i>	<i>BROWNIAN</i>	<i>CG-DNA</i>	<i>CG-SPICA</i>	<i>CLASS2</i>
<i>COLLOID</i>	<i>COLVARS</i>	<i>COMPRESS</i>	<i>CORESHELL</i>	<i>DIELECTRIC</i>	<i>DIFFRACTION</i>
<i>DIPOLE</i>	<i>DPD-BASIC</i>	<i>DPD-MESO</i>	<i>DPD-REACT</i>	<i>DPD-SMOOTH</i>	<i>DRUDE</i>
<i>EFF</i>	<i>ELECTRODE</i>	<i>EXTRA-COMPUTE</i>	<i>EXTRA-DUMP</i>	<i>EXTRA-FIX</i>	<i>EXTRA-MOLECULE</i>
<i>EXTRA-PAIR</i>	<i>FEP</i>	<i>GPU</i>	<i>GRANULAR</i>	<i>H5MD</i>	<i>INTEL</i>
<i>INTER-LAYER</i>	<i>KIM</i>	<i>KOKKOS</i>	<i>KSPACE</i>	<i>LATBOLTZ</i>	<i>LATTE</i>
<i>MACHDYN</i>	<i>MANIFOLD</i>	<i>MANYBODY</i>	<i>MC</i>	<i>MDI</i>	<i>MEAM</i>
<i>MESONT</i>	<i>MGPT</i>	<i>MISC</i>	<i>ML-HDNNP</i>	<i>ML-IAP</i>	<i>ML-PACE</i>
<i>ML-QUIP</i>	<i>ML-RANN</i>	<i>ML-SNAP</i>	<i>MOFFF</i>	<i>MOLECULE</i>	<i>MOLFILE</i>
<i>MPIIO</i>	<i>MSCG</i>	<i>NETCDF</i>	<i>OPENMP</i>	<i>OPT</i>	<i>ORIENT</i>
<i>PERI</i>	<i>PHONON</i>	<i>PLUGIN</i>	<i>PLUMED</i>	<i>POEMS</i>	<i>PTM</i>
<i>PYTHON</i>	<i>QEQ</i>	<i>QMMM</i>	<i>QTB</i>	<i>REACTION</i>	<i>REAXFF</i>
<i>REPLICA</i>	<i>RIGID</i>	<i>SCAFACOS</i>	<i>SHOCK</i>	<i>SMTBQ</i>	<i>SPH</i>
<i>SPIN</i>	<i>SRD</i>	<i>TALLY</i>	<i>UEF</i>	<i>VORONOI</i>	<i>VTK</i>
<i>YAFF</i>					

## 6.2.1 ADIOS package

### Contents:

ADIOS is a high-performance I/O library. This package implements the *dump atom/adios*, *dump custom/adios* and *read\_dump ... format adios* commands to write and read data using the ADIOS library.

**Authors:** Norbert Podhorszki (ORNL) from the ADIOS developer team.

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/ADIOS: filenames -> commands
- src/ADIOS/README
- examples/PACKAGES/adios
- <https://github.com/ornladios/ADIOS2>
- *dump atom/adios*
- *dump custom/adios*
- *read\_dump*

## 6.2.2 AMOEBA package

### Contents:

Implementation of the AMOEBA and HIPPO polarized force fields originally developed by Jay Ponder's group at the U Washington at St Louis. The LAMMPS implementation is based on Fortran 90 code provided by the Ponder group in their [Tinker MD](#) software.

**Authors:** Josh Rackers and Steve Plimpton (Sandia), Trung Nguyen (U Chicago)

### Supporting info:

- `src/AMOEBA`: filenames -> commands
  - *AMOEBA and HIPPO howto*
  - *pair\_style amoeba*
  - *pair\_style hippo*
  - *atom\_style amoeba*
  - *angle\_style amoeba*
  - *improper\_style amoeba*
  - *fix amoeba/bitorsion*
  - *fix amoeba/pitorsion*
  - `tools/tinker/tinker2Imp.py`
  - `examples/amoeba`
- 

## 6.2.3 ASPHERE package

### Contents:

Computes, time-integration fixes, and pair styles for aspherical particle models including ellipsoids, 2d lines, and 3d triangles.

### Supporting info:

- `src/ASPHERE`: filenames -> commands
  - *Howto spherical*
  - *pair\_style gayberne*
  - *pair\_style resquared*
  - `doc/PDF/pair_gayberne_extra.pdf`
  - `doc/PDF/pair_resquared_extra.pdf`
  - `examples/ASPHERE`
  - `examples/ellipse`
  - <https://www.lammps.org/movies.html#line>
  - <https://www.lammps.org/movies.html#tri>
-

## 6.2.4 ATC package

### Contents:

ATC stands for atoms-to-continuum. This package implements a *fix atc* command to either couple molecular dynamics with continuum finite element equations or perform on-the-fly conversion of atomic information to continuum fields.

**Authors:** Reese Jones, Jeremy Templeton, Jon Zimmerman (Sandia).

### Install:

This package has *specific installation instructions* on the *Build extras* page. The ATC package requires that also the **MANYBODY** package is installed.

### Supporting info:

- src/ATC: filenames -> commands
  - src/ATC/README
  - *fix atc*
  - examples/PACKAGES/atc
  - <https://www.lammps.org/pictures.html#atc>
- 

## 6.2.5 AWPMD package

### Contents:

AWPMD stands for Antisymmetrized Wave Packet Molecular Dynamics. This package implements an atom, pair, and fix style which allows electrons to be treated as explicit particles in a classical molecular dynamics model.

**Author:** Ilya Valuev (JIHT, Russia).

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/AWPMD: filenames -> commands
  - src/AWPMD/README
  - *pair\_style awpmd/cut*
  - examples/PACKAGES/awpmd
- 

## 6.2.6 BOCS package

### Contents:

This package provides *fix bocs*, a modified version of *fix npt* which includes the pressure correction to the barostat as outlined in:

N. J. H. Dunn and W. G. Noid, “Bottom-up coarse-grained models that accurately describe the structure, pressure, and compressibility of molecular liquids”, J. Chem. Phys. 143, 243148 (2015).

**Authors:** Nicholas J. H. Dunn and Michael R. DeLyser (The Pennsylvania State University)

**Supporting info:**

The BOCS package for LAMMPS is part of the BOCS software package: <https://github.com/noid-group/BOCS>

See the following reference for information about the entire package:

Dunn, NJH; Lebold, KM; DeLyser, MR; Rudzinski, JF; Noid, WG. “BOCS: Bottom-Up Open-Source Coarse-Graining Software.” J. Phys. Chem. B. 122, 13, 3363-3377 (2018).

Example inputs are in the examples/PACKAGES/bocs folder.

---

## 6.2.7 BODY package

**Contents:**

Body-style particles with internal structure. Computes, time-integration fixes, pair styles, as well as the body styles themselves. See the *Howto body* page for an overview.

**Supporting info:**

- src/BODY filenames -> commands
  - *Howto\_body*
  - *atom\_style body*
  - *fix nve/body*
  - *pair\_style body/nparticle*
  - examples/body
- 

## 6.2.8 BPM package

**Contents:**

Pair styles, bond styles, fixes, and computes for bonded particle models for mesoscale simulations of solids and fracture. See the *Howto bpm* page for an overview.

**Authors:** Joel T. Clemmer (Sandia National Labs)

New in version 4May2022.

**Supporting info:**

- src/BPM filenames -> commands
  - *Howto\_bpm*
  - *atom\_style bpm/sphere*
  - *bond\_style bpm/rotational*
  - *bond\_style bpm/spring*
  - *compute nbond/atom*
  - *fix nve/bpm/sphere*
  - *pair\_style bpm/spring*
-

- examples/bpm
- 

## 6.2.9 BROWNIAN package

### Contents:

This package provides *fix brownian*, *fix brownian/sphere*, and *fix brownian/asphere* as well as *fix propel/self* which allow to do Brownian Dynamics time integration of point, spherical and aspherical particles and also support self-propelled particles.

**Authors:** Sam Cameron (University of Bristol), Stefan Paquay (while at Brandeis University) (initial version of *fix propel/self*)

Example inputs are in the examples/PACKAGES/brownian folder.

---

## 6.2.10 CG-DNA package

### Contents:

Several pair styles, bond styles, and integration fixes for coarse-grained modelling of single- and double-stranded DNA and RNA based on the oxDNA and oxRNA model of Doye, Louis and Ouldridge. The package includes Langevin-type rigid-body integrators with improved stability.

**Author:** Oliver Henrich (University of Strathclyde, Glasgow).

### Install:

The CG-DNA package requires that also the [MOLECULE](#) and [ASPHERE](#) packages are installed.

### Supporting info:

- src/CG-DNA: filenames -> commands
  - /src/CG-DNA/README
  - *pair\_style oxdna/\**
  - *pair\_style oxdna2/\**
  - *pair\_style oxrna2/\**
  - *bond\_style oxdna/\**
  - *bond\_style oxdna2/\**
  - *bond\_style oxrna2/\**
  - *fix nve/dotc/langevin*
-



### 6.2.11 CG-SPICA package

**Contents:**

Several pair styles and an angle style which implement the coarse-grained SPICA (formerly called SDK) model which enables simulation of biological or soft material systems.

**Original Author:** Axel Kohlmeyer (Temple U).

**Maintainers:** Yusuke Miyazaki and Wataru Shinoda (Okayama U).

**Supporting info:**

- src/CG-SPICA: filenames -> commands
  - src/CG-SPICA/README
  - *pair\_style lj/spica/\**
  - *angle\_style spica*
  - examples/PACKAGES/cgspica
  - <https://www.lammps.org/pictures.html#cg>
  - <https://www.spica-ff.org/>
- 

### 6.2.12 CLASS2 package

**Contents:**

Bond, angle, dihedral, improper, and pair styles for the COMPASS CLASS2 molecular force field.

**Supporting info:**

- src/CLASS2: filenames -> commands
  - *bond\_style class2*
  - *angle\_style class2*
  - *dihedral\_style class2*
  - *improper\_style class2*
  - *pair\_style lj/class2*
- 

### 6.2.13 COLLOID package

**Contents:**

Coarse-grained finite-size colloidal particles. Pair styles and fix wall styles for colloidal interactions. Includes the Fast Lubrication Dynamics (FLD) method for hydrodynamic interactions, which is a simplified approximation to Stokesian dynamics.

**Authors:** This package includes Fast Lubrication Dynamics pair styles which were created by Amit Kumar and Michael Bybee from Jonathan Higdon's group at UIUC.

**Supporting info:**

- src/COLLOID: filenames -> commands
  - *fix wall/colloid*
  - *pair\_style colloid*
  - *pair\_style yukawa/colloid*
  - *pair\_style brownian*
  - *pair\_style lubricate*
  - *pair\_style lubricateU*
  - examples/colloid
  - examples/srd
- 

## 6.2.14 COLVARS package

### Contents:

COLVARS stands for collective variables, which can be used to implement various enhanced sampling methods, including Adaptive Biasing Force, Metadynamics, Steered MD, Umbrella Sampling and Restraints. A *fix colvars* command is implemented which wraps a COLVARS library, which implements these methods. simulations.

**Authors:** The COLVARS library is written and maintained by Giacomo Fiorin (ICMS, Temple University, Philadelphia, PA, USA) and Jerome Henin (LISM, CNRS, Marseille, France), originally for the NAMD MD code, but with portability in mind. Axel Kohlmeyer (Temple U) provided the interface to LAMMPS.

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/COLVARS: filenames -> commands
  - doc/PDF/colvars-refman-lammps.pdf
  - src/COLVARS/README
  - lib/colvars/README
  - *fix colvars*
  - examples/PACKAGES/colvars
- 

## 6.2.15 COMPRESS package

### Contents:

Compressed output of dump files via the zlib compression library, using dump styles with a “gz” in their style name.

To use this package you must have the zlib compression library available on your system.

**Author:** Axel Kohlmeyer (Temple U).

### Install:

This package has *specific installation instructions* on the *Build extras* page.

---

**Supporting info:**

- src/COMPRESS: filenames -> commands
  - src/COMPRESS/README
  - lib/compress/README
  - *dump atom/gz*
  - *dump cfg/gz*
  - *dump custom/gz*
  - *dump xyz/gz*
- 

### 6.2.16 CORESHELL package

**Contents:**

Compute and pair styles that implement the adiabatic core/shell model for polarizability. The pair styles augment Born, Buckingham, and Lennard-Jones styles with core/shell capabilities. The *compute temp/cs* command calculates the temperature of a system with core/shell particles. See the *Howto coreshell* page for an overview of how to use this package.

**Author:** Hendrik Heenen (Technical U of Munich).

**Supporting info:**

- src/CORESHELL: filenames -> commands
  - *Howto coreshell*
  - *Howto polarizable*
  - *compute temp/cs*
  - *pair\_style born/coul/long/cs*
  - *pair\_style buck/coul/long/cs*
  - *pair\_style lj/cut/coul/long/cs*
  - *examples/coreshell*
- 

### 6.2.17 DIELECTRIC package

**Contents:**

An atom style, multiple pair styles, several fixes, Kspace styles and a compute for simulating systems using boundary element solvers for computing the induced charges at the interface between two media with different dielectric constants.

**Install:**

To use this package, also the *KSPACE* and *EXTRA-PAIR* packages need to be installed.

**Author:** Trung Nguyen and Monica Olvera de la Cruz (Northwestern U)

**Supporting info:**

- src/DIELECTRIC: filenames -> commands

- *atom\_style dielectric*
  - *pair\_style coul/cut/dielectric*
  - *pair\_style coul/long/dielectric*
  - *pair\_style lj/cut/coul/cut/dielectric*
  - *pair\_style lj/cut/coul/debye/dielectric*
  - *pair\_style lj/cut/coul/long/dielectric*
  - *pair\_style lj/cut/coul/msm/dielectric*
  - *pair\_style ppm/dielectric*
  - *pair\_style ppm/disp/dielectric*
  - *pair\_style msm/dielectric*
  - *fix\_style polarize/bem/icc*
  - *fix\_style polarize/bem/gmres*
  - *fix\_style polarize/functional*
  - *compute efield/atom*
  - examples/PACKAGES/dielectric
- 

## 6.2.18 DIFFRACTION package

### Contents:

Two computes and a fix for calculating x-ray and electron diffraction intensities based on kinematic diffraction theory.

**Author:** Shawn Coleman while at the U Arkansas.

### Supporting info:

- src/DIFFRACTION: filenames -> commands
  - *compute saed*
  - *compute xrd*
  - *fix saed/vtk*
  - examples/PACKAGES/diffraction
- 

## 6.2.19 DIPOLE package

### Contents:

An atom style and several pair styles for point dipole models with short-range or long-range interactions.

### Supporting info:

- src/DIPOLE: filenames -> commands
  - *atom\_style dipole*
-

- *pair\_style lj/cut/dipole/cut*
  - *pair\_style lj/cut/dipole/long*
  - *pair\_style lj/long/dipole/long*
  - *angle\_style dipole*
  - examples/dipole
- 

## 6.2.20 DPD-BASIC package

### Contents:

Pair styles for the basic dissipative particle dynamics (DPD) method and DPD thermostatting.

**Author:** Kurt Smith (U Pittsburgh), Martin Svoboda, Martin Lisal (ICPF and UJEP)

### Supporting info:

- src/DPD-BASIC: filenames -> commands
  - *pair\_style dpd*
  - *pair\_style dpd/tstat*
  - *pair\_style dpd/ext*
  - *pair\_style dpd/ext/tstat*
  - examples/PACKAGES/dpd-basic
- 

## 6.2.21 DPD-MESO package

### Contents:

Several extensions of the dissipative particle dynamics (DPD) method. Specifically, energy-conserving DPD (eDPD) that can model non-isothermal processes, many-body DPD (mDPD) for simulating vapor-liquid coexistence, and transport DPD (tDPD) for modeling advection-diffusion-reaction systems. The equations of motion of these DPD extensions are integrated through a modified velocity-Verlet (MVV) algorithm.

**Author:** Zhen Li (Department of Mechanical Engineering, Clemson University)

### Supporting info:

- src/DPD-MESO: filenames -> commands
- src/DPD-MESO/README
- *atom\_style edpd*
- *pair\_style edpd*
- *pair\_style mdpd*
- *pair\_style tdpd*
- *fix mvv/dpd*
- examples/PACKAGES/mesodpd

- <https://www.lammps.org/movies.html#mesodpd>
- 

## 6.2.22 DPD-REACT package

### Contents:

DPD stands for dissipative particle dynamics. This package implements coarse-grained DPD-based models for energetic, reactive molecular crystalline materials. It includes many pair styles specific to these systems, including for reactive DPD, where each particle has internal state for multiple species and a coupled set of chemical reaction ODEs are integrated each timestep. Highly accurate time integrators for isothermal, isoenergetic, isobaric and isenthalpic conditions are included. These enable long timesteps via the Shardlow splitting algorithm.

**Authors:** Jim Larentzos (ARL), Tim Mattox (Engility Corp), and John Brennan (ARL).

### Supporting info:

- src/DPD-REACT: filenames -> commands
  - /src/DPD-REACT/README
  - *compute dpd*
  - *compute dpd/atom*
  - *fix eos/cv*
  - *fix eos/table*
  - *fix eos/table/rx*
  - *fix shardlow*
  - *fix rx*
  - *pair\_style table/rx*
  - *pair\_style dpd/fdt*
  - *pair\_style dpd/fdt/energy*
  - *pair\_style exp6/rx*
  - *pair\_style multi/lucy*
  - *pair\_style multi/lucy/rx*
  - examples/PACKAGES/dpd-react
- 

## 6.2.23 DPD-SMOOTH package

### Contents:

A pair style for smoothed dissipative particle dynamics (SDPD), which is an extension of smoothed particle hydrodynamics (SPH) to mesoscale where thermal fluctuations are important (see the *SPH package*). Also two fixes for moving and rigid body integration of SPH/SDPD particles (particles of atom\_style meso).

**Author:** Morteza Jalalvand (Institute for Advanced Studies in Basic Sciences, Iran).

### Supporting info:

- src/DPD-SMOOTH: filenames -> commands
  - src/DPD-SMOOTH/README
  - *pair\_style sdpd/taitwater/isothermal*
  - *fix meso/move*
  - *fix rigid/meso*
  - examples/PACKAGES/dpd-smooth
- 

### 6.2.24 DRUDE package

#### Contents:

Fixes, pair styles, and a compute to simulate thermalized Drude oscillators as a model of polarization. See the *Howto drude* and *Howto drude2* pages for an overview of how to use the package. There are auxiliary tools for using this package in tools/drude.

**Authors:** Alain Dequidt (U Clermont Auvergne), Julien Devemy (CNRS), and Agilio Padua (ENS de Lyon).

#### Supporting info:

- src/DRUDE: filenames -> commands
  - *Howto drude*
  - *Howto drude2*
  - *Howto polarizable*
  - src/DRUDE/README
  - *fix drude*
  - *fix drude/transform/\**
  - *compute temp/drude*
  - *pair\_style thole*
  - *pair\_style lj/cut/thole/long*
  - examples/PACKAGES/drude
  - tools/drude
- 

### 6.2.25 EFF package

#### Contents:

EFF stands for electron force field which allows a classical MD code to model electrons as particles of variable radius. This package contains atom, pair, fix and compute styles which implement the eFF as described in A. Jaramillo-Botero, J. Su, Q. An, and W.A. Goddard III, JCC, 2010. The eFF potential was first introduced by Su and Goddard, in 2007. There are auxiliary tools for using this package in tools/eff; see its README file.

**Author:** Andres Jaramillo-Botero (CalTech).

#### Supporting info:

- src/EFF: filenames -> commands
  - src/EFF/README
  - *atom\_style electron*
  - *fix nve/eff*
  - *fix nvt/eff*
  - *fix npt/eff*
  - *fix langevin/eff*
  - *compute temp/eff*
  - *pair\_style eff/cut*
  - *pair\_style eff/inline*
  - examples/PACKAGES/eff
  - tools/eff/README
  - tools/eff
  - <https://www.lammps.org/movies.html#eff>
- 

## 6.2.26 ELECTRODE package

### Contents:

The ELECTRODE package allows the user to enforce a constant potential method for groups of atoms that interact with the remaining atoms as electrolyte.

**Authors:** The ELECTRODE library is written and maintained by Ludwig Ahrens-Iwers (TUHH, Hamburg, Germany), Shern Tee (UQ, Brisbane, Australia) and Robert Meissner (TUHH, Hamburg, Germany).

New in version 4May2022.

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- *fix electrode/conp*
- 

## 6.2.27 EXTRA-COMPUTE package

### Contents:

Additional compute styles that are less commonly used.

### Supporting info:

- src/EXTRA-COMPUTE: filenames -> commands
  - *compute*
-



### 6.2.28 EXTRA-DUMP package

**Contents:**

Additional dump styles that are less commonly used.

**Supporting info:**

- src/EXTRA-DUMP: filenames -> commands
  - *dump*
- 

### 6.2.29 EXTRA-FIX package

**Contents:**

Additional fix styles that are less commonly used.

**Supporting info:**

- src/EXTRA-FIX: filenames -> commands
  - *fix*
- 

### 6.2.30 EXTRA-MOLECULE package

**Contents:**

Additional bond, angle, dihedral, and improper styles that are less commonly used.

**Install:**

To use this package, also the *MOLECULE* package needs to be installed.

**Supporting info:**

- src/EXTRA-MOLECULE: filenames -> commands
  - *molecular styles*
- 

### 6.2.31 EXTRA-PAIR package

**Contents:**

Additional pair styles that are less commonly used.

**Supporting info:**

- src/EXTRA-PAIR: filenames -> commands
  - *pair\_style*
-

## 6.2.32 FEP package

### Contents:

FEP stands for free energy perturbation. This package provides methods for performing FEP simulations by using a *fix adapt/fep* command with soft-core pair potentials, which have a “soft” in their style name. There are auxiliary tools for using this package in tools/fep; see its README file.

**Author:** Agilio Padua (ENS de Lyon)

### Supporting info:

- src/FEP: filenames -> commands
  - src/FEP/README
  - *fix adapt/fep*
  - *compute fep*
  - *pair\_style \*/soft*
  - examples/PACKAGES/fep
  - tools/fep/README
  - tools/fep
- 

## 6.2.33 GPU package

### Contents:

Dozens of pair styles and a version of the PPPM long-range Coulombic solver optimized for GPUs. All such styles have a “gpu” as a suffix in their style name. The GPU code can be compiled with either CUDA or OpenCL, however the OpenCL variants are no longer actively maintained and only the CUDA versions are regularly tested. The *GPU package* page gives details of what hardware and GPU software is required on your system, and details on how to build and use this package. Its styles can be invoked at run time via the “-sf gpu” or “-suffix gpu” *command-line switches*. See also the *KOKKOS* package, which has GPU-enabled styles.

**Authors:** Mike Brown (Intel) while at Sandia and ORNL and Trung Nguyen (Northwestern U) while at ORNL and later. AMD HIP support by Evgeny Kuznetsov, Vladimir Stegailov, and Vsevolod Nikolskiy (HSE University).

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/GPU: filenames -> commands
- src/GPU/README
- lib/gpu/README
- *Accelerator packages*
- *GPU package*
- *Section 2.6 -sf gpu*
- *Section 2.6 -pk gpu*
- *package gpu*

- *Commands* pages (*pair*, *kpace*) for styles followed by (g)
  - *Benchmarks* page of website
- 

### 6.2.34 GRANULAR package

#### Contents:

Pair styles and fixes for finite-size granular particles, which interact with each other and boundaries via frictional and dissipative potentials.

#### Supporting info:

- src/GRANULAR: filenames -> commands
  - *Howto granular*
  - *fix pour*
  - *fix wall/gran*
  - *pair\_style gran/hooke*
  - *pair\_style gran/hertz/history*
  - examples/granregion
  - examples/pour
  - bench/in.chute
  - <https://www.lammps.org/pictures.html#jamming>
  - <https://www.lammps.org/movies.html#hopper>
  - <https://www.lammps.org/movies.html#dem>
  - <https://www.lammps.org/movies.html#brazil>
  - <https://www.lammps.org/movies.html#granregion>
- 

### 6.2.35 H5MD package

#### Contents:

H5MD stands for HDF5 for MD. [HDF5](#) is a portable, binary, self-describing file format, used by many scientific simulations. H5MD is a format for molecular simulations, built on top of HDF5. This package implements a *dump h5md* command to output LAMMPS snapshots in this format.

To use this package you must have the HDF5 library available on your system.

**Author:** Pierre de Buyl (KU Leuven) created both the package and the H5MD format.

#### Install:

This package has *specific installation instructions* on the *Build extras* page.

#### Supporting info:

- src/H5MD: filenames -> commands

- [src/H5MD/README](#)
  - [lib/h5md/README](#)
  - [dump h5md](#)
- 

## 6.2.36 INTEL package

### Contents:

Dozens of pair, fix, bond, angle, dihedral, improper, and kspace styles which are optimized for Intel CPUs and KNLs (Knights Landing). All of them have an “intel” in their style name. The [INTEL package](#) page gives details of what hardware and compilers are required on your system, and how to build and use this package. Its styles can be invoked at run time via the “-sf intel” or “-suffix intel” *command-line switches*. Also see the [KOKKOS](#), [OPT](#), and [OPENMP](#) packages, which have styles optimized for CPUs and KNLs.

You need to have an Intel compiler, version 14 or higher to take full advantage of this package. While compilation with GNU compilers is supported, performance will be sub-optimal.

---

**Note:** the INTEL package contains styles that require using the -restrict flag, when compiling with Intel compilers.

---

**Author:** Mike Brown (Intel).

### Install:

This package has *specific installation instructions* on the [Build extras](#) page.

### Supporting info:

- [src/INTEL: filenames -> commands](#)
  - [src/INTEL/README](#)
  - [Accelerator packages](#)
  - [INTEL package](#)
  - [Section 2.6 -sf intel](#)
  - [Section 2.6 -pk intel](#)
  - [package intel](#)
  - Search the [commands](#) pages ([fix](#), [compute](#), [pair](#), [bond](#), [angle](#), [dihedral](#), [improper](#), [kspace](#)) for styles followed by (i)
  - [src/INTEL/TEST](#)
  - [Benchmarks](#) page of website
-

## 6.2.37 INTERLAYER package

### Contents:

A collection of pair styles specifically to be used for modeling layered materials, most commonly graphene sheets (or equivalents).

### Supporting info:

- src/INTERLAYER: filenames -> commands
  - *Pair style* page
  - examples/PACKAGES/interlayer
- 

## 6.2.38 KIM package

### Contents:

This package contains a command with a set of sub-commands that serve as a wrapper on the [Open Knowledgebase of Interatomic Models \(OpenKIM\)](#) repository of interatomic models (IMs) enabling compatible ones to be used in LAMMPS simulations.

This includes *kim init*, and *kim interactions* commands to select, initialize and instantiate the IM, a *kim query* command to perform web queries for material property predictions of OpenKIM IMs, a *kim param* command to access KIM Model Parameters from LAMMPS, and a *kim property* command to write material properties computed in LAMMPS to standard KIM property instance format.

Support for KIM IMs that conform to the [KIM Application Programming Interface \(API\)](#) is provided by the *pair\_style kim* command.

---

**Note:** The command *pair\_style kim* is called by *kim interactions* and is not recommended to be directly used in input scripts.

---

To use this package you must have the KIM API library available on your system. The KIM API is available for download on the [OpenKIM website](#). When installing LAMMPS from binary, the kim-api package is a dependency that is automatically downloaded and installed.

Information about the KIM project can be found at its website: <https://openkim.org>. The KIM project is led by Ellad Tadmor and Ryan Elliott (U Minnesota) and is funded by the [National Science Foundation](#).

**Authors:** Ryan Elliott (U Minnesota) is the main developer for the KIM API and the *pair\_style kim* command. Yaser Afshar (U Minnesota), Axel Kohlmeyer (Temple U), Ellad Tadmor (U Minnesota), and Daniel Karls (U Minnesota) contributed to the *kim command* interface in close collaboration with Ryan Elliott.

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- *kim command*
  - *pair\_style kim*
  - src/KIM: filenames -> commands
  - src/KIM/README
-

- [lib/kim/README](#)
  - [examples/kim](#)
- 

## 6.2.39 KOKKOS package

### Contents:

Dozens of atom, pair, bond, angle, dihedral, improper, fix, compute styles adapted to compile using the Kokkos library which can convert them to OpenMP or CUDA code so that they run efficiently on multicore CPUs, KNLs, or GPUs. All the styles have a “kk” as a suffix in their style name. The *KOKKOS package* page gives details of what hardware and software is required on your system, and how to build and use this package. Its styles can be invoked at run time via the “-sf kk” or “-suffix kk” *command-line switches*. Also see the *GPU*, *OPT*, *INTEL*, and *OPENMP* packages, which have styles optimized for CPUs, KNLs, and GPUs.

You must have a C++14 compatible compiler to use this package. KOKKOS makes extensive use of advanced C++ features, which can expose compiler bugs, especially when compiling for maximum performance at high optimization levels. Please see the file [lib/kokkos/README](#) for a list of compilers and their respective platforms, that are known to work.

**Authors:** The KOKKOS package was created primarily by Christian Trott and Stan Moore (Sandia), with contributions from other folks as well. It uses the open-source [Kokkos library](#) which was developed by Carter Edwards, Christian Trott, and others at Sandia, and which is included in the LAMMPS distribution in [lib/kokkos](#).

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- [src/KOKKOS: filenames -> commands](#)
  - [src/KOKKOS/README](#)
  - [lib/kokkos/README](#)
  - *Accelerator packages*
  - *KOKKOS package*
  - *Section 2.6 -k on ...*
  - *Section 2.6 -sf kk*
  - *Section 2.6 -pk kokkos*
  - *package kokkos*
  - Search the *commands* pages (*fix, compute, pair, bond, angle, dihedral, improper, kspace*) for styles followed by (k)
  - [Benchmarks page of website](#)
-

## 6.2.40 KSPACE package

### Contents:

A variety of long-range Coulombic solvers, as well as pair styles which compute the corresponding short-range pairwise Coulombic interactions. These include Ewald, particle-particle particle-mesh (PPPM), and multilevel summation method (MSM) solvers.

### Install:

Building with this package requires a 1d FFT library be present on your system for use by the PPPM solvers. This can be the KISS FFT library provided with LAMMPS, third party libraries like FFTW, or a vendor-supplied FFT library. See the *Build settings* page for details on how to select different FFT options for your LAMPMS build.

### Supporting info:

- `src/KSPACE`: filenames -> commands
  - *kspace\_style*
  - `doc/PDF/kspace.pdf`
  - *Howto tip3p*
  - *Howto tip4p*
  - *Howto spc*
  - *pair\_style coul*
  - Search the *pair style* page for styles with “long” or “msm” in name
  - `examples/peptide`
  - `bench/in.rhodo`
- 

## 6.2.41 LATBOLTZ package

### Contents:

Fixes which implement a background Lattice-Boltzmann (LB) fluid, which can be used to model MD particles influenced by hydrodynamic forces.

**Authors:** Frances Mackay and Colin Denniston (University of Western Ontario).

### Install:

The LATBOLTZ package requires that LAMMPS is build in *MPI parallel mode*.

### Supporting info:

- `src/LATBOLTZ`: filenames -> commands
  - `src/LATBOLTZ/README`
  - *fix lb/fluid*
  - *fix lb/momentum*
  - *fix lb/viscous*
  - `examples/PACKAGES/latboltz`
-

## 6.2.42 LATTE package

### Contents:

A fix command which wraps the LATTE DFTB code, so that molecular dynamics can be run with LAMMPS using density-functional tight-binding quantum forces calculated by LATTE.

More information on LATTE can be found at this website: <https://github.com/lanl/LATTE>. A brief technical description is given with the *fix latte* command.

**Authors:** Christian Negre (LANL) and Steve Plimpton (Sandia). LATTE itself is developed at Los Alamos National Laboratory by Marc Cawkwell, Anders Niklasson, and Christian Negre.

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/LATTE: filenames -> commands
  - src/LATTE/README
  - lib/latte/README
  - *fix latte*
  - examples/latte
  - LAMMPS-LATTE tutorial
- 

## 6.2.43 MACHDYN package

### Contents:

An atom style, fixes, computes, and several pair styles which implements smoothed Mach dynamics (SMD) for solids, which is a model related to smoothed particle hydrodynamics (SPH) for liquids (see the *SPH package*).

This package solves solids mechanics problems via a state of the art stabilized meshless method with hourglass control. It can specify hydrostatic interactions independently from material strength models, i.e. pressure and deviatoric stresses are separated. It provides many material models (Johnson-Cook, plasticity with hardening, Mie-Grueneisen, Polynomial EOS) and allows new material models to be added. It implements rigid boundary conditions (walls) which can be specified as surface geometries from \*.STL files.

**Author:** Georg Ganzenmuller (Fraunhofer-Institute for High-Speed Dynamics, Ernst Mach Institute, Germany).

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/MACHDYN: filenames -> commands
  - src/MACHDYN/README
  - doc/PDF/MACHDYN\_LAMMPS\_userguide.pdf
  - examples/PACKAGES/machdyn
  - <https://www.lammps.org/movies.html#smd>
-



## 6.2.44 MANIFOLD package

### Contents:

Several fixes and a “manifold” class which enable simulations of particles constrained to a manifold (a 2D surface within the 3D simulation box). This is done by applying the RATTLE constraint algorithm to formulate single-particle constraint functions  $g(x_i, y_i, z_i) = 0$  and their derivative (i.e. the normal of the manifold)  $n = \text{grad}(g)$ .

**Author:** Stefan Paquay (until 2017: Eindhoven University of Technology (TU/e), The Netherlands; since 2017: Brandeis University, Waltham, MA, USA)

### Supporting info:

- src/MANIFOLD: filenames -> commands
  - src/MANIFOLD/README
  - *Howto manifold*
  - *fix manifoldforce*
  - *fix nve/manifold/rattle*
  - *fix nvt/manifold/rattle*
  - examples/PACKAGES/manifold
  - <https://www.lammps.org/movies.html#manifold>
- 

## 6.2.45 MANYBODY package

### Contents:

A variety of many-body and bond-order potentials. These include (AI)REBO, BOP, EAM, EIM, Stillinger-Weber, and Tersoff potentials.

### Supporting info:

- src/MANYBODY: filenames -> commands
  - *Pair style* page
  - examples/comb
  - examples/eim
  - examples/nb3d
  - examples/shear
  - examples/streitz
  - examples/vashishta
  - bench/in.eam
-

## 6.2.46 MC package

### Contents:

Several fixes and a pair style that have Monte Carlo (MC) or MC-like attributes. These include fixes for creating, breaking, and swapping bonds, for performing atomic swaps, and performing grand-canonical MC (GCMC) or similar processes in conjunction with dynamics.

### Supporting info:

- src/MC: filenames -> commands
  - *fix atom/swap*
  - *fix bond/break*
  - *fix bond/create*
  - *fix bond/create/angle*
  - *fix bond/swap*
  - *fix charge/regulation*
  - *fix gcmc*
  - *fix tfmc*
  - *fix widom*
  - *pair\_style dsmc*
  - <https://www.lammps.org/movies.html#gcmc>
- 

## 6.2.47 MDI package

### Contents:

A LAMMPS command and fixes to allow client-server coupling of LAMMPS to other atomic or molecular simulation codes or materials modeling workflows via the [MolSSI Driver Interface \(MDI\)](#) library.

**Author:** Taylor Barnes - MolSSI, [taylor.a.barnes at gmail.com](mailto:taylor.a.barnes@gmail.com)

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/MDI/README
  - lib/mdi/README
  - *Howto MDI*
  - *mdi*
  - *fix mdi/qm*
  - examples/PACKAGES/mdi
-

## 6.2.48 MEAM package

### Contents:

A pair style for the modified embedded atom (MEAM) potential translated from the Fortran version in the (obsolete) MEAM package to plain C++. The MEAM fully replaces the MEAM package, which has been removed from LAMMPS after the 12 December 2018 version.

**Author:** Sebastian Huetter, (Otto-von-Guericke University Magdeburg) based on the Fortran version of Greg Wagner (Northwestern U) while at Sandia.

### Supporting info:

- src/MEAM: filenames -> commands
  - src/MEAM/README
  - *pair\_style meam*
  - examples/meam
- 

## 6.2.49 MESONT package

### Contents:

MESONT is a LAMMPS package for simulation of nanomechanics of nanotubes (NTs). The model is based on a coarse-grained representation of NTs as “flexible cylinders” consisting of a variable number of segments. Internal interactions within a NT and the van der Waals interaction between the tubes are described by a mesoscopic force field designed and parameterized based on the results of atomic-level molecular dynamics simulations. The description of the force field is provided in the papers listed below.

This package contains two independent implementations of this model: *pair\_style mesont/tpm* is the original implementation of the model based on a Fortran library in the `lib/mesont` folder. The second implementation is provided by the *mesocnt* styles (*bond\_style mesocnt*, *angle\_style mesocnt* and *pair\_style mesocnt*). The *mesocnt* implementation has the same features as the original implementation with the addition of friction, but is directly implemented in C++, interfaces more cleanly with general LAMMPS functionality, and is typically faster. It also does not require its own atom style and can be installed without any external libraries.

### Download of potential files:

The potential files for these pair styles are *very* large and thus are not included in the regular downloaded packages of LAMMPS or the git repositories. Instead, they will be automatically downloaded from a web server when the package is installed for the first time.

### Authors of the \*mesont\* styles:

Maxim V. Shugaev (University of Virginia), Alexey N. Volkov (University of Alabama), Leonid V. Zhigilei (University of Virginia)

**Author of the \*mesocnt\* styles:** Philipp Kloza (U Cambridge)

### Supporting info:

- src/MESONT: filenames -> commands
- src/MESONT/README
- *atom\_style mesont*
- *pair\_style mesont/tpm*

- *compute mesont*
  - *bond\_style mesocnt*
  - *angle\_style mesocnt*
  - *pair\_style mesocnt*
  - examples/PACKAGES/mesont
  - tools/mesont
- 

### 6.2.50 MGPT package

#### Contents:

A pair style which provides a fast implementation of the quantum-based MGPT multi-ion potentials. The MGPT or model GPT method derives from first-principles DFT-based generalized pseudopotential theory (GPT) through a series of systematic approximations valid for mid-period transition metals with nearly half-filled d bands. The MGPT method was originally developed by John Moriarty at LLNL. The pair style in this package calculates forces and energies using an optimized matrix-MGPT algorithm due to Tomas Opperstrup at LLNL.

**Authors:** Tomas Opperstrup and John Moriarty (LLNL).

#### Supporting info:

- src/MGPT: filenames -> commands
  - src/MGPT/README
  - *pair\_style mgpt*
  - examples/PACKAGES/mgpt
- 

### 6.2.51 MISC package

#### Contents:

A variety of compute, fix, pair, bond styles with specialized capabilities that don't align with other packages. Do a directory listing, "ls src/MISC", to see the list of commands.

**Note:** the MISC package contains styles that require using the -restrict flag, when compiling with Intel compilers.

---

#### Supporting info:

- src/MISC: filenames -> commands
  - *bond\_style special*
  - *compute viscosity/cos*
  - *fix accelerate/cos*
  - *fix imd*
  - *fix ipi*
  - *pair\_style agni*
-

- *pair\_style list*
  - *pair\_style srp*
  - *pair\_style tracker*
- 

## 6.2.52 ML-HDNNP package

### Contents:

A *pair\_style hdnnp* command which allows to use high-dimensional neural network potentials (HDNNPs), a form of machine learning potentials. HDNNPs must be carefully trained prior to their application in a molecular dynamics simulation.

To use this package you must have the `n2p2` library installed and compiled on your system.

**Author:** Andreas Singraber

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/ML-HDNNP: filenames -> commands
  - src/ML-HDNNP/README
  - lib/hdnnp/README
  - *pair\_style hdnnp*
  - examples/PACKAGES/hdnnp
- 

## 6.2.53 ML-IAP package

### Contents:

A general interface for machine-learning interatomic potentials, including PyTorch.

### Install:

To use this package, also the *ML-SNAP* package needs to be installed. To make the *mliappy* model available, also the *PYTHON* package needs to be installed, the version of Python must be 3.6 or later, and the *cython* software must be installed.

**Author:** Aidan Thompson (Sandia), Nicholas Lubbers (LANL).

### Supporting info:

- src/ML-IAP: filenames -> commands
- src/ML-IAP/README.md
- *pair\_style mliap*
- *compute\_style mliap*
- examples/mliap (see README)

When built with the *mliappy* model this package includes an extension for coupling with Python models, including PyTorch. In this case, the Python interpreter linked to LAMMPS will need the *cython* and *numpy* modules installed. The provided examples build models with PyTorch, which would therefore also need to be installed to run those examples.

---

## 6.2.54 ML-PACE package

### Contents:

A pair style for the Atomic Cluster Expansion potential (ACE). ACE is a methodology for deriving a highly accurate classical potential fit to a large archive of quantum mechanical (DFT) data. The ML-PACE package provides an efficient implementation for running simulations with ACE potentials.

### Authors:

This package was written by Yury Lysogorskiy<sup>1</sup>, Cas van der Oord<sup>2</sup>, Anton Bochkarev<sup>1</sup>, Sarath Menon<sup>1</sup>, Matteo Rinaldi<sup>1</sup>, Thomas Hammerschmidt<sup>1</sup>, Matous Mrovec<sup>1</sup>, Aidan Thompson<sup>3</sup>, Gabor Csanyi<sup>2</sup>, Christoph Ortner<sup>4</sup>, Ralf Drautz<sup>1</sup>.

<sup>1</sup>: Ruhr-University Bochum, Bochum, Germany

<sup>2</sup>: University of Cambridge, Cambridge, United Kingdom

<sup>3</sup>: Sandia National Laboratories, Albuquerque, New Mexico, USA

<sup>4</sup>: University of British Columbia, Vancouver, BC, Canada

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- `src/ML-PACE: filenames -> commands`
  - *pair\_style pace*
  - `examples/PACKAGES/pace`
- 

## 6.2.55 ML-QUIP package

### Contents:

A *pair\_style quip* command which wraps the [QUIP libAtoms library](#), which includes a variety of interatomic potentials, including Gaussian Approximation Potential (GAP) models developed by the Cambridge University group.

To use this package you must have the QUIP libAtoms library available on your system.

**Author:** Albert Bartok (Cambridge University)

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- `src/ML-QUIP: filenames -> commands`
  - `src/ML-QUIP/README`
-

- *pair\_style quip*
  - examples/PACKAGES/quip
- 

### 6.2.56 ML-RANN package

#### Contents:

A pair style for using rapid atomistic neural network (RANN) potentials. These neural network potentials work by first generating a series of symmetry functions from the neighbor list and then using these values as the input layer of a neural network.

#### Authors:

This package was written by Christopher Barrett with contributions by Doyl Dickel, Mississippi State University.

#### Supporting info:

- src/ML-RANN: filenames -> commands
  - *pair\_style rann*
  - examples/PACKAGES/rann
- 

### 6.2.57 ML-SNAP package

#### Contents:

A pair style for the spectral neighbor analysis potential (SNAP). SNAP is methodology for deriving a highly accurate classical potential fit to a large archive of quantum mechanical (DFT) data. Also several computes which analyze attributes of the potential.

**Author:** Aidan Thompson (Sandia).

#### Supporting info:

- src/ML-SNAP: filenames -> commands
  - *pair\_style snap*
  - *compute sna/atom*
  - *compute sna/grid*
  - *compute sna/grid/local*
  - *compute snad/atom*
  - *compute snav/atom*
  - examples/snap
-

## 6.2.58 MOFFF package

### Contents:

Pair, angle and improper styles needed to employ the MOF-FF force field by Schmid and coworkers with LAMMPS. MOF-FF is a first principles derived force field with the primary aim to simulate MOFs and related porous framework materials, using spherical Gaussian charges. It is described in S. Bureekaew et al., Phys. Stat. Sol. B 2013, 250, 1128-1141. For the usage of MOF-FF see the example in the example directory as well as the [MOF+](#) website.

**Author:** Hendrik Heenen (Technical U of Munich), Rochus Schmid (Ruhr-University Bochum).

### Supporting info:

- src/MOFFF: filenames -> commands
  - src/MOFFF/README
  - *pair\_style buck6d/coul/gauss*
  - *angle\_style class2*
  - *angle\_style cosine/buck6d*
  - *improper\_style inversion/harmonic*
  - examples/PACKAGES/mofff
- 

## 6.2.59 MOLECULE package

### Contents:

A large number of atom, pair, bond, angle, dihedral, improper styles that are used to model molecular systems with fixed covalent bonds. The pair styles include the Dreiding (hydrogen-bonding) and CHARMM force fields, and a TIP4P water model.

### Supporting info:

- src/MOLECULE: filenames -> commands
- *atom\_style*
- *bond\_style*
- *angle\_style*
- *dihedral\_style*
- *improper\_style*
- *pair\_style hbond/dreiding/lj*
- *pair\_style lj/charmm/coul/charmm*
- *Howto bioFF*
- examples/cmap
- examples/dreiding
- examples/micelle,
- examples/peptide
- bench/in.chain



- [bench/in.rhodo](#)
- 

## 6.2.60 MOLFILE package

### Contents:

A *dump molfile* command which uses molfile plugins that are bundled with the VMD molecular visualization and analysis program, to enable LAMMPS to dump snapshots in formats compatible with various molecular simulation tools.

To use this package you must have the desired VMD plugins available on your system.

Note that this package only provides the interface code, not the plugins themselves, which will be accessed when requesting a specific plugin via the *dump molfile* command. Plugins can be obtained from a VMD installation which has to match the platform that you are using to compile LAMMPS for. By adding plugins to VMD, support for new file formats can be added to LAMMPS (or VMD or other programs that use them) without having to re-compile the application itself. More information about the VMD molfile plugins can be found at <http://www.ks.uiuc.edu/Research/vmd/plugins/molfile>.

**Author:** Axel Kohlmeyer (Temple U).

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/MOLFILE: filenames -> commands
  - src/MOLFILE/README
  - lib/molfile/README
  - *dump molfile*
- 

## 6.2.61 MPIIO package

### Contents:

Support for parallel output/input of dump and restart files via the MPIIO library. It adds *dump styles* with a “mpio” in their style name. Restart files with an “.mpio” suffix are also written and read in parallel.

**Warning:** The MPIIO package is currently unmaintained and has become unreliable. Use with caution.

### Install:

The MPIIO package requires that LAMMPS is build in *MPI parallel mode*.

### Supporting info:

- src/MPIIO: filenames -> commands
  - *dump*
  - *restart*
  - *write\_restart*
-

- *read\_restart*
- 

## 6.2.62 MSCG package

### Contents:

A *fix mscg* command which can parameterize a Multi-Scale Coarse-Graining (MSCG) model using the open-source MS-CG library.

To use this package you must have the MS-CG library available on your system.

**Authors:** The fix was written by Lauren Abbott (Sandia). The MS-CG library was developed by Jacob Wagner in Greg Voth's group at the University of Chicago.

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/MSCG: filenames -> commands
  - src/MSCG/README
  - lib/mscg/README
  - examples/mscg
- 

## 6.2.63 NETCDF package

### Contents:

Dump styles for writing NetCDF formatted dump files. NetCDF is a portable, binary, self-describing file format developed on top of HDF5. The file contents follow the AMBER NetCDF trajectory conventions (<http://ambermd.org/netcdf/nctraj.xhtml>), but include extensions.

To use this package you must have the NetCDF library available on your system.

Note that NetCDF files can be directly visualized with the following tools:

- **Ovito** (Ovito supports the AMBER convention and the extensions mentioned above)
- **VMD**

**Author:** Lars Pastewka (Karlsruhe Institute of Technology).

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/NETCDF: filenames -> commands
  - src/NETCDF/README
  - lib/netcdf/README
  - *dump netcdf*
-

## 6.2.64 OPENMP package

### Contents:

Hundreds of pair, fix, compute, bond, angle, dihedral, improper, and kspace styles which are altered to enable threading on many-core CPUs via OpenMP directives. All of them have an “omp” in their style name. The *OPENMP package* page gives details of what hardware and compilers are required on your system, and how to build and use this package. Its styles can be invoked at run time via the “-sf omp” or “-suffix omp” *command-line switches*. Also see the *KOKKOS*, *OPT*, and *INTEL* packages, which have styles optimized for CPUs.

**Author:** Axel Kohlmeyer (Temple U).

---

**Note:** To enable multi-threading support the compile flag “-fopenmp” and the link flag “-fopenmp” (for GNU compilers, you have to look up the equivalent flags for other compilers) must be used to build LAMMPS. When using Intel compilers, also the “-restrict” flag is required. The OPENMP package can be compiled without enabling OpenMP; then all code will be compiled as serial and the only improvement over the regular styles are some data access optimization. These flags should be added to the CCFLAGS and LINKFLAGS lines of your Makefile.machine. See src/MAKE/OPTIONS/Makefile.omp for an example.

---

Once you have an appropriate Makefile.machine, you can install/un-install the package and build LAMMPS in the usual manner:

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/OPENMP: filenames -> commands
  - src/OPENMP/README
  - *Accelerator packages*
  - *OPENMP package*
  - *Command line option -suffix/-sf omp*
  - *Command line option -package/-pk omp*
  - *package omp*
  - Search the *commands* pages (*fix*, *compute*, *pair*, *bond*, *angle*, *dihedral*, *improper*, *kspace*) for styles followed by (o)
  - *Benchmarks page of website*
- 

## 6.2.65 OPT package

### Contents:

A handful of pair styles which are optimized for improved CPU performance on single or multiple cores. These include EAM, LJ, CHARMM, and Morse potentials. The styles have an “opt” suffix in their style name. The *OPT package* page gives details of how to build and use this package. Its styles can be invoked at run time via the “-sf opt” or “-suffix opt” *command-line switches*. See also the *KOKKOS*, *INTEL*, and *OPENMP* packages, which have styles optimized for CPU performance.

**Authors:** James Fischer (High Performance Technologies), David Richie, and Vincent Natoli (Stone Ridge Technology).

**Install:**

This package has *specific installation instructions* on the *Build extras* page.

**Supporting info:**

- src/OPT: filenames -> commands
- *Accelerator packages*
- *OPT package*
- *Section 2.6 -sf opt*
- Search the *pair style* page for styles followed by (t)
- *Benchmarks page* of website

## 6.2.66 ORIENT package

**Contents:**

A few fixes that apply orientation dependent forces for studying grain boundary migration.

**Supporting info:**

- src/ORIENT: filenames -> commands
  - *fix orient/bcc*
  - *fix orient/fcc*
  - *fix orient/eco*
- 

## 6.2.67 PERI package

**Contents:**

An atom style, several pair styles which implement different Peridynamics materials models, and several computes which calculate diagnostics. Peridynamics is a particle-based meshless continuum model.

**Authors:** The original package was created by Mike Parks (Sandia). Additional Peridynamics models were added by Rezwanur Rahman and John Foster (UTSA).

**Supporting info:**

- src/PERI: filenames -> commands
- doc/PDF/PDLammps\_overview.pdf
- doc/PDF/PDLammps\_EPS.pdf
- doc/PDF/PDLammps\_VES.pdf
- *atom\_style peri*
- *pair\_style peri/\**
- *compute damage/atom*

- *compute plasticity/atom*
  - *examples/peri*
  - <https://www.lammps.org/movies.html#peri>
- 

## 6.2.68 PHONON package

### Contents:

A *fix phonon* command that calculates dynamical matrices, which can then be used to compute phonon dispersion relations, directly from molecular dynamics simulations. And a *dynamical\_matrix* as well as a *third\_order* command to compute the dynamical matrix and third order tensor from finite differences.

### Install:

The PHONON package requires that also the **KSPACE** package is installed.

**Authors:** Ling-Ti Kong (Shanghai Jiao Tong University) for “fix phonon” and Charlie Sievers (UC Davis) for “dynamical\_matrix” and “third\_order”

### Supporting info:

- *src/PHONON*: filenames -> commands
  - *src/PHONON/README*
  - *fix phonon*
  - *dynamical\_matrix*
  - *third\_order*
  - *examples/PACKAGES/phonon*
- 

## 6.2.69 PLUGIN package

### Contents:

A *plugin* command that can load and unload several kind of styles in LAMMPS from shared object files at runtime without having to recompile and relink LAMMPS.

When the environment variable `LAMMPS_PLUGIN_PATH` is set, then LAMMPS will search the directory (or directories) listed in this path for files with names that end in `plugin.so` (e.g. `helloplugin.so`) and will try to load the contained plugins automatically at start-up.

**Authors:** Axel Kohlmeyer (Temple U)

### Supporting info:

- *src/PLUGIN*: filenames -> commands
  - *plugin command*
  - *Information on writing plugins*
  - *examples/plugin*
-

## 6.2.70 PLUMED package

### Contents:

The fix *plumed* command allows you to use the PLUMED free energy plugin for molecular dynamics to analyze and bias your LAMMPS trajectory on the fly. The PLUMED library is called from within the LAMMPS input script by using the *fix plumed* command.

**Authors:** The **PLUMED library** is written and maintained by Massimiliano Bonomi, Giovanni Bussi, Carlo Camilioni, and Gareth Tribello.

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/PLUMED/README
  - lib/plumed/README
  - *fix plumed*
  - examples/PACKAGES/plumed
- 

## 6.2.71 POEMS package

### Contents:

A fix that wraps the Parallelizable Open source Efficient Multibody Software (POEMS) library, which is able to simulate the dynamics of articulated body systems. These are systems with multiple rigid bodies (collections of particles) whose motion is coupled by connections at hinge points.

**Author:** Rudra Mukherjee (JPL) while at RPI.

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- src/POEMS: filenames -> commands
  - src/POEMS/README
  - lib/poems/README
  - *fix poems*
  - examples/rigid
-

### 6.2.72 PTM package

**Contents:**

A *compute ptm/atom* command that calculates local structure characterization using the Polyhedral Template Matching methodology.

**Author:** Peter Mahler Larsen (MIT).

**Supporting info:**

- src/PTM: filenames not starting with ptm\_ -> commands
  - src/PTM: filenames starting with ptm\_ -> supporting code
  - src/PTM/LICENSE
  - *compute ptm/atom*
- 

### 6.2.73 PYTHON package

**Contents:**

A *python* command which allow you to execute Python code from a LAMMPS input script. The code can be in a separate file or embedded in the input script itself. See the *Python call* page for an overview of using Python from LAMMPS in this manner and all the *Python* manual pages for other ways to use LAMMPS and Python together.

---

**Note:** Building with the PYTHON package assumes you have a Python development environment (headers and libraries) available on your system, which needs to be either Python version 2.7 or Python 3.5 and later.

---

**Install:**

This package has *specific installation instructions* on the *Build extras* page.

**Supporting info:**

- src/PYTHON: filenames -> commands
  - *Python call*
  - lib/python/README
  - examples/python
- 

### 6.2.74 QEQ package

**Contents:**

Several fixes for performing charge equilibration (QEq) via different algorithms. These can be used with pair styles that perform QEq as part of their formulation.

**Supporting info:**

- src/QEQ: filenames -> commands
  - *fix qeq\**
-

- [examples/qeq](#)
  - [examples/streitz](#)
- 

### 6.2.75 QMMM package

#### Contents:

A *fix qmmm* command which allows LAMMPS to be used as the MM code in a QM/MM simulation. This is currently only available in combination with the [Quantum ESPRESSO](#) package.

To use this package you must have Quantum ESPRESSO (QE) available on your system and include its coupling library in the compilation and then compile LAMMPS as a library. For QM/MM calculations you then build a custom binary with MPI support, that sets up 3 partitions with MPI sub-communicators (for inter- and intra-partition communication) and then calls the corresponding library interfaces on each partition (2x LAMMPS and 1x QE).

The current implementation supports an ONIOM style mechanical coupling and a multi-pole based electrostatic coupling to the Quantum ESPRESSO plane wave DFT package. The QM/MM interface has been written in a manner that coupling to other QM codes should be possible without changes to LAMMPS itself.

**Authors:** Axel Kohlmeyer (Temple U), Mariella Ippolito and Carlo Cavazzoni (CINECA, Italy)

#### Install:

This package has *specific installation instructions* on the [Build extras](#) page.

#### Supporting info:

- [src/QMMM: filenames -> commands](#)
  - [src/QMMM/README](#)
  - [lib/qmmm/README](#)
  - [fix phonon](#)
  - [lib/qmmm/example-ec/README](#)
  - [lib/qmmm/example-mc/README](#)
- 

### 6.2.76 QTB package

#### Contents:

Two fixes which provide a self-consistent quantum treatment of vibrational modes in a classical molecular dynamics simulation. By coupling the MD simulation to a colored thermostat, it introduces zero point energy into the system, altering the energy power spectrum and the heat capacity to account for their quantum nature. This is useful when modeling systems at temperatures lower than their classical limits or when temperatures ramp across the classical limits in a simulation.

**Author:** Yuan Shen (Stanford U).

#### Supporting info:

- [src/QTB: filenames -> commands](#)
  - [src/QTB/README](#)
  - [fix qtb](#)
-



- *fix qbmsst*
  - examples/PACKAGES/qtb
- 

### 6.2.77 REACTION package

#### Contents:

This package allows for complex bond topology changes (reactions) during a running MD simulation, when using classical force fields. Topology changes are defined in pre- and post-reaction molecule templates and can include creation and deletion of bonds, angles, dihedrals, impropers, atom types, bond types, angle types, dihedral types, improper types, and/or atomic charges. Other options currently available include reaction constraints (e.g. angle and Arrhenius constraints), deletion of reaction byproducts or other small molecules, and chiral-sensitive reactions.

**Author:** Jacob R. Gissinger (CU Boulder) while at NASA Langley Research Center.

#### Supporting info:

- src/REACTION: filenames -> commands
  - src/REACTION/README
  - *fix bond/react*
  - examples/PACKAGES/reaction
  - 2017 LAMMPS Workshop
  - 2019 LAMMPS Workshop
  - reactor.org
- 

### 6.2.78 REAXFF package

#### Contents:

A pair style which implements the ReaxFF potential in C/C++. ReaxFF is a universal reactive force field. See the src/REAXFF/README file for more info on differences between the two packages. Also two fixes for monitoring molecules as bonds are created and destroyed.

**Author:** Hasan Metin Aktulga (MSU) while at Purdue University.

#### Supporting info:

- src/REAXFF: filenames -> commands
  - src/REAXFF/README
  - *pair\_style reaxff*
  - *fix reaxff/bonds*
  - *fix reaxff/species*
  - examples/reaxff
-

## 6.2.79 REPLICA package

### Contents:

A collection of multi-replica methods which can be used when running multiple LAMMPS simulations (replicas). See the *Howto replica* page for an overview of how to run multi-replica simulations in LAMMPS. Methods in the package include nudged elastic band (NEB), parallel replica dynamics (PRD), temperature accelerated dynamics (TAD), parallel tempering, and a verlet/split algorithm for performing long-range Coulombics on one set of processors, and the remainder of the force field calculation on another set.

### Supporting info:

- src/REPLICA: filenames -> commands
  - *Howto replica*
  - *neb*
  - *prd*
  - *tad*
  - *temper*,
  - *temper/npt*,
  - *temper/grem*,
  - *run\_style verlet/split*
  - examples/neb
  - examples/prd
  - examples/tad
  - examples/PACKAGES/grem
- 

## 6.2.80 RIGID package

### Contents:

Fixes which enforce rigid constraints on collections of atoms or particles. This includes SHAKE and RATTLE, as well as various rigid-body integrators for a few large bodies or many small bodies. Also several computes which calculate properties of rigid bodies.

### Supporting info:

- src/RIGID: filenames -> commands
- *compute erotate/rigid*
- *fix shake*
- *fix rattle*
- *fix rigid/\**
- examples/ASPHERE
- examples/rigid
- bench/in.rhodo

- <https://www.lammps.org/movies.html#box>
  - <https://www.lammps.org/movies.html#star>
- 

### 6.2.81 SCAFACOS package

#### Contents:

A KSpace style which wraps the ScaFaCoS Coulomb solver library to compute long-range Coulombic interactions.

To use this package you must have the ScaFaCoS library available on your system.

**Author:** Rene Halver (JSC) wrote the scafacos LAMMPS command.

ScaFaCoS itself was developed by a consortium of German research facilities with a BMBF (German Ministry of Science and Education) funded project in 2009-2012. Participants of the consortium were the Universities of Bonn, Chemnitz, Stuttgart, and Wuppertal as well as the Forschungszentrum Juelich.

#### Install:

This package has *specific installation instructions* on the *Build extras* page. The SCAFACOS package requires that LAMMPS is build in *MPI parallel mode*.

#### Supporting info:

- src/SCAFACOS: filenames -> commands
  - src/SCAFACOS/README
  - *kpace\_style scafacos*
  - *kpace\_modify*
  - examples/PACKAGES/sfacos
- 

### 6.2.82 SHOCK package

#### Contents:

Fixes for running impact simulations where a shock-wave passes through a material.

#### Supporting info:

- src/SHOCK: filenames -> commands
  - *fix append/atoms*
  - *fix msst*
  - *fix nphug*
  - *fix wall/piston*
  - examples/hugonostat
  - examples/msst
-

### 6.2.83 SMTBQ package

**Contents:**

Pair styles which implement Second Moment Tight Binding models. One with QEq charge equilibration (SMTBQ) for the description of ionocovalent bonds in oxides, and two more as plain SMATB models.

**Authors:** SMTBQ: Nicolas Salles, Emile Maras, Olivier Politano, and Robert Tetot (LAAS-CNRS, France); SMATB: Daniele Rapetti (Politecnico di Torino)

**Supporting info:**

- src/SMTBQ: filenames -> commands
  - src/SMTBQ/README
  - *pair\_style smtbq*
  - *pair\_style smatb*, *pair\_style smatb/single*
  - examples/PACKAGES/smtbq
- 

### 6.2.84 SPH package

**Contents:**

An atom style, fixes, computes, and several pair styles which implements smoothed particle hydrodynamics (SPH) for liquids. See the related *MACHDYN package* for smooth Mach dynamics (SMD) for solids.

This package contains ideal gas, Lennard-Jones equation of states, Tait, and full support for complete (i.e. internal-energy dependent) equations of state. It allows for plain or Monaghans XSPH integration of the equations of motion. It has options for density continuity or density summation to propagate the density field. It has *set* command options to set the internal energy and density of particles from the input script and allows the same quantities to be output with thermodynamic output or to dump files via the *compute property/atom* command.

**Author:** Georg Ganzenmuller (Fraunhofer-Institute for High-Speed Dynamics, Ernst Mach Institute, Germany).

**Supporting info:**

- src/SPH: filenames -> commands
  - src/SPH/README
  - doc/PDF/SPH\_LAMMPS\_userguide.pdf
  - examples/PACKAGES/sph
  - <https://www.lammps.org/movies.html#sph>
-

## 6.2.85 SPIN package

### Contents:

Model atomic magnetic spins classically, coupled to atoms moving in the usual manner via MD. Various pair, fix, and compute styles.

**Author:** Julien Tranchida (Sandia).

### Supporting info:

- src/SPIN: filenames -> commands
  - *Howto spins*
  - *pair\_style spin/dipole/cut*
  - *pair\_style spin/dipole/long*
  - *pair\_style spin/dmi*
  - *pair\_style spin/exchange*
  - *pair\_style spin/exchange/biquadratic*
  - *pair\_style spin/magelec*
  - *pair\_style spin/neel*
  - *fix nve/spin*
  - *fix langevin/spin*
  - *fix precession/spin*
  - *compute spin*
  - *neb/spin*
  - examples/SPIN
- 

## 6.2.86 SRD package

### Contents:

A pair of fixes which implement the Stochastic Rotation Dynamics (SRD) method for coarse-graining of a solvent, typically around large colloidal particles.

### Supporting info:

- src/SRD: filenames -> commands
- *fix srd*
- *fix wall/srd*
- examples/srd
- examples/ASPHERE
- <https://www.lammps.org/movies.html#tri>
- <https://www.lammps.org/movies.html#line>
- <https://www.lammps.org/movies.html#poly>

## 6.2.87 TALLY package

### Contents:

Several compute styles that can be called when pairwise interactions are calculated to tally information (forces, heat flux, energy, stress, etc) about individual interactions.

**Author:** Axel Kohlmeyer (Temple U).

### Supporting info:

- src/TALLY: filenames -> commands
  - src/TALLY/README
  - *compute \*/tally*
  - examples/PACKAGES/tally
- 

## 6.2.88 UEF package

### Contents:

A fix style for the integration of the equations of motion under extensional flow with proper boundary conditions, as well as several supporting compute styles and an output option.

**Author:** David Nicholson (MIT).

### Supporting info:

- src/UEF: filenames -> commands
  - src/UEF/README
  - *fix nvt/uef*
  - *fix npt/uef*
  - *compute pressure/uef*
  - *compute temp/uef*
  - *dump cfg/uef*
  - examples/uef
-

## 6.2.89 VORONOI package

### Contents:

A `compute` command which calculates the Voronoi tessellation of a collection of atoms by wrapping the [Voro++ library](#). This can be used to calculate the local volume of each atom or its near neighbors.

To use this package you must have the Voro++ library available on your system.

**Author:** Daniel Schwen (INL) while at LANL. The open-source Voro++ library was written by Chris Rycroft (Harvard U) while at UC Berkeley and LBNL.

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- `src/VORONOI`: filenames -> commands
  - `src/VORONOI/README`
  - `lib/voronoi/README`
  - `compute voronoi/atom`
  - `examples/voronoi`
- 

## 6.2.90 VTK package

### Contents:

A `dump vtk` command which outputs snapshot info in the [VTK format](#), enabling visualization by [Paraview](#) or other visualization packages.

To use this package you must have VTK library available on your system.

**Authors:** Richard Berger (JKU) and Daniel Queteschiner (DCS Computing).

### Install:

This package has *specific installation instructions* on the *Build extras* page.

### Supporting info:

- `src/VTK`: filenames -> commands
  - `src/VTK/README`
  - `lib/vtk/README`
  - `dump vtk`
-

## 6.2.91 YAFF package

### Contents:

Some potentials that are also implemented in the Yet Another Force Field (YAFF) code. The expressions and their use are discussed in the following papers

- Vanduyfhuys et al., J. Comput. Chem., 36 (13), 1015-1027 (2015) [link](#)
- Vanduyfhuys et al., J. Comput. Chem., 39 (16), 999-1011 (2018) [link](#)

which discuss the QuickFF methodology.

**Author:** Steven Vandenbrande.

### Supporting info:

- src/YAFF/README
- *angle\_style cross*
- *angle\_style mm3*
- *bond\_style mm3*
- *improper\_style distharm*
- *improper\_style sqdistharm*
- *pair\_style mm3/switch3/coulgauss/long*
- *pair\_style lj/switch3/coulgauss/long*
- examples/PACKAGES/yaff



## ACCELERATE PERFORMANCE

This section describes various methods for improving LAMMPS performance for different classes of problems running on different kinds of machines.

There are two thrusts to the discussion that follows. The first is using code options that implement alternate algorithms that can speed-up a simulation. The second is to use one of the several accelerator packages provided with LAMMPS that contain code optimized for certain kinds of hardware, including multi-core CPUs, GPUs, and Intel Xeon Phi co-processors.

The [Benchmark page](#) of the LAMMPS website gives performance results for the various accelerator packages discussed on the [Accelerator packages](#) page, for several of the standard LAMMPS benchmark problems, as a function of problem size and number of compute nodes, on different hardware platforms.

### 7.1 Benchmarks

Current LAMMPS performance is discussed on the [Benchmarks page](#) of the [LAMMPS website](#) where timings and parallel efficiency are listed. The page has several sections, which are briefly described below:

- CPU performance on 5 standard problems, strong and weak scaling
- GPU and Xeon Phi performance on same and related problems
- Comparison of cost of interatomic potentials
- Performance of huge, billion-atom problems

The 5 standard problems are as follow:

1. LJ = atomic fluid, Lennard-Jones potential with 2.5 sigma cutoff (55 neighbors per atom), NVE integration
2. Chain = bead-spring polymer melt of 100-mer chains, FENE bonds and LJ pairwise interactions with a  $2^{1/6}$  sigma cutoff (5 neighbors per atom), NVE integration
3. EAM = metallic solid, Cu EAM potential with 4.95 Angstrom cutoff (45 neighbors per atom), NVE integration
4. Chute = granular chute flow, frictional history potential with 1.1 sigma cutoff (7 neighbors per atom), NVE integration
5. Rhodo = rhodopsin protein in solvated lipid bilayer, CHARMM force field with a 10 Angstrom LJ cutoff (440 neighbors per atom), particle-particle particle-mesh (PPPM) for long-range Coulombics, NPT integration

Input files for these 5 problems are provided in the bench directory of the LAMMPS distribution. Each has 32,000 atoms and runs for 100 timesteps. The size of the problem (number of atoms) can be varied using command-line switches as described in the bench/README file. This is an easy way to test performance and either strong or weak scalability on your machine.

The bench directory includes a few log.\* files that show performance of these 5 problems on 1 or 4 cores of Linux desktop. The bench/FERMI and bench/KEPLER directories have input files and scripts and instructions for running the same (or similar) problems using OpenMP or GPU or Xeon Phi acceleration options. See the README files in those directories and the *Accelerator packages* pages for instructions on how to build LAMMPS and run on that kind of hardware.

The bench/POTENTIALS directory has input files which correspond to the table of results on the [Potentials](#) section of the Benchmarks web page. So you can also run those test problems on your machine.

The [billion-atom](#) section of the Benchmarks web page has performance data for very large benchmark runs of simple Lennard-Jones (LJ) models, which use the bench/in.lj input script.

---

For all the benchmarks, a useful metric is the CPU cost per atom per timestep. Since performance scales roughly linearly with problem size and timesteps for all LAMMPS models (i.e. interatomic or coarse-grained potentials), the run time of any problem using the same model (atom style, force field, cutoff, etc) can then be estimated.

Performance on a parallel machine can also be predicted from one-core or one-node timings if the parallel efficiency can be estimated. The communication bandwidth and latency of a particular parallel machine affects the efficiency. On most machines LAMMPS will give a parallel efficiency on these benchmarks above 50% so long as the number of atoms/core is a few 100 or greater, and closer to 100% for large numbers of atoms/core. This is for all-MPI mode with one MPI task per core. For nodes with accelerator options or hardware (OpenMP, GPU, Phi), you should first measure single node performance. Then you can estimate parallel performance for multi-node runs using the same logic as for all-MPI mode, except that now you will typically need many more atoms/node to achieve good scalability.

## 7.2 Measuring performance

Before trying to make your simulation run faster, you should understand how it currently performs and where the bottlenecks are.

The best way to do this is run the your system (actual number of atoms) for a modest number of timesteps (say 100 steps) on several different processor counts, including a single processor if possible. Do this for an equilibrium version of your system, so that the 100-step timings are representative of a much longer run. There is typically no need to run for 1000s of timesteps to get accurate timings; you can simply extrapolate from short runs.

For the set of runs, look at the timing data printed to the screen and log file at the end of each LAMMPS run. The [screen and logfile output](#) page gives an overview.

Running on one (or a few processors) should give a good estimate of the serial performance and what portions of the timestep are taking the most time. Running the same problem on a few different processor counts should give an estimate of parallel scalability. I.e. if the simulation runs 16x faster on 16 processors, its 100% parallel efficient; if it runs 8x faster on 16 processors, it's 50% efficient.

The most important data to look at in the timing info is the timing breakdown and relative percentages. For example, trying different options for speeding up the long-range solvers will have little impact if they only consume 10% of the run time. If the pairwise time is dominating, you may want to look at GPU or OMP versions of the pair style, as discussed below. Comparing how the percentages change as you increase the processor count gives you a sense of how different operations within the timestep are scaling. Note that if you are running with a Kspace solver, there is additional output on the breakdown of the Kspace time. For PPPM, this includes the fraction spent on FFTs, which can be communication intensive.

Another important detail in the timing info are the histograms of atoms counts and neighbor counts. If these vary widely across processors, you have a load-imbalance issue. This often results in inaccurate relative timing data, because processors have to wait when communication occurs for other processors to catch up. Thus the reported times for “Communication” or “Other” may be higher than they really are, due to load-imbalance. If this is an issue, you can use the [timer sync](#) command to obtain synchronized timings.

## 7.3 General tips

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**Note:** this page is still a work in progress

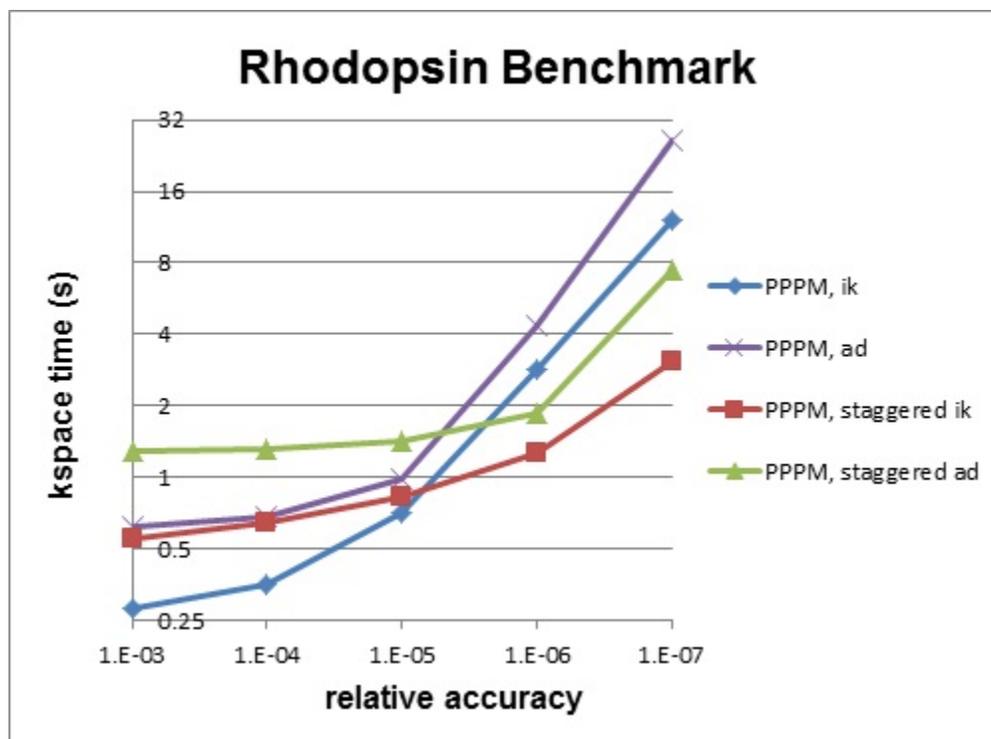
---

Here is a list of general ideas for improving simulation performance. Most of them are only applicable to certain models and certain bottlenecks in the current performance, so let the timing data you generate be your guide. It is hard, if not impossible, to predict how much difference these options will make, since it is a function of problem size, number of processors used, and your machine. There is no substitute for identifying performance bottlenecks, and trying out various options.

- rRESPA
- Two-FFT PPPM
- Staggered PPPM
- single vs double PPPM
- partial charge PPPM
- verlet/split run style
- processor command for proc layout and numa layout
- load-balancing: balance and fix balance

Two-FFT PPPM, also called *analytic differentiation* or *ad* PPPM, uses 2 FFTs instead of the 4 FFTs used by the default *ik differentiation* PPPM. However, 2-FFT PPPM also requires a slightly larger mesh size to achieve the same accuracy as 4-FFT PPPM. For problems where the FFT cost is the performance bottleneck (typically large problems running on many processors), 2-FFT PPPM may be faster than 4-FFT PPPM.

Staggered PPPM performs calculations using two different meshes, one shifted slightly with respect to the other. This can reduce force aliasing errors and increase the accuracy of the method, but also doubles the amount of work required. For high relative accuracy, using staggered PPPM allows one to half the mesh size in each dimension as compared to regular PPPM, which can give around a 4x speedup in the kspace time. However, for low relative accuracy, using staggered PPPM gives little benefit and can be up to 2x slower in the kspace time. For example, the rhodopsin benchmark was run on a single processor, and results for kspace time vs. relative accuracy for the different methods are shown in the figure below. For this system, staggered PPPM (using *ik differentiation*) becomes useful when using a relative accuracy of slightly greater than  $1e-5$  and above.



**Note:** Using staggered PPPM may not give the same increase in accuracy of energy and pressure as it does in forces, so some caution must be used if energy and/or pressure are quantities of interest, such as when using a barostat.

## 7.4 Accelerator packages

Accelerated versions of various *pair\_style*, *fixes*, *computes*, and other commands have been added to LAMMPS, which will typically run faster than the standard non-accelerated versions. Some require appropriate hardware to be present on your system, e.g. GPUs or Intel Xeon Phi co-processors.

All of these commands are in packages provided with LAMMPS. An overview of packages is give on the *Packages* doc pages.

These are the accelerator packages currently in LAMMPS:

<i>GPU Package</i>	for GPUs via CUDA, OpenCL, or ROCm HIP
<i>INTEL Package</i>	for Intel CPUs and Intel Xeon Phi
<i>KOKKOS Package</i>	for NVIDIA GPUs, Intel Xeon Phi, and OpenMP threading
<i>OPENMP Package</i>	for OpenMP threading and generic CPU optimizations
<i>OPT Package</i>	generic CPU optimizations

### 7.4.1 GPU package

The GPU package was developed by Mike Brown while at SNL and ORNL (now at Intel Corp.) and his collaborators, particularly Trung Nguyen (now at Northwestern). Support for AMD GPUs via HIP was added by Vsevolod Nikolskiy and coworkers at HSE University.

The GPU package provides GPU versions of many pair styles and for parts of the *k-space\_style pppm* for long-range Coulombics. It has the following general features:

- It is designed to exploit common GPU hardware configurations where one or more GPUs are coupled to many cores of one or more multi-core CPUs, e.g. within a node of a parallel machine.
- Atom-based data (e.g. coordinates, forces) are moved back-and-forth between the CPU(s) and GPU every timestep.
- Neighbor lists can be built on the CPU or on the GPU
- The charge assignment and force interpolation portions of PPPM can be run on the GPU. The FFT portion, which requires MPI communication between processors, runs on the CPU.
- Force computations of different style (pair vs. bond/angle/dihedral/improper) can be performed concurrently on the GPU and CPU(s), respectively.
- It allows for GPU computations to be performed in single or double precision, or in mixed-mode precision, where pairwise forces are computed in single precision, but accumulated into double-precision force vectors.
- LAMMPS-specific code is in the GPU package. It makes calls to a generic GPU library in the `lib/gpu` directory. This library provides either Nvidia support, AMD support, or more general OpenCL support (for Nvidia GPUs, AMD GPUs, Intel GPUs, and multi-core CPUs). so that the same functionality is supported on a variety of hardware.

#### Required hardware/software:

To compile and use this package in CUDA mode, you currently need to have an NVIDIA GPU and install the corresponding NVIDIA CUDA toolkit software on your system (this is only tested on Linux and unsupported on Windows):

- Check if you have an NVIDIA GPU: `cat /proc/driver/nvidia/gpus/*/information`
- Go to [http://www.nvidia.com/object/cuda\\_get.html](http://www.nvidia.com/object/cuda_get.html)
- Install a driver and toolkit appropriate for your system (SDK is not necessary)
- Run `lammmps/lib/gpu/nvc_get_devices` (after building the GPU library, see below) to list supported devices and properties

To compile and use this package in OpenCL mode, you currently need to have the OpenCL headers and the (vendor neutral) OpenCL library installed. In OpenCL mode, the acceleration depends on having an [OpenCL Installable Client Driver \(ICD\)](#) installed. There can be multiple of them for the same or different hardware (GPUs, CPUs, Accelerators) installed at the same time. OpenCL refers to those as ‘platforms’. The GPU library will try to auto-select the best suitable platform, but this can be overridden using the `platform` option of the `package` command. run `lammmps/lib/gpu/ocl_get_devices` to get a list of available platforms and devices with a suitable ICD available.

To compile and use this package for Intel GPUs, OpenCL or the Intel oneAPI HPC Toolkit can be installed using linux package managers. The latter also provides optimized C++, MPI, and many other libraries and tools. See:

- <https://software.intel.com/content/www/us/en/develop/tools/oneapi/hpc-toolkit/download.html>

If you do not have a discrete GPU card installed, this package can still provide significant speedups on some CPUs that include integrated GPUs. Additionally, for many macs, OpenCL is already included with the OS and Makefiles are available in the `lib/gpu` directory.

To compile and use this package in HIP mode, you have to have the AMD ROCm software installed. Versions of ROCm older than 3.5 are currently deprecated by AMD.

**Building LAMMPS with the GPU package:**

See the *Build extras* page for instructions.

**Run with the GPU package from the command line:**

The `mpirun` or `mpiexec` command sets the total number of MPI tasks used by LAMMPS (one or multiple per compute node) and the number of MPI tasks used per node. E.g. the `mpirun` command in MPICH does this via its `-np` and `-ppn` switches. Ditto for OpenMPI via `-np` and `-npernode`.

When using the GPU package, you cannot assign more than one GPU to a single MPI task. However multiple MPI tasks can share the same GPU, and in many cases it will be more efficient to run this way. Likewise it may be more efficient to use less MPI tasks/node than the available # of CPU cores. Assignment of multiple MPI tasks to a GPU will happen automatically if you create more MPI tasks/node than there are GPUs/node. E.g. with 8 MPI tasks/node and 2 GPUs, each GPU will be shared by 4 MPI tasks.

The GPU package also has limited support for OpenMP for both multi-threading and vectorization of routines that are run on the CPUs. This requires that the GPU library and LAMMPS are built with flags to enable OpenMP support (e.g. `-fopenmp`). Some styles for time integration are also available in the GPU package. These run completely on the CPUs in full double precision, but exploit multi-threading and vectorization for faster performance.

Use the “`-sf gpu`” *command-line switch*, which will automatically append “`gpu`” to styles that support it. Use the “`-pk gpu Ng`” *command-line switch* to set `Ng` = # of GPUs/node to use. If `Ng` is 0, the number is selected automatically as the number of matching GPUs that have the highest number of compute cores.

```

mpirun -sf gpu -pk gpu 1 -in in.script           # 1 MPI task uses 1 GPU
mpirun -np 12 lmp_machine -sf gpu -pk gpu 2 -in in.script # 12 MPI tasks share 2 GPUs on a single 16-core (or whatever) node
mpirun -np 48 -ppn 12 lmp_machine -sf gpu -pk gpu 2 -in in.script # ditto on 4 16-core nodes

```

Note that if the “`-sf gpu`” switch is used, it also issues a default `package gpu 0` command, which will result in automatic selection of the number of GPUs to use.

Using the “`-pk`” switch explicitly allows for setting of the number of GPUs/node to use and additional options. Its syntax is the same as the “`package gpu`” command. See the *package* command page for details, including the default values used for all its options if it is not specified.

Note that the default for the `package gpu` command is to set the Newton flag to “off” pairwise interactions. It does not affect the setting for bonded interactions (LAMMPS default is “on”). The “off” setting for pairwise interaction is currently required for GPU package pair styles.

**Or run with the GPU package by editing an input script:**

The discussion above for the `mpirun/mpiexec` command, MPI tasks/node, and use of multiple MPI tasks/GPU is the same.

Use the *suffix gpu* command, or you can explicitly add an “`gpu`” suffix to individual styles in your input script, e.g.

```
pair_style lj/cut/gpu 2.5
```

You must also use the `package gpu` command to enable the GPU package, unless the “`-sf gpu`” or “`-pk gpu`” *command-line switches* were used. It specifies the number of GPUs/node to use, as well as other options.

**Speed-ups to expect:**

The performance of a GPU versus a multi-core CPU is a function of your hardware, which pair style is used, the number of atoms/GPU, and the precision used on the GPU (double, single, mixed). Using the GPU package in OpenCL mode on CPUs (which uses vectorization and multithreading) is usually resulting in inferior performance compared to using LAMMPS’ native threading and vectorization support in the OPENMP and INTEL packages.

See the [Benchmark page](#) of the LAMMPS website for performance of the GPU package on various hardware, including the Titan HPC platform at ORNL.

You should also experiment with how many MPI tasks per GPU to use to give the best performance for your problem and machine. This is also a function of the problem size and the pair style being using. Likewise, you should experiment with the precision setting for the GPU library to see if single or mixed precision will give accurate results, since they will typically be faster.

MPI parallelism typically outperforms OpenMP parallelism, but in some cases using fewer MPI tasks and multiple OpenMP threads with the GPU package can give better performance. 3-body potentials can often perform better with multiple OMP threads because the inter-process communication is higher for these styles with the GPU package in order to allow deterministic results.

#### **Guidelines for best performance:**

- Using multiple MPI tasks per GPU will often give the best performance, as allowed by most multi-core CPU/GPU configurations.
- If the number of particles per MPI task is small (e.g. 100s of particles), it can be more efficient to run with fewer MPI tasks per GPU, even if you do not use all the cores on the compute node.
- The *package gpu* command has several options for tuning performance. Neighbor lists can be built on the GPU or CPU. Force calculations can be dynamically balanced across the CPU cores and GPUs. GPU-specific settings can be made which can be optimized for different hardware. See the *package* command page for details.
- As described by the *package gpu* command, GPU accelerated pair styles can perform computations asynchronously with CPU computations. The “Pair” time reported by LAMMPS will be the maximum of the time required to complete the CPU pair style computations and the time required to complete the GPU pair style computations. Any time spent for GPU-enabled pair styles for computations that run simultaneously with *bond*, *angle*, *dihedral*, *improper*, and *long-range* calculations will not be included in the “Pair” time.
- Since only part of the *pppm kspace* style is GPU accelerated, it may be faster to only use GPU acceleration for Pair styles with long-range electrostatics. See the “pair/only” keyword of the *package* command for a shortcut to do that. The work between *kspace* on the CPU and non-bonded interactions on the GPU can be balanced through adjusting the coulomb cutoff without loss of accuracy.
- When the *mode* setting for the *package gpu* command is *force/neighbor*, the time for neighbor list calculations on the GPU will be added into the “Pair” time, not the “Neighbor” time. An additional breakdown of the times required for various tasks on the GPU (data copy, neighbor calculations, force computations, etc) are output only with the LAMMPS screen output (not in the log file) at the end of each run. These timings represent total time spent on the GPU for each routine, regardless of asynchronous CPU calculations.
- The output section “GPU Time Info (average)” reports “Max Mem / Proc”. This is the maximum memory used at one time on the GPU for data storage by a single MPI process.

## Restrictions

None.

### 7.4.2 INTEL package

The INTEL package is maintained by Mike Brown at Intel Corporation. It provides two methods for accelerating simulations, depending on the hardware you have. The first is acceleration on Intel CPUs by running in single, mixed, or double precision with vectorization. The second is acceleration on Intel Xeon Phi co-processors via offloading neighbor list and non-bonded force calculations to the Phi. The same C++ code is used in both cases. When offloading to a co-processor from a CPU, the same routine is run twice, once on the CPU and once with an offload flag. This allows LAMMPS to run on the CPU cores and co-processor cores simultaneously.

#### Currently Available INTEL Styles

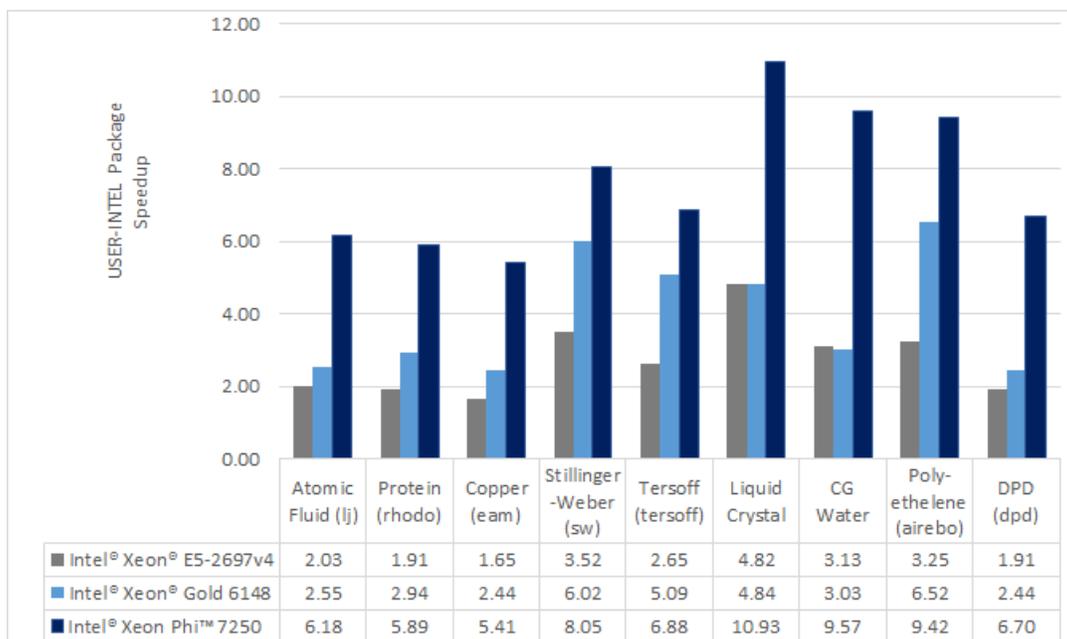
- Angle Styles: charmm, harmonic
- Bond Styles: fene, fourier, harmonic
- Dihedral Styles: charmm, fourier, harmonic, opl
- Fixes: nve, npt, nvt, nvt/sllod, nve/asphere
- Improper Styles: cvff, harmonic
- Pair Styles: airebo, airebo/morse, buck/coul/cut, buck/coul/long, buck, dpd, eam, eam/alloy, eam/fs, gayberne, lj/charmm/coul/charmm, lj/charmm/coul/long, lj/cut, lj/cut/coul/long, lj/long/coul/long, rebo, sw, tersoff
- K-Space Styles: pppm, pppm/disp

<p><b>Warning:</b> None of the styles in the INTEL package currently support computing per-atom stress. If any compute or fix in your input requires it, LAMMPS will abort with an error message.</p>
---

#### Speed-up to expect

The speedup will depend on your simulation, the hardware, which styles are used, the number of atoms, and the floating-point precision mode. Performance improvements are shown compared to LAMMPS *without using other acceleration packages* as these are under active development (and subject to performance changes). The measurements were performed using the input files available in the src/INTEL/TEST directory with the provided run script. These are scalable in size; the results given are with 512K particles (524K for Liquid Crystal). Most of the simulations are standard LAMMPS benchmarks (indicated by the filename extension in parenthesis) with modifications to the run length and to add a warm-up run (for use with offload benchmarks).





Results are speedups obtained on Intel Xeon E5-2697v4 processors (code-named Broadwell), Intel Xeon Phi 7250 processors (code-named Knights Landing), and Intel Xeon Gold 6148 processors (code-named Skylake) with “June 2017” LAMMPS built with Intel Parallel Studio 2017 update 2. Results are with 1 MPI task per physical core. See [src/INTEL/TEST/README](#) for the raw simulation rates and instructions to reproduce.

## Accuracy and order of operations

In most molecular dynamics software, parallelization parameters (# of MPI, OpenMP, and vectorization) can change the results due to changing the order of operations with finite-precision calculations. The INTEL package is deterministic. This means that the results should be reproducible from run to run with the *same* parallel configurations and when using deterministic libraries or library settings (MPI, OpenMP, FFT). However, there are differences in the INTEL package that can change the order of operations compared to LAMMPS without acceleration:

- Neighbor lists can be created in a different order
- Bins used for sorting atoms can be oriented differently
- The default stencil order for PPPM is 7. By default, LAMMPS will calculate other PPPM parameters to fit the desired accuracy with this order
- The *newton* setting applies to all atoms, not just atoms shared between MPI tasks
- Vectorization can change the order for adding pairwise forces
- When using the `-DLMP_USE_MKL_RNG` define (all included intel optimized makefiles do) at build time, the random number generator for dissipative particle dynamics (pair style dpd/intel) uses the Mersenne Twister generator included in the Intel MKL library (that should be more robust than the default Masaglia random number generator)

The precision mode (described below) used with the INTEL package can change the *accuracy* of the calculations. For the default *mixed* precision option, calculations between pairs or triplets of atoms are performed in single precision, intended to be within the inherent error of MD simulations. All accumulation is performed in double precision to prevent the error from growing with the number of atoms in the simulation. *Single* precision mode should not be used without appropriate validation.

## Quick Start for Experienced Users

LAMMPS should be built with the INTEL package installed. Simulations should be run with 1 MPI task per physical core, not *hardware thread*.

- Edit src/MAKE/OPTIONS/Makefile.intel\_cpu\_intelmpi as necessary.
- Set the environment variable KMP\_BLOCKTIME=0
- “-pk intel 0 omp \$t -sf intel” added to LAMMPS command-line
- \$t should be 2 for Intel Xeon CPUs and 2 or 4 for Intel Xeon Phi
- For some of the simple 2-body potentials without long-range electrostatics, performance and scalability can be better with the “newton off” setting added to the input script
- For simulations on higher node counts, add “processors \* \* \* grid numa” to the beginning of the input script for better scalability
- If using *kpace\_style pppm* in the input script, add “kpace\_modify diff ad” for better performance

For Intel Xeon Phi CPUs:

- Runs should be performed using MCDRAM.

For simulations using *kpace\_style pppm* on Intel CPUs supporting AVX-512:

- Add “kpace\_modify diff ad” to the input script
- The command-line option should be changed to “-pk intel 0 omp \$r lrt yes -sf intel” where \$r is the number of threads minus 1.
- Do not use thread affinity (set KMP\_AFFINITY=none)
- The “newton off” setting may provide better scalability

For Intel Xeon Phi co-processors (Offload):

- Edit src/MAKE/OPTIONS/Makefile.intel\_co-processor as necessary
- “-pk intel N omp 1” added to command-line where N is the number of co-processors per node.

---

## Required hardware/software

When using Intel compilers version 16.0 or later is required.

In order to use offload to co-processors, an Intel Xeon Phi co-processor and an Intel compiler are required.

Although any compiler can be used with the INTEL package, currently, vectorization directives are disabled by default when not using Intel compilers due to lack of standard support and observations of decreased performance. The OpenMP standard now supports directives for vectorization and we plan to transition the code to this standard once it is available in most compilers. We expect this to allow improved performance and support with other compilers.

For Intel Xeon Phi x200 series processors (code-named Knights Landing), there are multiple configuration options for the hardware. For best performance, we recommend that the MCDRAM is configured in “Flat” mode and with the cluster mode set to “Quadrant” or “SNC4”. “Cache” mode can also be used, although the performance might be slightly lower.

## Notes about Simultaneous Multithreading

Modern CPUs often support Simultaneous Multithreading (SMT). On Intel processors, this is called Hyper-Threading (HT) technology. SMT is hardware support for running multiple threads efficiently on a single core. *Hardware threads* or *logical cores* are often used to refer to the number of threads that are supported in hardware. For example, the Intel Xeon E5-2697v4 processor is described as having 36 cores and 72 threads. This means that 36 MPI processes or OpenMP threads can run simultaneously on separate cores, but that up to 72 MPI processes or OpenMP threads can be running on the CPU without costly operating system context switches.

Molecular dynamics simulations will often run faster when making use of SMT. If a thread becomes stalled, for example because it is waiting on data that has not yet arrived from memory, another thread can start running so that the CPU pipeline is still being used efficiently. Although benefits can be seen by launching a MPI task for every hardware thread, for multinode simulations, we recommend that OpenMP threads are used for SMT instead, either with the INTEL package, *OPENMP package*, or *KOKKOS package*. In the example above, up to 36X speedups can be observed by using all 36 physical cores with LAMMPS. By using all 72 hardware threads, an additional 10-30% performance gain can be achieved.

The BIOS on many platforms allows SMT to be disabled, however, we do not recommend this on modern processors as there is little to no benefit for any software package in most cases. The operating system will report every hardware thread as a separate core allowing one to determine the number of hardware threads available. On Linux systems, this information can normally be obtained with:

```
cat /proc/cpuinfo
```

## Building LAMMPS with the INTEL package

See the *Build extras* page for instructions. Some additional details are covered here.

For building with make, several example Makefiles for building with the Intel compiler are included with LAMMPS in the src/MAKE/OPTIONS/ directory:

```
Makefile.intel_cpu_intelmpi # Intel Compiler, Intel MPI, No Offload
Makefile.knl                # Intel Compiler, Intel MPI, No Offload
Makefile.intel_cpu_mpich   # Intel Compiler, MPICH, No Offload
Makefile.intel_cpu_openmpi # Intel Compiler, OpenMPI, No Offload
Makefile.intel_co-processor # Intel Compiler, Intel MPI, Offload
```

Makefile.knl is identical to Makefile.intel\_cpu\_intelmpi except that it explicitly specifies that vectorization should be for Intel Xeon Phi x200 processors making it easier to cross-compile. For users with recent installations of Intel Parallel Studio, the process can be as simple as:

```
make yes-intel
source /opt/intel/parallel_studio_xe_2016.3.067/psxevars.sh
# or psxevars.csh for C-shell
make intel_cpu_intelmpi
```

Note that if you build with support for a Phi co-processor, the same binary can be used on nodes with or without co-processors installed. However, if you do not have co-processors on your system, building without offload support will produce a smaller binary.

The general requirements for Makefiles with the INTEL package are as follows. When using Intel compilers, “-restrict” is required and “-qopenmp” is highly recommended for CCFLAGS and LINKFLAGS. CCFLAGS should include “-DLMP\_INTEL\_USELRT” (unless POSIX Threads are not supported in the build environment) and “-DLMP\_USE\_MKL\_RNG” (unless Intel Math Kernel Library (MKL) is not available in the build environment). For Intel compilers, LIB should include “-ltbbmalloc” or if the library is not available, “-DLMP\_INTEL\_NO\_TBB” can be

added to CCFLAGS. For builds supporting offload, “-DLMP\_INTEL\_OFFLOAD” is required for CCFLAGS and “-qoffload” is required for LINKFLAGS. Other recommended CCFLAG options for best performance are “-O2 -fno-alias -ansi-alias -qoverride-limits fp-model fast=2 -no-prec-div”.

---

**Note:** See the src/INTEL/README file for additional flags that might be needed for best performance on Intel server processors code-named “Skylake”.

---

---

**Note:** The vectorization and math capabilities can differ depending on the CPU. For Intel compilers, the “-x” flag specifies the type of processor for which to optimize. “-xHost” specifies that the compiler should build for the processor used for compiling. For Intel Xeon Phi x200 series processors, this option is “-xMIC-AVX512”. For fourth generation Intel Xeon (v4/Broadwell) processors, “-xCORE-AVX2” should be used. For older Intel Xeon processors, “-xAVX” will perform best in general for the different simulations in LAMMPS. The default in most of the example Makefiles is to use “-xHost”, however this should not be used when cross-compiling.

---

### Running LAMMPS with the INTEL package

Running LAMMPS with the INTEL package is similar to normal use with the exceptions that one should 1) specify that LAMMPS should use the INTEL package, 2) specify the number of OpenMP threads, and 3) optionally specify the specific LAMMPS styles that should use the INTEL package. 1) and 2) can be performed from the command-line or by editing the input script. 3) requires editing the input script. Advanced performance tuning options are also described below to get the best performance.

When running on a single node (including runs using offload to a co-processor), best performance is normally obtained by using 1 MPI task per physical core and additional OpenMP threads with SMT. For Intel Xeon processors, 2 OpenMP threads should be used for SMT. For Intel Xeon Phi CPUs, 2 or 4 OpenMP threads should be used (best choice depends on the simulation). In cases where the user specifies that LRT mode is used (described below), 1 or 3 OpenMP threads should be used. For multi-node runs, using 1 MPI task per physical core will often perform best, however, depending on the machine and scale, users might get better performance by decreasing the number of MPI tasks and using more OpenMP threads. For performance, the product of the number of MPI tasks and OpenMP threads should not exceed the number of available hardware threads in almost all cases.

---

**Note:** Setting core affinity is often used to pin MPI tasks and OpenMP threads to a core or group of cores so that memory access can be uniform. Unless disabled at build time, affinity for MPI tasks and OpenMP threads on the host (CPU) will be set by default on the host *when using offload to a co-processor*. In this case, it is unnecessary to use other methods to control affinity (e.g. taskset, numactl, I\_MPI\_PIN\_DOMAIN, etc.). This can be disabled with the *no\_affinity* option to the *package intel* command or by disabling the option at build time (by adding -DINTEL\_OFFLOAD\_NOAFFINITY to the CCFLAGS line of your Makefile). Disabling this option is not recommended, especially when running on a machine with Intel Hyper-Threading technology disabled.

---

## Run with the INTEL package from the command line

To enable INTEL optimizations for all available styles used in the input script, the “-sf intel” *command-line switch* can be used without any requirement for editing the input script. This switch will automatically append “intel” to styles that support it. It also invokes a default command: *package intel 1*. This package command is used to set options for the INTEL package. The default package command will specify that INTEL calculations are performed in mixed precision, that the number of OpenMP threads is specified by the OMP\_NUM\_THREADS environment variable, and that if co-processors are present and the binary was built with offload support, that 1 co-processor per node will be used with automatic balancing of work between the CPU and the co-processor.

You can specify different options for the INTEL package by using the “-pk intel Nphi” *command-line switch* with keyword/value pairs as specified in the documentation. Here, Nphi = # of Xeon Phi co-processors/node (ignored without offload support). Common options to the INTEL package include *omp* to override any OMP\_NUM\_THREADS setting and specify the number of OpenMP threads, *mode* to set the floating-point precision mode, and *lrt* to enable Long-Range Thread mode as described below. See the *package intel* command for details, including the default values used for all its options if not specified, and how to set the number of OpenMP threads via the OMP\_NUM\_THREADS environment variable if desired.

Examples (see documentation for your MPI/Machine for differences in launching MPI applications):

```
mpirun -np 72 -ppn 36 lmp_machine -sf intel -in in.script
↳ # 2 nodes, 36 MPI tasks/node, $OMP_NUM_THREADS OpenMP Threads
mpirun -np 72 -ppn 36 lmp_machine -sf intel -in in.script -pk intel 0 omp 2 mode double
↳ # Don't use any co-processors that might be available, use 2 OpenMP threads for each
↳ task, use double precision
```

## Or run with the INTEL package by editing an input script

As an alternative to adding command-line arguments, the input script can be edited to enable the INTEL package. This requires adding the *package intel* command to the top of the input script. For the second example above, this would be:

```
package intel 0 omp 2 mode double
```

To enable the INTEL package only for individual styles, you can add an “intel” suffix to the individual style, e.g.:

```
pair_style lj/cut/intel 2.5
```

Alternatively, the *suffix intel* command can be added to the input script to enable INTEL styles for the commands that follow in the input script.

## Tuning for Performance

**Note:** The INTEL package will perform better with modifications to the input script when *PPPM* is used: *k-space\_modify diff ad* should be added to the input script.

Long-Range Thread (LRT) mode is an option to the *package intel* command that can improve performance when using *PPPM* for long-range electrostatics on processors with SMT. It generates an extra pthread for each MPI task. The thread is dedicated to performing some of the PPPM calculations and MPI communications. This feature requires setting the pre-processor flag -DLMP\_INTEL\_USELRT in the makefile when compiling LAMMPS. It is unset in the default makefiles (*Makefile.mpi* and *Makefile.serial*) but it is set in all makefiles tuned for the INTEL package. On Intel Xeon Phi x200 series CPUs, the LRT feature will likely improve performance, even on a single node. On Intel Xeon processors, using this mode might result in better performance when using multiple nodes, depending on the

specific machine configuration. To enable LRT mode, specify that the number of OpenMP threads is one less than would normally be used for the run and add the “lrt yes” option to the “-pk” command-line suffix or “package intel” command. For example, if a run would normally perform best with “-pk intel 0 omp 4”, instead use “-pk intel 0 omp 3 lrt yes”. When using LRT, you should set the environment variable “KMP\_AFFINITY=none”. LRT mode is not supported when using offload.

---

**Note:** Changing the *newton* setting to off can improve performance and/or scalability for simple 2-body potentials such as lj/cut or when using LRT mode on processors supporting AVX-512.

---

Not all styles are supported in the INTEL package. You can mix the INTEL package with styles from the *OPT* package or the *OPENMP* package. Of course, this requires that these packages were installed at build time. This can be performed automatically by using “-sf hybrid intel opt” or “-sf hybrid intel omp” command-line options. Alternatively, the “opt” and “omp” suffixes can be appended manually in the input script. For the latter, the *package omp* command must be in the input script or the “-pk omp Nt” *command-line switch* must be used where Nt is the number of OpenMP threads. The number of OpenMP threads should not be set differently for the different packages. Note that the *suffix hybrid intel omp* command can also be used within the input script to automatically append the “omp” suffix to styles when INTEL styles are not available.

---

**Note:** For simulations on higher node counts, add *processors \* \* \* grid numa* to the beginning of the input script for better scalability.

---

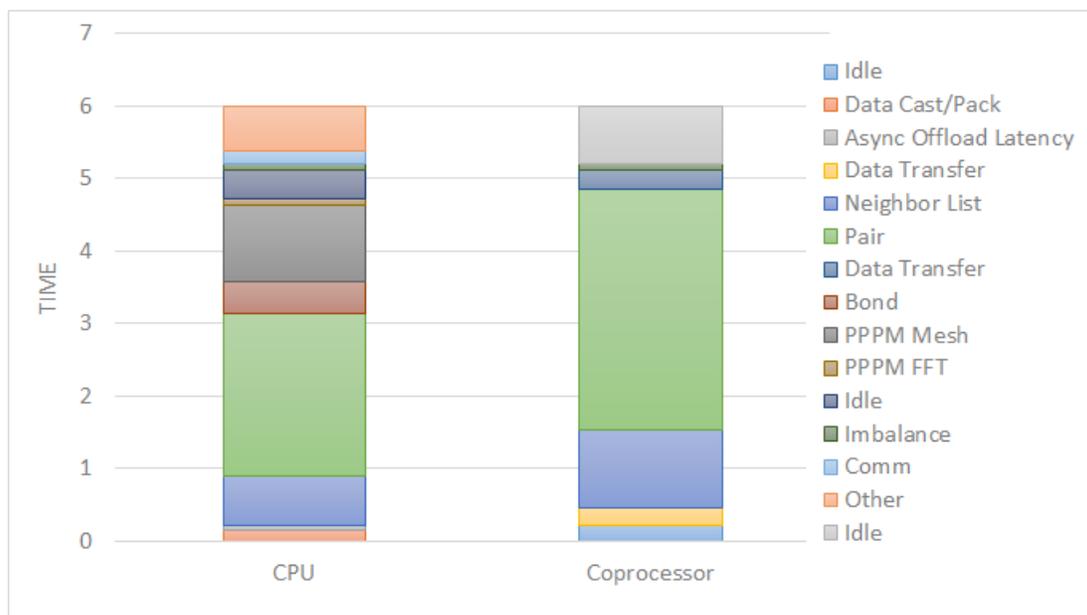
When running on many nodes, performance might be better when using fewer OpenMP threads and more MPI tasks. This will depend on the simulation and the machine. Using the *verlet/split* run style might also give better performance for simulations with *PPPM* electrostatics. Note that this is an alternative to LRT mode and the two cannot be used together.

Currently, when using Intel MPI with Intel Xeon Phi x200 series CPUs, better performance might be obtained by setting the environment variable “I\_MPI\_SHM\_LMT=shm” for Linux kernels that do not yet have full support for AVX-512. Runs on Intel Xeon Phi x200 series processors will always perform better using MCDRAM. Please consult your system documentation for the best approach to specify that MPI runs are performed in MCDRAM.

## Tuning for Offload Performance

The default settings for offload should give good performance.

When using LAMMPS with offload to Intel co-processors, best performance will typically be achieved with concurrent calculations performed on both the CPU and the co-processor. This is achieved by offloading only a fraction of the neighbor and pair computations to the co-processor or using *hybrid* pair styles where only one style uses the “intel” suffix. For simulations with long-range electrostatics or bond, angle, dihedral, improper calculations, computation and data transfer to the co-processor will run concurrently with computations and MPI communications for these calculations on the host CPU. This is illustrated in the figure below for the rhodopsin protein benchmark running on E5-2697v2 processors with a Intel Xeon Phi 7120p co-processor. In this plot, the vertical axis is time and routines running at the same time are running concurrently on both the host and the co-processor.



The fraction of the offloaded work is controlled by the *balance* keyword in the *package intel* command. A balance of 0 runs all calculations on the CPU. A balance of 1 runs all supported calculations on the co-processor. A balance of 0.5 runs half of the calculations on the co-processor. Setting the balance to -1 (the default) will enable dynamic load balancing that continuously adjusts the fraction of offloaded work throughout the simulation. Because data transfer cannot be timed, this option typically produces results within 5 to 10 percent of the optimal fixed balance.

If running short benchmark runs with dynamic load balancing, adding a short warm-up run (10-20 steps) will allow the load-balancer to find a near-optimal setting that will carry over to additional runs.

The default for the *package intel* command is to have all the MPI tasks on a given compute node use a single Xeon Phi co-processor. In general, running with a large number of MPI tasks on each node will perform best with offload. Each MPI task will automatically get affinity to a subset of the hardware threads available on the co-processor. For example, if your card has 61 cores, with 60 cores available for offload and 4 hardware threads per core (240 total threads), running with 24 MPI tasks per node will cause each MPI task to use a subset of 10 threads on the co-processor. Fine tuning of the number of threads to use per MPI task or the number of threads to use per core can be accomplished with keyword settings of the *package intel* command.

The INTEL package has two modes for deciding which atoms will be handled by the co-processor. This choice is controlled with the *ghost* keyword of the *package intel* command. When set to 0, ghost atoms (atoms at the borders between MPI tasks) are not offloaded to the card. This allows for overlap of MPI communication of forces with computation on the co-processor when the *newton* setting is “on”. The default is dependent on the style being used, however, better performance may be achieved by setting this option explicitly.

When using offload with CPU Hyper-Threading disabled, it may help performance to use fewer MPI tasks and OpenMP threads than available cores. This is due to the fact that additional threads are generated internally to handle the asynchronous offload tasks.

If pair computations are being offloaded to an Intel Xeon Phi co-processor, a diagnostic line is printed to the screen (not to the log file), during the setup phase of a run, indicating that offload mode is being used and indicating the number of co-processor threads per MPI task. Additionally, an offload timing summary is printed at the end of each run. When offloading, the frequency for *atom sorting* is changed to 1 so that the per-atom data is effectively sorted at every rebuild of the neighbor lists. All the available co-processor threads on each Phi will be divided among MPI tasks, unless the *tp task* option of the “-pk intel” *command-line switch* is used to limit the co-processor threads per MPI task.

## Restrictions

When offloading to a co-processor, *hybrid* styles that require skip lists for neighbor builds cannot be offloaded. Using *hybrid/overlay* is allowed. Only one intel accelerated style may be used with hybrid styles when offloading. *Special\_bonds* exclusion lists are not currently supported with offload, however, the same effect can often be accomplished by setting cutoffs for excluded atom types to 0. None of the pair styles in the INTEL package currently support the “inner”, “middle”, “outer” options for rRESPA integration via the *run\_style respa* command; only the “pair” option is supported.

## References

- Brown, W.M., Carrillo, J.-M.Y., Mishra, B., Gavhane, N., Thakkar, F.M., De Kraker, A.R., Yamada, M., Ang, J.A., Plimpton, S.J., “Optimizing Classical Molecular Dynamics in LAMMPS”, in Intel Xeon Phi Processor High Performance Programming: Knights Landing Edition, J. Jeffers, J. Reinders, A. Sodani, Eds. Morgan Kaufmann.
- Brown, W. M., Semin, A., Hebenstreit, M., Khvostov, S., Raman, K., Plimpton, S.J. *Increasing Molecular Dynamics Simulation Rates with an 8-Fold Increase in Electrical Power Efficiency*. 2016 High Performance Computing, Networking, Storage and Analysis, SC16: International Conference (pp. 82-95).
- Brown, W.M., Carrillo, J.-M.Y., Gavhane, N., Thakkar, F.M., Plimpton, S.J. *Optimizing Legacy Molecular Dynamics Software with Directive-Based Offload*. Computer Physics Communications. 2015. 195: p. 95-101.

### 7.4.3 KOKKOS package

Kokkos is a templated C++ library that provides abstractions to allow a single implementation of an application kernel (e.g. a pair style) to run efficiently on different kinds of hardware, such as GPUs, Intel Xeon Phis, or many-core CPUs. Kokkos maps the C++ kernel onto different back end languages such as CUDA, OpenMP, or Pthreads. The Kokkos library also provides data abstractions to adjust (at compile time) the memory layout of data structures like 2d and 3d arrays to optimize performance on different hardware. For more information on Kokkos, see [the Kokkos GitHub page](#).

The LAMMPS KOKKOS package contains versions of pair, fix, and atom styles that use data structures and macros provided by the Kokkos library, which is included with LAMMPS in `/lib/kokkos`. The KOKKOS package was developed primarily by Christian Trott (Sandia) and Stan Moore (Sandia) with contributions of various styles by others, including Sikandar Mashayak (UIUC), Ray Shan (Sandia), and Dan Ibanez (Sandia). For more information on developing using Kokkos abstractions see the Kokkos [Wiki](#).

Kokkos currently provides support for 4 modes of execution (per MPI task). These are Serial (MPI-only for CPUs and Intel Phi), OpenMP (threading for many-core CPUs and Intel Phi), CUDA (for NVIDIA GPUs) and HIP (for AMD GPUs). You choose the mode at build time to produce an executable compatible with a specific hardware.

---

#### C++14 support

Kokkos requires using a compiler that supports the c++14 standard. For some compilers, it may be necessary to add a flag to enable c++14 support. For example, the GNU compiler uses the `-std=c++14` flag. For a list of compilers that have been tested with the Kokkos library, see the Kokkos [README](#).

---

#### NVIDIA CUDA support

To build with Kokkos support for NVIDIA GPUs, the NVIDIA CUDA toolkit software version 9.0 or later must be installed on your system. See the discussion for the *GPU package* for details of how to check and do this.

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### CUDA and MPI library compatibility

Kokkos with CUDA currently implicitly assumes that the MPI library is GPU-aware. This is not always the case, especially when using pre-compiled MPI libraries provided by a Linux distribution. This is not a problem when using only a single GPU with a single MPI rank. When running with multiple MPI ranks, you may see segmentation faults without GPU-aware MPI support. These can be avoided by adding the flags `-pk kokkos gpu/aware off` to the LAMMPS command line or by using the command `package kokkos gpu/aware off` in the input file.

---

### AMD GPU support

To build with Kokkos the HIPCC compiler from the AMD ROCm software version 3.5 or later is required. Supporting this Kokkos mode in LAMMPS is still work in progress. Please contact the LAMMPS developers if you run into problems.

---

### Building LAMMPS with the KOKKOS package

See the [Build extras](#) page for instructions.

### Running LAMMPS with the KOKKOS package

All Kokkos operations occur within the context of an individual MPI task running on a single node of the machine. The total number of MPI tasks used by LAMMPS (one or multiple per compute node) is set in the usual manner via the `mpirun` or `mpiexec` commands, and is independent of Kokkos. E.g. the `mpirun` command in OpenMPI does this via its `-np` and `-ppn` switches. Ditto for MPICH via `-np` and `-ppn`.

### Running on a multi-core CPU

Here is a quick overview of how to use the KOKKOS package for CPU acceleration, assuming one or more 16-core nodes.

```
mpirun -np 16 lmp_kokkos_mpi_only -k on -sf kk -in in.lj      # 1 node, 16 MPI tasks/
↳node, no multi-threading
mpirun -np 2 -ppn 1 lmp_kokkos_omp -k on t 16 -sf kk -in in.lj # 2 nodes, 1 MPI task/
↳node, 16 threads/task
mpirun -np 2 lmp_kokkos_omp -k on t 8 -sf kk -in in.lj      # 1 node, 2 MPI tasks/
↳node, 8 threads/task
mpirun -np 32 -ppn 4 lmp_kokkos_omp -k on t 4 -sf kk -in in.lj # 8 nodes, 4 MPI tasks/
↳node, 4 threads/task
```

To run using the KOKKOS package, use the “`-k on`”, “`-sf kk`” and “`-pk kokkos`” *command-line switches* in your `mpirun` command. You must use the “`-k on`” *command-line switch* to enable the KOKKOS package. It takes additional arguments for hardware settings appropriate to your system. For OpenMP use:

```
-k on t Nt
```

The “`t Nt`” option specifies how many OpenMP threads per MPI task to use with a node. The default is `Nt = 1`, which is MPI-only mode. Note that the product of MPI tasks \* OpenMP threads/task should not exceed the physical number of cores (on a node), otherwise performance will suffer. If Hyper-Threading (HT) is enabled, then the product of MPI tasks \* OpenMP threads/task should not exceed the physical number of cores \* hardware threads. The “`-k on`”

switch also issues a “package kokkos” command (with no additional arguments) which sets various KOKKOS options to default values, as discussed on the [package](#) command doc page.

The “-sf kk” *command-line switch* will automatically append the “/kk” suffix to styles that support it. In this manner no modification to the input script is needed. Alternatively, one can run with the KOKKOS package by editing the input script as described below.

---

**Note:** When using a single OpenMP thread, the Kokkos Serial back end (i.e. Makefile.kokkos\_mpi\_only) will give better performance than the OpenMP back end (i.e. Makefile.kokkos\_omp) because some of the overhead to make the code thread-safe is removed.

---

---

**Note:** Use the “-pk kokkos” *command-line switch* to change the default *package kokkos* options. See its doc page for details and default settings. Experimenting with its options can provide a speed-up for specific calculations. For example:

---

```
mpirun -np 16 lmp_kokkos_mpi_only -k on -sf kk -pk kokkos newton on neigh half comm no -  
↪ in in.lj          # Newton on, Half neighbor list, non-threaded comm
```

If the *newton* command is used in the input script, it can also override the Newton flag defaults.

For half neighbor lists and OpenMP, the KOKKOS package uses data duplication (i.e. thread-private arrays) by default to avoid thread-level write conflicts in the force arrays (and other data structures as necessary). Data duplication is typically fastest for small numbers of threads (i.e. 8 or less) but does increase memory footprint and is not scalable to large numbers of threads. An alternative to data duplication is to use thread-level atomic operations which do not require data duplication. The use of atomic operations can be enforced by compiling LAMMPS with the “-DLMP\_KOKKOS\_USE\_ATOMICS” pre-processor flag. Most but not all Kokkos-enabled pair\_styles support data duplication. Alternatively, full neighbor lists avoid the need for duplication or atomic operations but require more compute operations per atom. When using the Kokkos Serial back end or the OpenMP back end with a single thread, no duplication or atomic operations are used. For CUDA and half neighbor lists, the KOKKOS package always uses atomic operations.

## CPU Cores, Sockets and Thread Affinity

When using multi-threading, it is important for performance to bind both MPI tasks to physical cores, and threads to physical cores, so they do not migrate during a simulation.

If you are not certain MPI tasks are being bound (check the defaults for your MPI installation), binding can be forced with these flags:

```
OpenMPI 1.8: mpirun -np 2 --bind-to socket --map-by socket ./lmp_openmpi ...  
Mvapich2 2.0: mpiexec -np 2 --bind-to socket --map-by socket ./lmp_mvapich ...
```

For binding threads with KOKKOS OpenMP, use thread affinity environment variables to force binding. With OpenMP 3.1 (gcc 4.7 or later, intel 12 or later) setting the environment variable `OMP_PROC_BIND=true` should be sufficient. In general, for best performance with OpenMP 4.0 or later set `OMP_PROC_BIND=spread` and `OMP_PLACES=threads`. For binding threads with the KOKKOS pthreads option, compile LAMMPS with the hwloc or libnuma support enabled as described in the [extra build options page](#).

## Running on Knight’s Landing (KNL) Intel Xeon Phi

Here is a quick overview of how to use the KOKKOS package for the Intel Knight’s Landing (KNL) Xeon Phi:

KNL Intel Phi chips have 68 physical cores. Typically 1 to 4 cores are reserved for the OS, and only 64 or 66 cores are used. Each core has 4 Hyper-Threads, so there are effectively  $N = 256$  ( $4 \times 64$ ) or  $N = 264$  ( $4 \times 66$ ) cores to run on. The product of MPI tasks \* OpenMP threads/task should not exceed this limit, otherwise performance will suffer. Note that with the KOKKOS package you do not need to specify how many KNLs there are per node; each KNL is simply treated as running some number of MPI tasks.

Examples of mpirun commands that follow these rules are shown below.

```
# Running on an Intel KNL node with 68 cores (272 threads/node via 4x hardware
↳threading):
mpirun -np 64 lmp_kokkos_phi -k on t 4 -sf kk -in in.lj      # 1 node, 64 MPI tasks/node,
↳ 4 threads/task
mpirun -np 66 lmp_kokkos_phi -k on t 4 -sf kk -in in.lj      # 1 node, 66 MPI tasks/node,
↳ 4 threads/task
mpirun -np 32 lmp_kokkos_phi -k on t 8 -sf kk -in in.lj      # 1 node, 32 MPI tasks/node,
↳ 8 threads/task
mpirun -np 512 -ppn 64 lmp_kokkos_phi -k on t 4 -sf kk -in in.lj # 8 nodes, 64 MPI
↳tasks/node, 4 threads/task
```

The `-np` setting of the `mpirun` command sets the number of MPI tasks/node. The “`-k on t Nt`” command-line switch sets the number of threads/task as `Nt`. The product of these two values should be `N`, i.e. 256 or 264.

**Note:** The default for the *package kokkos* command when running on KNL is to use “half” neighbor lists and set the Newton flag to “on” for both pairwise and bonded interactions. This will typically be best for many-body potentials. For simpler pairwise potentials, it may be faster to use a “full” neighbor list with Newton flag to “off”. Use the “`-pk kokkos`” *command-line switch* to change the default *package kokkos* options. See its page for details and default settings. Experimenting with its options can provide a speed-up for specific calculations. For example:

```
mpirun -np 64 lmp_kokkos_phi -k on t 4 -sf kk -pk kokkos comm host -in in.reax      #
↳Newton on, half neighbor list, threaded comm
mpirun -np 64 lmp_kokkos_phi -k on t 4 -sf kk -pk kokkos newton off neigh full comm no -
↳in in.lj      # Newton off, full neighbor list, non-threaded comm
```

**Note:** MPI tasks and threads should be bound to cores as described above for CPUs.

**Note:** To build with Kokkos support for Intel Xeon Phi co-processors such as Knight’s Corner (KNC), your system must be configured to use them in “native” mode, not “offload” mode like the INTEL package supports.

## Running on GPUs

Use the “-k” *command-line switch* to specify the number of GPUs per node. Typically the -np setting of the mpirun command should set the number of MPI tasks/node to be equal to the number of physical GPUs on the node. You can assign multiple MPI tasks to the same GPU with the KOKKOS package, but this is usually only faster if some portions of the input script have not been ported to use Kokkos. In this case, also packing/unpacking communication buffers on the host may give speedup (see the KOKKOS *package* command). Using CUDA MPS is recommended in this scenario.

Using a GPU-aware MPI library is highly recommended. GPU-aware MPI use can be avoided by using *-pk kokkos gpu/aware off*. As above for multi-core CPUs (and no GPU), if N is the number of physical cores/node, then the number of MPI tasks/node should not exceed N.

```
-k on g Ng
```

Here are examples of how to use the KOKKOS package for GPUs, assuming one or more nodes, each with two GPUs:

```
mpirun -np 2 lmp_kokkos_cuda_openmpi -k on g 2 -sf kk -in in.lj # 1 node, 2
↳MPI tasks/node, 2 GPUs/node
mpirun -np 32 -ppn 2 lmp_kokkos_cuda_openmpi -k on g 2 -sf kk -in in.lj # 16 nodes, 2
↳MPI tasks/node, 2 GPUs/node (32 GPUs total)
```

---

**Note:** The default for the *package kokkos* command when running on GPUs is to use “full” neighbor lists and set the Newton flag to “off” for both pairwise and bonded interactions, along with threaded communication. When running on Maxwell or Kepler GPUs, this will typically be best. For Pascal GPUs and beyond, using “half” neighbor lists and setting the Newton flag to “on” may be faster. For many pair styles, setting the neighbor binsize equal to twice the CPU default value will give speedup, which is the default when running on GPUs. Use the “-pk kokkos” *command-line switch* to change the default *package kokkos* options. See its page for details and default settings. Experimenting with its options can provide a speed-up for specific calculations. For example:

```
mpirun -np 2 lmp_kokkos_cuda_openmpi -k on g 2 -sf kk -pk kokkos newton on neigh half
↳binsize 2.8 -in in.lj # Newton on, half neighbor list, set binsize = neighbor
↳ghost cutoff
```

---

**Note:** When using a GPU, you will achieve the best performance if your input script does not use fix or compute styles which are not yet Kokkos-enabled. This allows data to stay on the GPU for multiple timesteps, without being copied back to the host CPU. Invoking a non-Kokkos fix or compute, or performing I/O for *thermo* or *dump* output will cause data to be copied back to the CPU incurring a performance penalty.

---

**Note:** To get an accurate timing breakdown between time spend in pair, kspace, etc., you must set the environment variable CUDA\_LAUNCH\_BLOCKING=1. However, this will reduce performance and is not recommended for production runs.

---

## Run with the KOKKOS package by editing an input script

Alternatively the effect of the “-sf” or “-pk” switches can be duplicated by adding the *package kokkos* or *suffix kk* commands to your input script.

The discussion above for building LAMMPS with the KOKKOS package, the mpirun/mpiexec command, and setting appropriate thread are the same.

You must still use the “-k on” *command-line switch* to enable the KOKKOS package, and specify its additional arguments for hardware options appropriate to your system, as documented above.

You can use the *suffix kk* command, or you can explicitly add a “kk” suffix to individual styles in your input script, e.g.

```
pair_style lj/cut/kk 2.5
```

You only need to use the *package kokkos* command if you wish to change any of its option defaults, as set by the “-k on” *command-line switch*.

### Using OpenMP threading and CUDA together:

With the KOKKOS package, both OpenMP multi-threading and GPUs can be compiled and used together in a few special cases. In the makefile for the conventional build, the KOKKOS\_DEVICES variable must include both, “Cuda” and “OpenMP”, as is the case for /src/MAKE/OPTIONS/Makefile.kokkos\_cuda\_mpi.

```
KOKKOS_DEVICES=Cuda,OpenMP
```

When building with CMake you need to enable both features as it is done in the kokkos-cuda.cmake CMake preset file.

```
cmake ../cmake -DKokkos_ENABLE_CUDA=yes -DKokkos_ENABLE_OPENMP=yes
```

The suffix “/kk” is equivalent to “/kk/device”, and for Kokkos CUDA, using the “-sf kk” in the command line gives the default CUDA version everywhere. However, if the “/kk/host” suffix is added to a specific style in the input script, the Kokkos OpenMP (CPU) version of that specific style will be used instead. Set the number of OpenMP threads as “t Nt” and the number of GPUs as “g Ng”

```
-k on t Nt g Ng
```

For example, the command to run with 1 GPU and 8 OpenMP threads is then:

```
mpiexec -np 1 lmp_kokkos_cuda_openmpi -in in.lj -k on g 1 t 8 -sf kk
```

Conversely, if the “-sf kk/host” is used in the command line and then the “/kk” or “/kk/device” suffix is added to a specific style in your input script, then only that specific style will run on the GPU while everything else will run on the CPU in OpenMP mode. Note that the execution of the CPU and GPU styles will NOT overlap, except for a special case:

A kspace style and/or molecular topology (bonds, angles, etc.) running on the host CPU can overlap with a pair style running on the GPU. First compile with “-default-stream per-thread” added to CCFLAGS in the Kokkos CUDA Makefile. Then explicitly use the “/kk/host” suffix for kspace and bonds, angles, etc. in the input file and the “kk” suffix (equal to “kk/device”) on the command line. Also make sure the environment variable CUDA\_LAUNCH\_BLOCKING is not set to “1” so CPU/GPU overlap can occur.

## Performance to expect

The performance of KOKKOS running in different modes is a function of your hardware, which KOKKOS-enable styles are used, and the problem size.

Generally speaking, the following rules of thumb apply:

- When running on CPUs only, with a single thread per MPI task, performance of a KOKKOS style is somewhere between the standard (un-accelerated) styles (MPI-only mode), and those provided by the OPENMP package. However the difference between all 3 is small (less than 20%).
- When running on CPUs only, with multiple threads per MPI task, performance of a KOKKOS style is a bit slower than the OPENMP package.
- When running large number of atoms per GPU, KOKKOS is typically faster than the GPU package when compiled for double precision. The benefit of using single or mixed precision with the GPU package depends significantly on the hardware in use and the simulated system and pair style.
- When running on Intel hardware, KOKKOS is not as fast as the INTEL package, which is optimized for x86 hardware (not just from Intel) and compilation with the Intel compilers. The INTEL package also can increase the vector length of vector instructions by switching to single or mixed precision mode.

See the [Benchmark page](#) of the LAMMPS website for performance of the KOKKOS package on different hardware.

## Advanced Kokkos options

There are other allowed options when building with the KOKKOS package that can improve performance or assist in debugging or profiling. They are explained on the *KOKKOS section of the build extras* doc page,

## Restrictions

Currently, there are no precision options with the KOKKOS package. All compilation and computation is performed in double precision.

## 7.4.4 OPENMP package

The OPENMP package was developed by Axel Kohlmeyer at Temple University. It provides optimized and multi-threaded versions of many pair styles, nearly all bonded styles (bond, angle, dihedral, improper), several Kspace styles, and a few fix styles. It uses the OpenMP interface for multi-threading, but can also be compiled without OpenMP support, providing optimized serial styles in that case.

## Required hardware/software

To enable multi-threading, your compiler must support the OpenMP interface. You should have one or more multi-core CPUs, as multiple threads can only be launched by each MPI task on the local node (using shared memory).

## Building LAMMPS with the OPENMP package

See the *Build extras* page for instructions.

## Run with the OPENMP package from the command line

These examples assume one or more 16-core nodes.

```
env OMP_NUM_THREADS=16 lmp_omp -sf omp -in in.script           # 1 MPI task, 16 threads,
↳according to OMP_NUM_THREADS
lmp_mpi -sf omp -in in.script                                 # 1 MPI task, no threads,
↳optimized kernels
mpirun -np 4 lmp_omp -sf omp -pk omp 4 -in in.script         # 4 MPI tasks, 4 threads/
↳task
mpirun -np 32 -ppn 4 lmp_omp -sf omp -pk omp 4 -in in.script # 8 nodes, 4 MPI tasks/
↳node, 4 threads/task
```

The `mpirun` or `mpiexec` command sets the total number of MPI tasks used by LAMMPS (one or multiple per compute node) and the number of MPI tasks used per node. E.g. the `mpirun` command in MPICH does this via its `-np` and `-ppn` switches. Ditto for OpenMPI via `-np` and `-npernode`.

You need to choose how many OpenMP threads per MPI task will be used by the OPENMP package. Note that the product of MPI tasks \* threads/task should not exceed the physical number of cores (on a node), otherwise performance will suffer.

As in the lines above, use the “`-sf omp`” *command-line switch*, which will automatically append “`omp`” to styles that support it. The “`-sf omp`” switch also issues a default *package omp 0* command, which will set the number of threads per MPI task via the `OMP_NUM_THREADS` environment variable.

You can also use the “`-pk omp Nt`” *command-line switch*, to explicitly set  $N_t = \#$  of OpenMP threads per MPI task to use, as well as additional options. Its syntax is the same as the *package omp* command whose page gives details, including the default values used if it is not specified. It also gives more details on how to set the number of threads via the `OMP_NUM_THREADS` environment variable.

## Or run with the OPENMP package by editing an input script

The discussion above for the `mpirun/mpiexec` command, MPI tasks/node, and threads/MPI task is the same.

Use the *suffix omp* command, or you can explicitly add an “`omp`” suffix to individual styles in your input script, e.g.

```
pair_style lj/cut/omp 2.5
```

You must also use the *package omp* command to enable the OPENMP package. When you do this you also specify how many threads per MPI task to use. The command page explains other options and how to set the number of threads via the `OMP_NUM_THREADS` environment variable.

## Speed-up to expect

Depending on which styles are accelerated, you should look for a reduction in the “Pair time”, “Bond time”, “KSpace time”, and “Loop time” values printed at the end of a run.

You may see a small performance advantage (5 to 20%) when running a OPENMP style (in serial or parallel) with a single thread per MPI task, versus running standard LAMMPS with its standard un-accelerated styles (in serial or all-MPI parallelization with 1 task/core). This is because many of the OPENMP styles contain similar optimizations to those used in the OPT package, described in *the OPT package* doc page.

With multiple threads/task, the optimal choice of number of MPI tasks/node and OpenMP threads/task can vary a lot and should always be tested via benchmark runs for a specific simulation running on a specific machine, paying attention to guidelines discussed in the next sub-section.

A description of the multi-threading strategy used in the OPENMP package and some performance examples are [presented here](#).

## Guidelines for best performance

For many problems on current generation CPUs, running the OPENMP package with a single thread/task is faster than running with multiple threads/task. This is because the MPI parallelization in LAMMPS is often more efficient than multi-threading as implemented in the OPENMP package. The parallel efficiency (in a threaded sense) also varies for different OPENMP styles.

Using multiple threads/task can be more effective under the following circumstances:

- Individual compute nodes have a significant number of CPU cores but the CPU itself has limited memory bandwidth, e.g. for Intel Xeon 53xx (Clovertown) and 54xx (Harpertown) quad-core processors. Running one MPI task per CPU core will result in significant performance degradation, so that running with 4 or even only 2 MPI tasks per node is faster. Running in hybrid MPI+OpenMP mode will reduce the inter-node communication bandwidth contention in the same way, but offers an additional speedup by utilizing the otherwise idle CPU cores.
- The interconnect used for MPI communication does not provide sufficient bandwidth for a large number of MPI tasks per node. For example, this applies to running over gigabit ethernet or on Cray XT4 or XT5 series supercomputers. As in the aforementioned case, this effect worsens when using an increasing number of nodes.
- The system has a spatially inhomogeneous particle density which does not map well to the *domain decomposition scheme* or *load-balancing* options that LAMMPS provides. This is because multi-threading achieves parallelism over the number of particles, not via their distribution in space.
- A machine is being used in “capability mode”, i.e. near the point where MPI parallelism is maxed out. For example, this can happen when using the *PPPM solver* for long-range electrostatics on large numbers of nodes. The scaling of the KSpace calculation (see the *kpace\_style* command) becomes the performance-limiting factor. Using multi-threading allows less MPI tasks to be invoked and can speed-up the long-range solver, while increasing overall performance by parallelizing the pairwise and bonded calculations via OpenMP. Likewise additional speedup can be sometimes be achieved by increasing the length of the Coulombic cutoff and thus reducing the work done by the long-range solver. Using the *run\_style verlet/split* command, which is compatible with the OPENMP package, is an alternative way to reduce the number of MPI tasks assigned to the KSpace calculation.

Additional performance tips are as follows:

- The best parallel efficiency from *omp* styles is typically achieved when there is at least one MPI task per physical CPU chip, i.e. socket or die.
- It is usually most efficient to restrict threading to a single socket, i.e. use one or more MPI task per socket.
- NOTE: By default, several current MPI implementations use a processor affinity setting that restricts each MPI task to a single CPU core. Using multi-threading in this mode will force all threads to share the one core and thus is likely to be counterproductive. Instead, binding MPI tasks to a (multi-core) socket, should solve this issue.



## Restrictions

None.

## 7.4.5 OPT package

The OPT package was developed by James Fischer (High Performance Technologies), David Richie, and Vincent Natoli (Stone Ridge Technologies). It contains a handful of pair styles whose compute() methods were rewritten in C++ templated form to reduce the overhead due to if tests and other conditional code.

### Required hardware/software

Any hardware. Any compiler.

### Building LAMMPS with the OPT package

See the *Build extras* page for instructions.

### Run with the OPT package from the command line

```
lmp_mpi -sf opt -in in.script           # run in serial
mpirun -np 4 lmp_mpi -sf opt -in in.script # run in parallel
```

Use the “-sf opt” *command-line switch*, which will automatically append “opt” to styles that support it.

### Or run with the OPT package by editing an input script

Use the *suffix opt* command, or you can explicitly add an “opt” suffix to individual styles in your input script, e.g.

```
pair_style lj/cut/opt 2.5
```

### Speed-up to expect

You should see a reduction in the “Pair time” value printed at the end of a run. On most machines for reasonable problem sizes, it will be a 5 to 20% savings.

### Guidelines for best performance

Just try out an OPT pair style to see how it performs.

## Restrictions

None.

Inverting this list, LAMMPS currently has acceleration support for three kinds of hardware, via the listed packages:

Many-core CPUs	<i>INTEL, KOKKOS, OPENMP, OPT</i> packages
GPUs	<i>GPU, KOKKOS</i> packages
Intel Phi/AVX	<i>INTEL, KOKKOS</i> packages

Which package is fastest for your hardware may depend on the size problem you are running and what commands (accelerated and non-accelerated) are invoked by your input script. While these doc pages include performance guidelines, there is no substitute for trying out the different packages appropriate to your hardware.

Any accelerated style has the same name as the corresponding standard style, except that a suffix is appended. Otherwise, the syntax for the command that uses the style is identical, their functionality is the same, and the numerical results it produces should also be the same, except for precision and round-off effects.

For example, all of these styles are accelerated variants of the Lennard-Jones *pair\_style lj/cut*:

- *pair\_style lj/cut/gpu*
- *pair\_style lj/cut/intel*
- *pair\_style lj/cut/kk*
- *pair\_style lj/cut/omp*
- *pair\_style lj/cut/opt*

To see what accelerate styles are currently available for a particular style, find the style name in the *Commands* style pages (*fix,compute,pair,etc*) and see what suffixes are listed (g,i,k,o,t) with it. The doc pages for individual commands (e.g. *pair lj/cut* or *fix nve*) also list any accelerated variants available for that style.

To use an accelerator package in LAMMPS, and one or more of the styles it provides, follow these general steps. Details vary from package to package and are explained in the individual accelerator doc pages, listed above:

build the accelerator library	only for GPU package
install the accelerator package	make yes-opt, make yes-intel, etc
add compile/link flags to Makefile.machine in src/MAKE	only for INTEL, KOKKOS, OPENMP, OPT packages
re-build LAMMPS	make machine
prepare and test a regular LAMMPS simulation	Imp_machine -in in.script; mpirun -np 32 Imp_machine -in in.script
enable specific accelerator support via ‘-k on’ <i>command-line switch</i> ,	only needed for KOKKOS package
set any needed options for the package via “-pk” <i>command-line switch</i> or <i>package</i> command,	only if defaults need to be changed
use accelerated styles in your input via “-sf” <i>command-line switch</i> or <i>suffix</i> command	Imp_machine -in in.script -sf gpu

Note that the first 4 steps can be done as a single command with suitable make command invocations. This is discussed on the *Packages* doc pages, and its use is illustrated in the individual accelerator sections. Typically these steps only need to be done once, to create an executable that uses one or more accelerator packages.

The last 4 steps can all be done from the command-line when LAMMPS is launched, without changing your input script, as illustrated in the individual accelerator sections. Or you can add *package* and *suffix* commands to your input script.

---

**Note:** With a few exceptions, you can build a single LAMMPS executable with all its accelerator packages installed. Note however that the INTEL and KOKKOS packages require you to choose one of their hardware options when building for a specific platform. I.e. CPU or Phi option for the INTEL package. Or the OpenMP, Cuda, or Phi option for the KOKKOS package.

---

These are the exceptions. You cannot build a single executable with:

- both the INTEL Phi and KOKKOS Phi options
- the INTEL Phi or Kokkos Phi option, and the GPU package

See the examples/accelerate/README and make.list files for sample Make.py commands that build LAMMPS with any or all of the accelerator packages. As an example, here is a command that builds with all the GPU related packages installed (GPU, KOKKOS with Cuda), including settings to build the needed auxiliary GPU libraries for Kepler GPUs:

```
Make.py -j 16 -p omp gpu kokkos -cc nvcc wrap=mpi -gpu mode=double arch=35 -kokkos_
↳ cuda arch=35 lib=all file mpi
```

The examples/accelerate directory also has input scripts that can be used with all of the accelerator packages. See its README file for details.

Likewise, the bench directory has FERMI and KEPLER and PHI sub-directories with Make.py commands and input scripts for using all the accelerator packages on various machines. See the README files in those directories.

As mentioned above, the [Benchmark page](#) of the LAMMPS website gives performance results for the various accelerator packages for several of the standard LAMMPS benchmark problems, as a function of problem size and number of compute nodes, on different hardware platforms.

Here is a brief summary of what the various packages provide. Details are in the individual accelerator sections.

- Styles with a “gpu” suffix are part of the GPU package and can be run on Intel, NVIDIA, or AMD GPUs. The speed-up on a GPU depends on a variety of factors, discussed in the accelerator sections.
- Styles with an “intel” suffix are part of the INTEL package. These styles support vectorized single and mixed precision calculations, in addition to full double precision. In extreme cases, this can provide speedups over 3.5x on CPUs. The package also supports acceleration in “offload” mode to Intel(R) Xeon Phi(TM) co-processors. This can result in additional speedup over 2x depending on the hardware configuration.
- Styles with a “kk” suffix are part of the KOKKOS package, and can be run using OpenMP on multicore CPUs, on an NVIDIA or AMD GPU, or on an Intel Xeon Phi in “native” mode. The speed-up depends on a variety of factors, as discussed on the KOKKOS accelerator page.
- Styles with an “omp” suffix are part of the OPENMP package and allow a pair-style to be run in multi-threaded mode using OpenMP. This can be useful on nodes with high-core counts when using less MPI processes than cores is advantageous, e.g. when running with PPPM so that FFTs are run on fewer MPI processors or when the many MPI tasks would overload the available bandwidth for communication.
- Styles with an “opt” suffix are part of the OPT package and typically speed-up the pairwise calculations of your simulation by 5-25% on a CPU.

The individual accelerator package doc pages explain:

- what hardware and software the accelerated package requires
- how to build LAMMPS with the accelerated package
- how to run with the accelerated package either via command-line switches or modifying the input script
- speed-ups to expect
- guidelines for best performance

- restrictions

## 7.5 Comparison of various accelerator packages

The next section compares and contrasts the various accelerator options, since there are multiple ways to perform OpenMP threading, run on GPUs, optimize for vector units on CPUs and run on Intel Xeon Phi (co-)processors.

All of these packages can accelerate a LAMMPS calculation taking advantage of hardware features, but they do it in different ways and acceleration is not always guaranteed.

As a consequence, for a particular simulation on specific hardware, one package may be faster than the other. We give some guidelines below, but the best way to determine which package is faster for your input script is to try multiple of them on your machine and experiment with available performance tuning settings. See the benchmarking section below for examples where this has been done.

### Guidelines for using each package optimally:

- Both, the GPU and the KOKKOS package allows you to assign multiple MPI ranks (= CPU cores) to the same GPU. For the GPU package, this can lead to a speedup through better utilization of the GPU (by overlapping computation and data transfer) and more efficient computation of the non-GPU accelerated parts of LAMMPS through MPI parallelization, as all system data is maintained and updated on the host. For KOKKOS, there is less to no benefit from this, due to its different memory management model, which tries to retain data on the GPU.
- The GPU package moves per-atom data (coordinates, forces, and (optionally) neighbor list data, if not computed on the GPU) between the CPU and GPU at every timestep. The KOKKOS/CUDA package only does this on timesteps when a CPU calculation is required (e.g. to invoke a fix or compute that is non-GPU-ized). Hence, if you can formulate your input script to only use GPU-ized fixes and computes, and avoid doing I/O too often (thermo output, dump file snapshots, restart files), then the data transfer cost of the KOKKOS/CUDA package can be very low, causing it to run faster than the GPU package.
- The GPU package is often faster than the KOKKOS/CUDA package, when the number of atoms per GPU is on the smaller side. The crossover point, in terms of atoms/GPU at which the KOKKOS/CUDA package becomes faster depends strongly on the pair style. For example, for a simple Lennard Jones system the crossover (in single precision) is often about 50K-100K atoms per GPU. When performing double precision calculations the crossover point can be significantly smaller.
- Both KOKKOS and GPU package compute bonded interactions (bonds, angles, etc) on the CPU. If the GPU package is running with several MPI processes assigned to one GPU, the cost of computing the bonded interactions is spread across more CPUs and hence the GPU package can run faster in these cases.
- When using LAMMPS with multiple MPI ranks assigned to the same GPU, its performance depends to some extent on the available bandwidth between the CPUs and the GPU. This can differ significantly based on the available bus technology, capability of the host CPU and mainboard, the wiring of the buses and whether switches are used to increase the number of available bus slots, or if GPUs are housed in an external enclosure. This can become quite complex.
- To achieve significant acceleration through GPUs, both KOKKOS and GPU package require capable GPUs with fast on-device memory and efficient data transfer rates. This requests capable upper mid-level to high-end (desktop) GPUs. Using lower performance GPUs (e.g. on laptops) may result in a slowdown instead.
- For the GPU package, specifically when running in parallel with MPI, it is often more efficient to exclude the PPPM kspace style from GPU acceleration and instead run it - concurrently with a GPU accelerated pair style - on the CPU. This can often be easily achieved with placing a *suffix off* command before and a *suffix on* command after the *kspace\_style pppm* command.
- The KOKKOS/OpenMP and OPENMP package have different thread management strategies, which should result in OPENMP being more efficient for a small number of threads with increasing overhead as the number of threads

per MPI rank grows. The KOKKOS/OpenMP kernels have less overhead in that case, but have lower performance with few threads.

- The INTEL package contains many options and settings for achieving additional performance on Intel hardware (CPU and accelerator cards), but to unlock this potential, an Intel compiler is required. The package code will compile with GNU gcc, but it will not be as efficient.

**Differences between the GPU and KOKKOS packages:**

- The GPU package accelerates only pair force, neighbor list, and (parts of) PPPM calculations. The KOKKOS package attempts to run most of the calculation on the GPU, but can transparently support non-accelerated code (with a performance penalty due to having data transfers between host and GPU).
- The GPU package requires neighbor lists to be built on the CPU when using exclusion lists, or a triclinic simulation box.
- The GPU package can be compiled for CUDA or OpenCL and thus supports both, NVIDIA and AMD GPUs well. On NVIDIA hardware, using CUDA is typically resulting in equal or better performance over OpenCL.
- OpenCL in the GPU package does theoretically also support Intel CPUs or Intel Xeon Phi, but the native support for those in KOKKOS (or INTEL) is superior.



## HOWTO DISCUSSIONS

These doc pages describe how to perform various tasks with LAMMPS, both for users and developers. The [glossary](#) website page also lists MD terminology with links to corresponding LAMMPS manual pages. The example input scripts included in the examples directory of the LAMMPS distribution and highlighted on the [Examples](#) doc page also show how to setup and run various kinds of simulations.

### 8.1 General howto

#### 8.1.1 Restart a simulation

There are 3 ways to continue a long LAMMPS simulation. Multiple `run` commands can be used in the same input script. Each run will continue from where the previous run left off. Or binary restart files can be saved to disk using the `restart` command. At a later time, these binary files can be read via a `read_restart` command in a new script. Or they can be converted to text data files using the `-r` *command-line switch* and read by a `read_data` command in a new script.

Here we give examples of 2 scripts that read either a binary restart file or a converted data file and then issue a new run command to continue where the previous run left off. They illustrate what settings must be made in the new script. Details are discussed in the documentation for the `read_restart` and `read_data` commands.

Look at the `in.chain` input script provided in the `bench` directory of the LAMMPS distribution to see the original script that these 2 scripts are based on. If that script had the line

```
restart          50 tmp.restart
```

added to it, it would produce 2 binary restart files (`tmp.restart.50` and `tmp.restart.100`) as it ran.

This script could be used to read the first restart file and re-run the last 50 timesteps:

```
read_restart     tmp.restart.50

neighbor        0.4 bin
neigh_modify    every 1 delay 1

fix             1 all nve
fix             2 all langevin 1.0 1.0 10.0 904297

timestep        0.012

run             50
```

Note that the following commands do not need to be repeated because their settings are included in the restart file: *units*, *atom\_style*, *special\_bonds*, *pair\_style*, *bond\_style*. However these commands do need to be used, since their settings are not in the restart file: *neighbor*, *fix*, *timestep*.

If you actually use this script to perform a restarted run, you will notice that the thermodynamic data match at step 50 (if you also put a “thermo 50” command in the original script), but do not match at step 100. This is because the *fix langevin* command uses random numbers in a way that does not allow for perfect restarts.

As an alternate approach, the restart file could be converted to a data file as follows:

```
lmp_g++ -r tmp.restart.50 tmp.restart.data
```

Then, this script could be used to re-run the last 50 steps:

```
units          lj
atom_style     bond
pair_style     lj/cut 1.12
pair_modify    shift yes
bond_style     fene
special_bonds  0.0 1.0 1.0

read_data      tmp.restart.data

neighbor       0.4 bin
neigh_modify   every 1 delay 1

fix           1 all nve
fix           2 all langevin 1.0 1.0 10.0 904297

timestep       0.012

reset_timestep 50
run           50
```

Note that nearly all the settings specified in the original *in.chain* script must be repeated, except the *pair\_coeff* and *bond\_coeff* commands since the new data file lists the force field coefficients. Also, the *reset\_timestep* command is used to tell LAMMPS the current timestep. This value is stored in restart files, but not in data files.

## 8.1.2 Visualize LAMMPS snapshots

LAMMPS itself does not do visualization, but snapshots from LAMMPS simulations can be visualized (and analyzed) in a variety of ways.

Mention *dump image* and *dump movie*.

LAMMPS snapshots are created by the *dump* command which can create files in several formats. The native LAMMPS dump format is a text file (see “*dump atom*” or “*dump custom*”) which can be visualized by several popular visualization tools. The *dump image* and *dump movie* styles can output internally rendered images and convert a sequence of them to a movie during the MD run. Several programs included with LAMMPS as auxiliary tools can convert between LAMMPS format files and other formats. See the *Tools* page for details.

A Python-based toolkit distributed by our group can read native LAMMPS dump files, including custom dump files with additional columns of user-specified atom information, and convert them to various formats or pipe them into visualization software directly. See the [Pizza.py WWW site](#) for details. Specifically, *Pizza.py* can convert LAMMPS dump files into PDB, XYZ, *EnSight*, and VTK formats. *Pizza.py* can pipe LAMMPS dump files directly into the



Raster3d and RasMol visualization programs. Pizza.py has tools that do interactive 3d OpenGL visualization and one that creates SVG images of dump file snapshots.

### 8.1.3 Run multiple simulations from one input script

This can be done in several ways. See the documentation for individual commands for more details on how these examples work.

If “multiple simulations” means continue a previous simulation for more timesteps, then you simply use the `run` command multiple times. For example, this script

```
units lj
atom_style atomic
read_data data.lj
run 10000
run 10000
run 10000
run 10000
run 10000
```

would run 5 successive simulations of the same system for a total of 50,000 timesteps.

If you wish to run totally different simulations, one after the other, the `clear` command can be used in between them to re-initialize LAMMPS. For example, this script

```
units lj
atom_style atomic
read_data data.lj
run 10000
clear
units lj
atom_style atomic
read_data data.lj.new
run 10000
```

would run 2 independent simulations, one after the other.

For large numbers of independent simulations, you can use `variables` and the `next` and `jump` commands to loop over the same input script multiple times with different settings. For example, this script, named `in.polymer`

```
variable d index run1 run2 run3 run4 run5 run6 run7 run8
shell cd $d
read_data data.polymer
run 10000
shell cd ..
clear
next d
jump in.polymer
```

would run 8 simulations in different directories, using a `data.polymer` file in each directory. The same concept could be used to run the same system at 8 different temperatures, using a temperature variable and storing the output in different log and dump files, for example

```
variable a loop 8
variable t index 0.8 0.85 0.9 0.95 1.0 1.05 1.1 1.15
```

(continues on next page)

(continued from previous page)

```
log log.$a
read data.polymer
velocity all create $t 352839
fix 1 all nvt $t $t 100.0
dump 1 all atom 1000 dump.$a
run 100000
clear
next t
next a
jump in.polymer
```

All of the above examples work whether you are running on 1 or multiple processors, but assumed you are running LAMMPS on a single partition of processors. LAMMPS can be run on multiple partitions via the *-partition command-line switch*.

In the last 2 examples, if LAMMPS were run on 3 partitions, the same scripts could be used if the “index” and “loop” variables were replaced with *universe*-style variables, as described in the *variable* command. Also, the “next t” and “next a” commands would need to be replaced with a single “next a t” command. With these modifications, the 8 simulations of each script would run on the 3 partitions one after the other until all were finished. Initially, 3 simulations would be started simultaneously, one on each partition. When one finished, that partition would then start the fourth simulation, and so forth, until all 8 were completed.

## 8.1.4 Multi-replica simulations

Several commands in LAMMPS run multi-replica simulations, meaning that multiple instances (replicas) of your simulation are run simultaneously, with small amounts of data exchanged between replicas periodically.

These are the relevant commands:

- *hyper* for bond boost hyperdynamics (HD)
- *neb* for nudged elastic band calculations (NEB)
- *neb\_spin* for magnetic nudged elastic band calculations
- *prd* for parallel replica dynamics (PRD)
- *tad* for temperature accelerated dynamics (TAD)
- *temper* for parallel tempering with fixed volume
- *temper/npt* for parallel tempering extended for NPT
- *temper/grem* for parallel tempering with generalized replica exchange (gREM)
- *fix pimd* for path-integral molecular dynamics (PIMD)

NEB is a method for finding transition states and barrier potential energies. HD, PRD, and TAD are methods for performing accelerated dynamics to find and perform infrequent events. Parallel tempering or replica exchange runs different replicas at a series of temperature to facilitate rare-event sampling. PIMD runs different replicas whose individual particles in different replicas are coupled together by springs to model a system of ring-polymers which can represent the quantum nature of atom cores.

These commands can only be used if LAMMPS was built with the REPLICA package. See the *Build package* page for more info.

In all these cases, you must run with one or more processors per replica. The processors assigned to each replica are determined at run-time by using the *-partition command-line switch* to launch LAMMPS on multiple partitions, which in this context are the same as replicas. E.g. these commands:

```
mpirun -np 16 lmp_linux -partition 8x2 -in in.temper
mpirun -np 8 lmp_linux -partition 8x1 -in in.neb
```

would each run 8 replicas, on either 16 or 8 processors. Note the use of the *-in command-line switch* to specify the input script which is required when running in multi-replica mode.

Also note that with MPI installed on a machine (e.g. your desktop), you can run on more (virtual) processors than you have physical processors. Thus the above commands could be run on a single-processor (or few-processor) desktop so that you can run a multi-replica simulation on more replicas than you have physical processors. This is useful for testing and debugging, since with most modern processors and MPI libraries the efficiency of a calculation can severely diminish when oversubscribing processors.

## 8.1.5 Library interface to LAMMPS

As described on the *Build basics* doc page, LAMMPS can be built as a static or shared library, so that it can be called by another code, used in a *coupled manner* with other codes, or driven through a *Python interface*.

At the core of LAMMPS is the LAMMPS class which encapsulates the state of the simulation program through the state of the various class instances that it is composed of. So a calculation using LAMMPS requires to create an instance of the LAMMPS class and then send it (text) commands, either individually or from a file, or perform other operations that modify the state stored inside that instance or drive simulations. This is essentially what the `src/main.cpp` file does as well for the standalone LAMMPS executable with reading commands either from an input file or stdin.

Creating a LAMMPS instance can be done by using C++ code directly or through a C-style interface library to LAMMPS that is provided in the files `src/library.cpp` and `library.h`. This *C language API*, can be used from C and C++, and is also the basis for the *Python* and *Fortran* interfaces or wrappers included in the LAMMPS source code.

The `examples/COUPLE` and `python/examples` directories contain some example programs written in C++, C, Fortran, and Python, which show how a driver code can link to LAMMPS as a library, run LAMMPS on a subset of processors (so the others are available to run some other code concurrently), grab data from LAMMPS, change it, and send it back into LAMMPS.

A detailed documentation of the available APIs and examples of how to use them can be found in the *Programmer Guide* section of this manual.

## 8.1.6 Coupling LAMMPS to other codes

LAMMPS is designed to allow it to be coupled to other codes. For example, a quantum mechanics code might compute forces on a subset of atoms and pass those forces to LAMMPS. Or a continuum finite element (FE) simulation might use atom positions as boundary conditions on FE nodal points, compute a FE solution, and return interpolated forces on MD atoms.

LAMMPS can be coupled to other codes in at least 4 ways. Each has advantages and disadvantages, which you will have to think about in the context of your application.

1. Define a new *fix* command that calls the other code. In this scenario, LAMMPS is the driver code. During timestepping, the *fix* is invoked, and can make library calls to the other code, which has been linked to LAMMPS as a library. This is the way the *LATTE* package, which performs density-functional tight-binding calculations using the *LATTE software* to compute forces, is hooked to LAMMPS. See the *fix latte* command for more details. Also see the *Modify* doc pages for info on how to add a new *fix* to LAMMPS.
2. Define a new LAMMPS command that calls the other code. This is conceptually similar to method (1), but in this case LAMMPS and the other code are on a more equal footing. Note that now the other code is not called during the timestepping of a LAMMPS run, but between runs. The LAMMPS input script can be used to alternate LAMMPS runs with calls to the other code, invoked via the new command. The *run* command facilitates this

with its *every* option, which makes it easy to run a few steps, invoke the command, run a few steps, invoke the command, etc.

In this scenario, the other code can be called as a library, as in 1., or it could be a stand-alone code, invoked by a `system()` call made by the command (assuming your parallel machine allows one or more processors to start up another program). In the latter case the stand-alone code could communicate with LAMMPS through files that the command writes and reads.

See the [Modify command](#) page for info on how to add a new command to LAMMPS.

3. Use LAMMPS as a library called by another code. In this case the other code is the driver and calls LAMMPS as needed. Or a wrapper code could link and call both LAMMPS and another code as libraries. Again, the *run* command has options that allow it to be invoked with minimal overhead (no setup or clean-up) if you wish to do multiple short runs, driven by another program. Details about using the library interface are given in the [library API](#) documentation.
4. Couple LAMMPS with another code in a client/server fashion, using using the [MDI Library](#) developed by the [Molecular Sciences Software Institute \(MolSSI\)](#) to run LAMMPS as either an MDI driver (client) or an MDI engine (server). The MDI driver issues commands to the MDI server to exchange data between them. See the [Howto mdi](#) page for more information about how LAMMPS can operate in either of these modes.

### 8.1.7 Using LAMMPS with the MDI library for code coupling

Client/server coupling of two (or more) codes is where one code is the “client” and sends request messages (data) to one (or more) “server” code(s). A server responds to each request with a reply message (data). This enables two (or more) codes to work in tandem to perform a simulation. In this context, LAMMPS can act as either a client or server code. It does this by using the [MolSSI Driver Interface \(MDI\) library](#), developed by the [Molecular Sciences Software Institute \(MolSSI\)](#), which is supported by the [MDI](#) package.

Alternate methods for code coupling with LAMMPS are described on the [Howto couple](#) doc page.

Some advantages of client/server coupling are that the codes can run as stand-alone executables; they need not be linked together. Thus neither code needs to have a library interface. This also makes it easy to run the two codes on different numbers of processors. If a message protocol (format and content) is defined for a particular kind of simulation, then in principle any code which implements the client-side protocol can be used in tandem with any code which implements the server-side protocol. Neither code needs to know what specific other code it is working with.

In MDI nomenclature, a client code is the “driver”, and a server code is an “engine”. One driver code can communicate with one or more instances of one or more engine codes. Driver and engine codes can be written in any language: C, C++, Fortran, Python, etc.

In addition to allowing driver and engine(s) to run as stand-alone executables, MDI also enables an engine to be a *plugin* to the client code. In this scenario, server code(s) are compiled as shared libraries, and one (or more) instances of the server are instantiated by the driver code. If the driver code runs in parallel, it can split its MPI communicator into multiple sub-communicators, and launch each plugin engine instance on a sub-communicator. Driver processors within that sub-communicator exchange messages with the corresponding engine instance, and can also send MPI messages to other processors in the driver. The driver code can also destroy engine instances and re-instantiate them. LAMMPS can operate as either a stand-alone or plugin MDI engine. When it operates as a driver, it can use either stand-alone or plugin MDI engines.

The way in which an MDI driver communicates with an MDI engine is by making `MDI_Send()` and `MDI_Recv()` calls, which are conceptually similar to `MPI_Send()` and `MPI_Recv()` calls. Each send or receive operation uses a string to identify the command name, and optionally some data, which can be a single value or vector of values of any data type. Inside the MDI library, data is exchanged between the driver and engine via MPI calls or sockets. This a run-time choice by the user.

The *MDI* package provides a *mdi engine* command which enables LAMMPS to operate as an MDI engine. Its doc page explains the variety of standard and custom MDI commands which the LAMMPS engine recognizes and can respond to.

The package also provides a *mdi plugin* command which enables LAMMPS to operate as an MDI driver and load an MDI engine as a plugin library.

The package also has a *fix mdi/qm* `<fix_mdi_qm>` command in which LAMMPS operates as an MDI driver in conjunction with a quantum mechanics code as an MDI engine. The `post_force()` method of the `fix_mdi_qm.cpp` file shows how a driver issues MDI commands to another code. This command can be used to couple to an MDI engine which is either a stand-alone code or a plugin library.

As explained on the *fix mdi/qm* `<fix_mdi_qm>` command doc page, it can be used to perform *ab initio* MD simulations or energy minimizations, or to evaluate the quantum energy and forces for a series of independent systems. The `examples/mdi` directory has example input scripts for all of these use cases.

The `examples/mdi` directory contains Python scripts and LAMMPS input script which use LAMMPS as either an MDI driver or engine or both. Currently, 5 example use cases are provided:

- Run *ab initio* MD (AIMD) using 2 instances of LAMMPS. As a driver LAMMPS performs the timestepping in either NVE or NPT mode. As an engine, LAMMPS computes forces and is a surrogate for a quantum code.
- As a driver, LAMMPS runs an MD simulation. Every N steps it passes the current snapshot to an MDI engine to evaluate the energy, virial, and peratom forces. As the engine LAMMPS is a surrogate for a quantum code.
- As a driver, LAMMPS loops over a series of data files and passes the configuration to an MDI engine to evaluate the energy, virial, and peratom forces. As the engine LAMMPS is a surrogate for a quantum code.
- A Python script driver invokes a sequence of unrelated LAMMPS calculations. Calculations can be single-point energy/force evaluations, MD runs, or energy minimizations.
- Run AIMD with a Python driver code and 2 LAMMPS instances as engines. The first LAMMPS instance performs MD timestepping. The second LAMMPS instance acts as a surrogate QM code to compute forces.

Note that in any of these example where LAMMPS is used as an engine, an actual QM code (which supports MDI) could be used in its place, without modifying the input scripts or launch commands, except to specify the name of the QM code.

The `examples/mdi/Run.sh` file illustrates how to launch both driver and engine codes so that they communicate using the MDI library via either MPI or sockets. Or using the engine as a stand-alone code or plugin library.

Currently there are at least two quantum DFT codes which have direct MDI support, [Quantum ESPRESSO \(QE\)](#) and [INQ](#). There are also several QM codes which have indirect support through QCEngine or i-PI. The former means they require a wrapper program (QCEngine) with MDI support which writes/read files to pass data to the quantum code itself. The list of QCEngine-supported and i-PI-supported quantum codes is on the [MDI webpage](#).

Here is how to build QE as a stand-alone `pw.x` file which can be used in stand-alone mode:

```
% git clone --branch mdi_plugin https://github.com/MolSSI-MDI/q-e.git <base_path>/q-e
% build the executable pw.x, following the `QE build guide <https://gitlab.com/QEF/q-e/-/
wikis/Developers/CMake-build-system>` _
```

Here is how to build QE as a shared library which can be used in plugin mode, which results in a `libqemdi.so` file in `<base_path>/q-e/MDI/src`:

```
% git clone --branch mdi_plugin https://github.com/MolSSI-MDI/q-e.git <base_path>/q-e
% cd <base_path>/q-e
% ./configure --enable-parallel --enable-openmp --enable-shared FFLAGS="-fPIC" FCFLAGS="-
↳fPIC" CFLAGS="-fPIC" foxflags="-fPIC" try_foxflags="-fPIC"
% make -j 4 mdi
```

INQ cannot be built as a stand-alone code; it is by design a library. Here is how to build INQ as a shared library which can be used in plugin mode, which results in a `libinqmdi.so` file in `<base_path>/inq/build/examples`:

```
% git clone --branch mdi --recurse-submodules https://gitlab.com/taylor-a-barnes/inq.git
↳<base_path>/inq
% cd <base_path>/inq
% mkdir -p build
% cd build
% ../configure --prefix=<install_path>/install
% make -j 4
% make install
```

### 8.1.8 Bonded particle models

The BPM package implements bonded particle models which can be used to simulate mesoscale solids. Solids are constructed as a collection of particles which each represent a coarse-grained region of space much larger than the atomistic scale. Particles within a solid region are then connected by a network of bonds to provide solid elasticity.

Unlike traditional bonds in molecular dynamics, the equilibrium bond length can vary between bonds. Bonds store the reference state. This includes setting the equilibrium length equal to the initial distance between the two particles but can also include data on the bond orientation for rotational models. This produces a stress free initial state. Furthermore, bonds are allowed to break under large strains producing fracture. The `examples/bpm` directory has sample input scripts for simulations of the fragmentation of an impacted plate and the pouring of extended, elastic bodies.

---

Bonds can be created using a *read data* or *create bonds* command. Alternatively, a *molecule* template with bonds can be used with *fix deposit* or *fix pour* to create solid grains.

In this implementation, bonds store their reference state when they are first computed in the setup of the first simulation run. Data is then preserved across run commands and is written to *binary restart files* such that restarting the system will not reset the reference state of a bond. Bonds that are created midway into a run, such as those created by pouring grains using *fix pour*, are initialized on that timestep.

As bonds can be broken between neighbor list builds, the *special\_bonds* command works differently for BPM bond styles. There are two possible settings which determine how pair interactions work between bonded particles. First, one can turn off all pair interactions between bonded particles. Unlike *bond quartic*, this is not done by subtracting pair forces during the bond computation but rather by dynamically updating the special bond list. This is the default behavior of BPM bond styles and is done by updating the 1-2 special bond list as bonds break. To do this, LAMMPS requires *newton* bond off such that all processors containing an atom know when a bond breaks. Additionally, one must do either (A) or (B).

A) Use the following special bond settings

```
special_bonds lj 0 1 1 coul 1 1 1
```

These settings accomplish two goals. First, they turn off 1-3 and 1-4 special bond lists, which are not currently supported for BPMs. As BPMs often have dense bond networks, generating 1-3 and 1-4 special bond lists is expensive. By setting the `lj` weight for 1-2 bonds to zero, this turns off pairwise interactions. Even though there

are no charges in BPM models, setting a nonzero coul weight for 1-2 bonds ensures all bonded neighbors are still included in the neighbor list in case bonds break between neighbor list builds.

- B) Alternatively, one can simply overlay pair interactions such that all bonded particles also feel pair interactions. This can be accomplished by using the *overlay/pair* keyword present in all bpm bond styles and by using the following special bond settings

```
special_bonds lj/coul 1 1 1
```

See the *Howto* page on broken bonds for more information.

Currently there are two types of bonds included in the BPM package. The first bond style, *bond bpm/spring*, only applies pairwise, central body forces. Point particles must have *bond atom style* and may be thought of as nodes in a spring network. Alternatively, the second bond style, *bond bpm/rotational*, resolves tangential forces and torques arising with the shearing, bending, and twisting of the bond due to rotation or displacement of particles. Particles are similar to those used in the *granular package*, *atom style sphere*. However, they must also track the current orientation of particles and store bonds and therefore use a *bpm/sphere atom style*. This also requires a unique integrator *fix nve/bpm/sphere* which numerically integrates orientation similar to *fix nve/asphere*.

To monitor the fracture of bonds in the system, all BPM bond styles have the ability to record instances of bond breakage to output using the *dump local* command. Additionally, one can use *compute nbond/atom* to tally the current number of bonds per atom.

In addition to bond styles, a new pair style *pair bpm/spring* was added to accompany the bpm/spring bond style. This pair style is simply a hookean repulsion with similar velocity damping as its sister bond style.

While LAMMPS has many utilities to create and delete bonds, *only* the following are currently compatible with BPM bond styles:

- *create\_bonds*
- *delete\_bonds*
- *fix bond/create*
- *fix bond/break*
- *fix bond/swap*

Note *create\_bonds* requires certain special\_bonds settings. To subtract pair interactions, one will need to switch between different special\_bonds settings in the input script. An example is found in examples/bpm/impact.

### 8.1.9 Broken Bonds

Typically, bond interactions persist for the duration of a simulation in LAMMPS. However, there are some exceptions that allow for bonds to break including the *quartic bond style* and the bond styles in the *BPM package* which contains the *bpm/spring* and *bpm/rotational* bond styles. In these cases, a bond can be broken if it is stretched beyond a user-defined threshold. LAMMPS accomplishes this by setting the bond type to zero such that the bond force is no longer computed.

Users are normally able to weight the contribution of pair forces to atoms that are bonded using the *special\_bonds command*. When bonds break, this is not always the case. For the quartic bond style, pair forces are always turned off between bonded particles. LAMMPS does this via a computational sleight-of-hand. It subtracts the pairwise interaction as part of the bond computation. When the bond breaks, the subtraction stops. For this to work, the pairwise interaction must always be computed by the *pair\_style* command, whether the bond is broken or not. This means that *special\_bonds*

must be set to 1,1,1. After the bond breaks, the pairwise interaction between the two atoms is turned on, since they are no longer bonded.

In the BPM package, one can either turn off all pair interactions between bonded particles or leave them on, overlaying pair forces on top of bond forces. To remove pair forces, the special bond list is dynamically updated. More details can be found on the [Howto BPM](#) page.

Bonds can also be broken by fixes which change bond topology, including [fix bond/break](#) and [fix bond/react](#). These fixes will automatically trigger a rebuild of the neighbor list and update special bond data structures when bonds are broken.

Note that when bonds are dumped to a file via the [dump local](#) command, bonds with type 0 are not included. The [delete\\_bonds](#) command can also be used to query the status of broken bonds or permanently delete them, e.g.:

```
delete_bonds all stats
delete_bonds all bond 0 remove
```

The compute [nbond/atom](#) can also be used to tally the current number of bonds per atom, excluding broken bonds.

## 8.2 Settings howto

### 8.2.1 2d simulations

Use the [dimension](#) command to specify a 2d simulation.

Make the simulation box periodic in z via the [boundary](#) command. This is the default.

If using the [create\\_box](#) command to define a simulation box, set the z dimensions narrow, but finite, so that the [create\\_atoms](#) command will fill the 3d simulation box with a single z plane of atoms - e.g.

```
create_box 1 -10 10 -10 10 -0.25 0.25
```

If using the [read data](#) command to read in a file of atom coordinates, set the “zlo zhi” values to be finite but narrow, similar to the [create\\_box](#) command settings just described. For each atom in the file, assign a z coordinate so it falls inside the z-boundaries of the box - e.g. 0.0.

Use the [fix enforce2d](#) command as the last defined fix to insure that the z-components of velocities and forces are zeroed out every timestep. The reason to make it the last fix is so that any forces induced by other fixes will be zeroed out.

Many of the example input scripts included in the LAMMPS distribution are for 2d models.

---

**Note:** Some models in LAMMPS treat particles as finite-size spheres, as opposed to point particles. See the [atom\\_style sphere](#) and [fix nve/sphere](#) commands for details. By default, for 2d simulations, such particles will still be modeled as 3d spheres, not 2d discs (circles), meaning their moment of inertia will be that of a sphere. If you wish to model them as 2d discs, see the [set density/disc](#) command and the [disc](#) option for the [fix nve/sphere](#), [fix nvt/sphere](#), [fix nph/sphere](#), [fix npt/sphere](#) commands.

---



## 8.2.2 Type labels

New in version 15Sep2022.

Each atom in LAMMPS has an associated numeric atom type. Similarly, each bond, angle, dihedral, and improper is assigned a bond type, angle type, and so on. The primary use of these types is to map potential (force field) parameters to the interactions of the atom, bond, angle, dihedral, and improper.

By default, type values are entered as integers from 1 to Ntypes wherever they appear in LAMMPS input or output files. The total number Ntypes for each interaction is “locked in” when the simulation box is created.

A recent addition to LAMMPS is the option to use strings - referred to as type labels - as an alternative. Using type labels instead of numeric types can be advantageous in various scenarios. For example, type labels can make inputs more readable and generic (i.e. usable through the *include command* for different systems with different numerical values assigned to types. This generality also applies to other inputs like data files read by *read\_data* or molecule template files read by the *molecule* command. See below for a list of other commands that can use type labels in different ways.

LAMMPS will *internally* continue to use numeric types, which means that many previous restrictions still apply. For example, the total number of types is locked in when creating the simulation box, and potential parameters for each type must be provided even if not used by any interactions.

A collection of type labels for all type-kinds (atom types, bond types, etc.) is stored as a “label map” which is simply a list of numeric types and their associated type labels. Within a type-kind, each type label must be unique. It can be assigned to only one numeric type. To read and write type labels to data files for a given type-kind, *all* associated numeric types need have a type label assigned. Partial maps can be saved with the *labelmap write* command and read back with the *include* command.

Valid type labels can contain most ASCII characters, but cannot start with a number, a ‘#’, or a ‘\*’. Also, labels must not contain whitespace characters. When using the *labelmap command* in the LAMMPS input, if certain characters appear in the type label, such as the single (‘) or double (‘’) quote or the ‘#’ character, the label must be put in either double, single, or triple (‘‘‘’) quotes. Triple quotes allow for the most generic type label strings, but they require to have a leading and trailing blank space. When defining type labels the blanks will be ignored. Example:

```
labelmap angle 1 "" " C1'-C2"-C3# ""
```

This command will map the string `C1'-C2"-C3#` to the angle type 1.

There are two ways to define label maps. One is via the *labelmap* command. The other is via the *read\_data* command. A data file can have sections such as *Atom Type Labels*, *Bond Type Labels*, etc., which assign type labels to numeric types. The label map can be written out to data files by the *write\_data* command. This map is also written to and read from restart files, by the *write\_restart* and *read\_restart* commands.

### Use of type labels in LAMMPS input or output

Many LAMMPS input script commands that take a numeric type as an argument can use the associated type label instead. If a type label is not defined for a particular numeric type, only its numeric type can be used.

This example assigns labels to the atom types, and then uses the type labels to redefine the pair coefficients.

```
pair_coeff 1 2 1.0 1.0          # numeric types
labelmap atom 1 C 2 H
pair_coeff C H 1.0 1.0        # type labels
```

Adding support for type labels to various commands is an ongoing project. If an input script command (or a section in a file read by a command) allows substituting a type label for a numeric type argument, it will be explicitly mentioned in that command's documentation page.

As a temporary measure, input script commands can take advantage of variables and how they can be expanded during processing of the input. The variables can use functions that will translate type label strings to their respective number as defined in the current label map. See the *variable* command for details.

For example, here is how the `pair_coeff` command could be used with type labels if it did not yet support them, either with an explicit variable command or an implicit variable used in the `pair_coeff` command.

```
labelmap atom 1 C 2 H
variable atom1 equal label2type(atom,C)
variable atom2 equal label2type(atom,H)
pair_coeff ${atom1} ${atom2} 1.0 1.0
```

```
labelmap atom 1 C 2 H
pair_coeff $(label2type(atom,C)) $(label2type(atom,H)) 80.0 1.2
```

---

### Commands that can use label types

Any workflow that involves reading multiple data files, molecule templates or a combination of the two can be streamlined by using type labels instead of numeric types, because types are automatically synced between the files. The creation of simulation-ready reaction templates for *fix bond/react* is much simpler when using type labels, and results in templates that can be used without modification in multiple simulations or different systems.

### 8.2.3 Triclinic (non-orthogonal) simulation boxes

By default, LAMMPS uses an orthogonal simulation box to encompass the particles. The *boundary* command sets the boundary conditions of the box (periodic, non-periodic, etc). The orthogonal box has its “origin” at (xlo,ylo,zlo) and is defined by 3 edge vectors starting from the origin given by  $\mathbf{a} = (xhi-xlo,0,0)$ ;  $\mathbf{b} = (0,yhi-ylo,0)$ ;  $\mathbf{c} = (0,0,zhi-zlo)$ . The 6 parameters (xlo,xhi,ylo,yhi,zlo,zhi) are defined at the time the simulation box is created, e.g. by the *create\_box* or *read\_data* or *read\_restart* commands. Additionally, LAMMPS defines box size parameters lx,ly,lz where lx = xhi-xlo, and similarly in the y and z dimensions. The 6 parameters, as well as lx,ly,lz, can be output via the *thermo\_style custom* command.

LAMMPS also allows simulations to be performed in triclinic (non-orthogonal) simulation boxes shaped as a parallelepiped with triclinic symmetry. The parallelepiped has its “origin” at (xlo,ylo,zlo) and is defined by 3 edge vectors starting from the origin given by  $\mathbf{a} = (xhi-xlo,0,0)$ ;  $\mathbf{b} = (xy,yhi-ylo,0)$ ;  $\mathbf{c} = (xz,yz,zhi-zlo)$ . xy,xz,yz can be 0.0 or positive or negative values and are called “tilt factors” because they are the amount of displacement applied to faces of an originally orthogonal box to transform it into the parallelepiped. In LAMMPS the triclinic simulation box edge vectors  $\mathbf{a}$ ,  $\mathbf{b}$ , and  $\mathbf{c}$  cannot be arbitrary vectors. As indicated,  $\mathbf{a}$  must lie on the positive x axis.  $\mathbf{b}$  must lie in the xy plane, with strictly positive y component.  $\mathbf{c}$  may have any orientation with strictly positive z component. The requirement that  $\mathbf{a}$ ,  $\mathbf{b}$ , and  $\mathbf{c}$  have strictly positive x, y, and z components, respectively, ensures that  $\mathbf{a}$ ,  $\mathbf{b}$ , and  $\mathbf{c}$  form a complete right-handed basis. These restrictions impose no loss of generality, since it is possible to rotate/invert any set of 3 crystal basis vectors so that they conform to the restrictions.

For example, assume that the 3 vectors  $\mathbf{A}, \mathbf{B}, \mathbf{C}$  are the edge vectors of a general parallelepiped, where there is no restriction on  $\mathbf{A}, \mathbf{B}, \mathbf{C}$  other than they form a complete right-handed basis i.e.  $\mathbf{A} \times \mathbf{B} \cdot \mathbf{C} > 0$ . The equivalent LAMMPS

$\mathbf{a}, \mathbf{b}, \mathbf{c}$  are a linear rotation of  $\mathbf{A}, \mathbf{B}$ , and  $\mathbf{C}$  and can be computed as follows:

$$\begin{aligned}
 (\mathbf{a} \ \mathbf{b} \ \mathbf{c}) &= \begin{pmatrix} a_x & b_x & c_x \\ 0 & b_y & c_y \\ 0 & 0 & c_z \end{pmatrix} \\
 a_x &= A \\
 b_x &= \mathbf{B} \cdot \hat{\mathbf{A}} = B \cos \gamma \\
 b_y &= |\hat{\mathbf{A}} \times \mathbf{B}| = B \sin \gamma = \sqrt{B^2 - b_x^2} \\
 c_x &= \mathbf{C} \cdot \hat{\mathbf{A}} = C \cos \beta \\
 c_y &= \mathbf{C} \cdot (\widehat{\mathbf{A} \times \mathbf{B}}) \times \hat{\mathbf{A}} = \frac{\mathbf{B} \cdot \mathbf{C} - b_x c_x}{b_y} \\
 c_z &= |\mathbf{C} \cdot (\widehat{\mathbf{A} \times \mathbf{B}})| = \sqrt{C^2 - c_x^2 - c_y^2}
 \end{aligned}$$

where  $A = |\mathbf{A}|$  indicates the scalar length of  $\mathbf{A}$ . The hat symbol (^) indicates the corresponding unit vector.  $\beta$  and  $\gamma$  are angles between the vectors described below. Note that by construction,  $\mathbf{a}, \mathbf{b}$ , and  $\mathbf{c}$  have strictly positive x, y, and z components, respectively. If it should happen that  $\mathbf{A}, \mathbf{B}$ , and  $\mathbf{C}$  form a left-handed basis, then the above equations are not valid for  $\mathbf{c}$ . In this case, it is necessary to first apply an inversion. This can be achieved by interchanging two basis vectors or by changing the sign of one of them.

For consistency, the same rotation/inversion applied to the basis vectors must also be applied to atom positions, velocities, and any other vector quantities. This can be conveniently achieved by first converting to fractional coordinates in the old basis and then converting to distance coordinates in the new basis. The transformation is given by the following equation:

$$\mathbf{x} = (\mathbf{a} \ \mathbf{b} \ \mathbf{c}) \cdot \frac{1}{V} \begin{pmatrix} \mathbf{B} \times \mathbf{C} \\ \mathbf{C} \times \mathbf{A} \\ \mathbf{A} \times \mathbf{B} \end{pmatrix} \cdot \mathbf{X}$$

where  $V$  is the volume of the box,  $\mathbf{X}$  is the original vector quantity and  $\mathbf{x}$  is the vector in the LAMMPS basis.

There is no requirement that a triclinic box be periodic in any dimension, though it typically should be in at least the second dimension of the tilt (y in xy) if you want to enforce a shift in periodic boundary conditions across that boundary. Some commands that work with triclinic boxes, e.g. the *fix deform* and *fix npt* commands, require periodicity or non-shrink-wrap boundary conditions in specific dimensions. See the command doc pages for details.

The 9 parameters (xlo,xhi,ylo,yhi,zlo,zhi,xy,xz,yz) are defined at the time the simulation box is created. This happens in one of 3 ways. If the *create\_box* command is used with a region of style *prism*, then a triclinic box is setup. See the *region* command for details. If the *read\_data* command is used to define the simulation box, and the header of the data file contains a line with the “xy xz yz” keyword, then a triclinic box is setup. See the *read\_data* command for details. Finally, if the *read\_restart* command reads a restart file which was written from a simulation using a triclinic box, then a triclinic box will be setup for the restarted simulation.

Note that you can define a triclinic box with all 3 tilt factors = 0.0, so that it is initially orthogonal. This is necessary if the box will become non-orthogonal, e.g. due to the *fix npt* or *fix deform* commands. Alternatively, you can use the *change\_box* command to convert a simulation box from orthogonal to triclinic and vice versa.

As with orthogonal boxes, LAMMPS defines triclinic box size parameters lx,ly,lz where lx = xhi-xlo, and similarly in the y and z dimensions. The 9 parameters, as well as lx,ly,lz, can be output via the *thermo\_style custom* command.

To avoid extremely tilted boxes (which would be computationally inefficient), LAMMPS normally requires that no tilt factor can skew the box more than half the distance of the parallel box length, which is the first dimension in the tilt factor (x for xz). This is required both when the simulation box is created, e.g. via the *create\_box* or *read\_data* commands, as well as when the box shape changes dynamically during a simulation, e.g. via the *fix deform* or *fix npt* commands.

For example, if  $xlo = 2$  and  $xhi = 12$ , then the  $x$  box length is 10 and the  $xy$  tilt factor must be between -5 and 5. Similarly, both  $xz$  and  $yz$  must be between  $-(xhi-xlo)/2$  and  $+(yhi-ylo)/2$ . Note that this is not a limitation, since if the maximum tilt factor is 5 (as in this example), then configurations with  $tilt = \dots, -15, -5, 5, 15, 25, \dots$  are geometrically all equivalent. If the box tilt exceeds this limit during a dynamics run (e.g. via the *fix deform* command), then the box is “flipped” to an equivalent shape with a tilt factor within the bounds, so the run can continue. See the *fix deform* page for further details.

One exception to this rule is if the first dimension in the tilt factor ( $x$  for  $xy$ ) is non-periodic. In that case, the limits on the tilt factor are not enforced, since flipping the box in that dimension does not change the atom positions due to non-periodicity. In this mode, if you tilt the system to extreme angles, the simulation will simply become inefficient, due to the highly skewed simulation box.

The limitation on not creating a simulation box with a tilt factor skewing the box more than half the distance of the parallel box length can be overridden via the *box* command. Setting the *tilt* keyword to *large* allows any tilt factors to be specified.

Box flips that may occur using the *fix deform* or *fix npt* commands can be turned off using the *flip no* option with either of the commands.

Note that if a simulation box has a large tilt factor, LAMMPS will run less efficiently, due to the large volume of communication needed to acquire ghost atoms around a processor’s irregular-shaped sub-domain. For extreme values of tilt, LAMMPS may also lose atoms and generate an error.

Triclinic crystal structures are often defined using three lattice constants  $a$ ,  $b$ , and  $c$ , and three angles  $\alpha$ ,  $\beta$ , and  $\gamma$ . Note that in this nomenclature, the  $a$ ,  $b$ , and  $c$  lattice constants are the scalar lengths of the edge vectors **a**, **b**, and **c** defined above. The relationship between these 6 quantities ( $a$ ,  $b$ ,  $c$ ,  $\alpha$ ,  $\beta$ ,  $\gamma$ ) and the LAMMPS box sizes ( $lx, ly, lz$ ) = ( $xhi-xlo, yhi-ylo, zhi-zlo$ ) and tilt factors ( $xy, xz, yz$ ) is as follows:

$$\begin{aligned} a &= lx \\ b^2 &= ly^2 + xy^2 \\ c^2 &= lz^2 + xz^2 + yz^2 \\ \cos \alpha &= \frac{xy * xz + ly * yz}{b * c} \\ \cos \beta &= \frac{xz}{c} \\ \cos \gamma &= \frac{xy}{b} \end{aligned}$$

The inverse relationship can be written as follows:

$$\begin{aligned} lx &= a \\ xy &= b \cos \gamma \\ xz &= c \cos \beta \\ ly^2 &= b^2 - xy^2 \\ yz &= \frac{b * c \cos \alpha - xy * xz}{ly} \\ lz^2 &= c^2 - xz^2 - yz^2 \end{aligned}$$

The values of  $a$ ,  $b$ ,  $c$ ,  $\alpha$ ,  $\beta$ , and  $\gamma$  can be printed out or accessed by computes using the *thermo\_style custom* keywords *cella*, *cellb*, *cellc*, *cellalpha*, *cellbeta*, *cellgamma*, respectively.

As discussed on the *dump* command doc page, when the BOX BOUNDS for a snapshot is written to a dump file for a triclinic box, an orthogonal bounding box which encloses the triclinic simulation box is output, along with the 3 tilt factors ( $xy$ ,  $xz$ ,  $yz$ ) of the triclinic box, formatted as follows:

```
ITEM: BOX BOUNDS xy xz yz
xlo_bound xhi_bound xy
ylo_bound yhi_bound xz
zlo_bound zhi_bound yz
```

This bounding box is convenient for many visualization programs and is calculated from the 9 triclinic box parameters (xlo,xhi,ylo,yhi,zlo,zhi,xy,xz,yz) as follows:

```
xlo_bound = xlo + MIN(0.0, xy, xz, xy+xz)
xhi_bound = xhi + MAX(0.0, xy, xz, xy+xz)
ylo_bound = ylo + MIN(0.0, yz)
yhi_bound = yhi + MAX(0.0, yz)
zlo_bound = zlo
zhi_bound = zhi
```

These formulas can be inverted if you need to convert the bounding box back into the triclinic box parameters, e.g.  $xlo = xlo\_bound - \text{MIN}(0.0, xy, xz, xy+xz)$ .

One use of triclinic simulation boxes is to model solid-state crystals with triclinic symmetry. The *lattice* command can be used with non-orthogonal basis vectors to define a lattice that will tile a triclinic simulation box via the *create\_atoms* command.

A second use is to run Parrinello-Rahman dynamics via the *fix npt* command, which will adjust the xy, xz, yz tilt factors to compensate for off-diagonal components of the pressure tensor. The analog for an *energy minimization* is the *fix box/relax* command.

A third use is to shear a bulk solid to study the response of the material. The *fix deform* command can be used for this purpose. It allows dynamic control of the xy, xz, yz tilt factors as a simulation runs. This is discussed in the next section on non-equilibrium MD (NEMD) simulations.

## 8.2.4 Thermostats

Thermostatting means controlling the temperature of particles in an MD simulation. *Barostatting* means controlling the pressure. Since the pressure includes a kinetic component due to particle velocities, both these operations require calculation of the temperature. Typically a target temperature (T) and/or pressure (P) is specified by the user, and the thermostat or barostat attempts to equilibrate the system to the requested T and/or P.

Thermostatting in LAMMPS is performed by *fixes*, or in one case by a pair style. Several thermostatting fixes are available: Nose-Hoover (nvt), Berendsen, CSVr, Langevin, and direct rescaling (temp/rescale). Dissipative particle dynamics (DPD) thermostatting can be invoked via the *dpd/tstat* pair style:

- *fix nvt*
- *fix nvt/sphere*
- *fix nvt/asphere*
- *fix nvt/sllod*
- *fix temp/berendsen*
- *fix temp/csvr*
- *fix langevin*
- *fix temp/rescale*
- *pair\_style dpd/tstat*

*Fix nvt* only thermostats the translational velocity of particles. *Fix nvt/sllod* also does this, except that it subtracts out a velocity bias due to a deforming box and integrates the SLLOD equations of motion. See the *Howto nemd* page for further details. *Fix nvt/sphere* and *fix nvt/asphere* thermostat not only translation velocities but also rotational velocities for spherical and aspherical particles.

---

**Note:** A recent (2017) book by (*Daivis and Todd*) discusses use of the SLLOD method and non-equilibrium MD (NEMD) thermostating generally, for both simple and complex fluids, e.g. molecular systems. The latter can be tricky to do correctly.

---

DPD thermostating alters pairwise interactions in a manner analogous to the per-particle thermostating of *fix langevin*.

Any of the thermostating fixes can be instructed to use custom temperature computes that remove bias which has two effects: first, the current calculated temperature, which is compared to the requested target temperature, is calculated with the velocity bias removed; second, the thermostat adjusts only the thermal temperature component of the particle's velocities, which are the velocities with the bias removed. The removed bias is then added back to the adjusted velocities. See the doc pages for the individual fixes and for the *fix\_modify* command for instructions on how to assign a temperature compute to a thermostating fix.

For example, you can apply a thermostat only to atoms in a spatial region by using it in conjunction with *compute temp/region*. Or you can apply a thermostat to only the x and z components of velocity by using it with *compute temp/partial*. Of you could thermostat only the thermal temperature of a streaming flow of particles without affecting the streaming velocity, by using *compute temp/profile*.

Below is a list of custom temperature computes that can be used like that:

- *compute temp/asphere command*
- *compute temp/body command*
- *compute temp/chunk command*
- *compute temp/com command*
- *compute temp/deform command*
- *compute temp/partial command*
- *compute temp/profile command*
- *compute temp/ramp command*
- *compute temp/region command*
- *compute temp/rotate command*
- *compute temp/sphere command*

---

**Note:** Only the nvt fixes perform time integration, meaning they update the velocities and positions of particles due to forces and velocities respectively. The other thermostat fixes only adjust velocities; they do NOT perform time integration updates. Thus they should be used in conjunction with a constant NVE integration fix such as these:

---

- *fix nve*
- *fix nve/sphere*
- *fix nve/asphere*

Thermodynamic output, which can be setup via the *thermo\_style* command, often includes temperature values. As explained on the page for the *thermo\_style* command, the default temperature is setup by the thermo command itself. It is NOT the temperature associated with any thermostating fix you have defined or with any compute you have defined

that calculates a temperature. The doc pages for the thermostating fixes explain the ID of the temperature compute they create. Thus if you want to view these temperatures, you need to specify them explicitly via the *thermo\_style custom* command. Or you can use the *thermo\_modify* command to re-define what temperature compute is used for default thermodynamic output.

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**(Daivis and Todd)** Daivis and Todd, Nonequilibrium Molecular Dynamics (book), Cambridge University Press, <https://doi.org/10.1017/9781139017848>, (2017).

## 8.2.5 Barostats

Barostatting means controlling the pressure in an MD simulation. *Thermostating* means controlling the temperature of the particles. Since the pressure includes a kinetic component due to particle velocities, both these operations require calculation of the temperature. Typically a target temperature (T) and/or pressure (P) is specified by the user, and the thermostat or barostat attempts to equilibrate the system to the requested T and/or P.

Barostatting in LAMMPS is performed by *fixes*. Two barostatting methods are currently available: Nose-Hoover (*npt* and *nph*) and Berendsen:

- *fix npt*
- *fix npt/sphere*
- *fix npt/asphere*
- *fix nph*
- *fix press/berendsen*

The *fix npt* commands include a Nose-Hoover thermostat and barostat. *Fix nph* is just a Nose/Hoover barostat; it does no thermostating. Both *fix nph* and *fix press/berendsen* can be used in conjunction with any of the thermostating fixes.

As with the *thermostats*, *fix npt* and *fix nph* only use translational motion of the particles in computing T and P and performing thermo/barostatting. *Fix npt/sphere* and *fix npt/asphere* thermo/barostat using not only translation velocities but also rotational velocities for spherical and aspherical particles.

All of the barostatting fixes use the *compute pressure* compute to calculate a current pressure. By default, this compute is created with a simple *compute temp* (see the last argument of the *compute pressure* command), which is used to calculate the kinetic component of the pressure. The barostatting fixes can also use temperature computes that remove bias for the purpose of computing the kinetic component which contributes to the current pressure. See the doc pages for the individual fixes and for the *fix\_modify* command for instructions on how to assign a temperature or pressure compute to a barostatting fix.

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**Note:** As with the thermostats, the Nose/Hoover methods (*fix npt* and *fix nph*) perform time integration. *Fix press/berendsen* does NOT, so it should be used with one of the constant NVE fixes or with one of the NVT fixes.

---

Thermodynamic output, which can be setup via the *thermo\_style* command, often includes pressure values. As explained on the page for the *thermo\_style* command, the default pressure is setup by the thermo command itself. It is NOT the pressure associated with any barostatting fix you have defined or with any compute you have defined that calculates a pressure. The doc pages for the barostatting fixes explain the ID of the pressure compute they create. Thus if you want to view these pressures, you need to specify them explicitly via the *thermo\_style custom* command. Or you can use the *thermo\_modify* command to re-define what pressure compute is used for default thermodynamic output.

## 8.2.6 Walls

Walls in an MD simulation are typically used to bound particle motion, i.e. to serve as a boundary condition.

Walls in LAMMPS can be of rough (made of particles) or idealized surfaces. Ideal walls can be smooth, generating forces only in the normal direction, or frictional, generating forces also in the tangential direction.

Rough walls, built of particles, can be created in various ways. The particles themselves can be generated like any other particle, via the *lattice* and *create\_atoms* commands, or read in via the *read\_data* command.

Their motion can be constrained by many different commands, so that they do not move at all, move together as a group at constant velocity or in response to a net force acting on them, move in a prescribed fashion (e.g. rotate around a point), etc. Note that if a time integration fix like *fix nve* or *fix nvt* is not used with the group that contains wall particles, their positions and velocities will not be updated.

- *fix aveforce* - set force on particles to average value, so they move together
- *fix setforce* - set force on particles to a value, e.g. 0.0
- *fix freeze* - freeze particles for use as granular walls
- *fix nve/noforce* - advect particles by their velocity, but without force
- *fix move* - prescribe motion of particles by a linear velocity, oscillation, rotation, variable

The *fix move* command offers the most generality, since the motion of individual particles can be specified with *variable* formula which depends on time and/or the particle position.

For rough walls, it may be useful to turn off pairwise interactions between wall particles via the *neigh\_modify exclude* command.

Rough walls can also be created by specifying frozen particles that do not move and do not interact with mobile particles, and then tethering other particles to the fixed particles, via a *bond*. The bonded particles do interact with other mobile particles.

Idealized walls can be specified via several *fix* commands. *Fix wall/gran* creates frictional walls for use with granular particles; all the other commands create smooth walls.

- *fix wall/reflect* - reflective flat walls
- *fix wall/lj93* - flat walls, with Lennard-Jones 9/3 potential
- *fix wall/lj126* - flat walls, with Lennard-Jones 12/6 potential
- *fix wall/colloid* - flat walls, with *pair\_style colloid* potential
- *fix wall/harmonic* - flat walls, with repulsive harmonic spring potential
- *fix wall/morse* - flat walls, with Morse potential
- *fix wall/region* - use region surface as wall
- *fix wall/gran* - flat or curved walls with *pair\_style granular* potential

The *lj93*, *lj126*, *colloid*, *harmonic*, and *morse* styles all allow the flat walls to move with a constant velocity, or oscillate in time. The *fix wall/region* command offers the most generality, since the region surface is treated as a wall, and the geometry of the region can be a simple primitive volume (e.g. a sphere, or cube, or plane), or a complex volume made from the union and intersection of primitive volumes. *Regions* can also specify a volume “interior” or “exterior” to the specified primitive shape or *union* or *intersection*. *Regions* can also be “dynamic” meaning they move with constant velocity, oscillate, or rotate.

The only frictional idealized walls currently in LAMMPS are flat or curved surfaces specified by the *fix wall/gran* command. At some point we plan to allow region surfaces to be used as frictional walls, as well as triangulated surfaces.



## 8.2.7 NEMD simulations

Non-equilibrium molecular dynamics or NEMD simulations are typically used to measure a fluid's rheological properties such as viscosity. In LAMMPS, such simulations can be performed by first setting up a non-orthogonal simulation box (see the preceding Howto section).

A shear strain can be applied to the simulation box at a desired strain rate by using the *fix deform* command. The *fix nvt/sllod* command can be used to thermostat the sheared fluid and integrate the SLLOD equations of motion for the system. Fix nvt/sllod uses *compute temp/deform* to compute a thermal temperature by subtracting out the streaming velocity of the shearing atoms. The velocity profile or other properties of the fluid can be monitored via the *fix ave/chunk* command.

---

**Note:** A recent (2017) book by (*Daivis and Todd*) discusses use of the SLLOD method and non-equilibrium MD (NEMD) thermostating generally, for both simple and complex fluids, e.g. molecular systems. The latter can be tricky to do correctly.

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As discussed in the previous section on non-orthogonal simulation boxes, the amount of tilt or skew that can be applied is limited by LAMMPS for computational efficiency to be 1/2 of the parallel box length. However, *fix deform* can continuously strain a box by an arbitrary amount. As discussed in the *fix deform* command, when the tilt value reaches a limit, the box is flipped to the opposite limit which is an equivalent tiling of periodic space. The strain rate can then continue to change as before. In a long NEMD simulation these box re-shaping events may occur many times.

In a NEMD simulation, the “remap” option of *fix deform* should be set to “remap v”, since that is what *fix nvt/sllod* assumes to generate a velocity profile consistent with the applied shear strain rate.

An alternative method for calculating viscosities is provided via the *fix viscosity* command.

NEMD simulations can also be used to measure transport properties of a fluid through a pore or channel. Simulations of steady-state flow can be performed using the *fix flow/gauss* command.

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**(Daivis and Todd)** Daivis and Todd, Nonequilibrium Molecular Dynamics (book), Cambridge University Press, <https://doi.org/10.1017/9781139017848>, (2017).

## 8.2.8 Long-range dispersion settings

The PPPM method computes interactions by splitting the pair potential into two parts, one of which is computed in a normal pairwise fashion, the so-called real-space part, and one of which is computed using the Fourier transform, the so called reciprocal-space or kspace part. For both parts, the potential is not computed exactly but is approximated. Thus, there is an error in both parts of the computation, the real-space and the kspace error. The just mentioned facts are true both for the PPPM for Coulomb as well as dispersion interactions. The deciding difference - and also the reason why the parameters for *pppm/disp* have to be selected with more care - is the impact of the errors on the results: The kspace error of the PPPM for Coulomb and dispersion interaction and the real-space error of the PPPM for Coulomb interaction have the character of noise. In contrast, the real-space error of the PPPM for dispersion has a clear physical interpretation: the underprediction of cohesion. As a consequence, the real-space error has a much stronger effect than the kspace error on simulation results for *pppm/disp*. Parameters must thus be chosen in a way that this error is much smaller than the kspace error.

When using *pppm/disp* and not making any specifications on the PPPM parameters via the *kspace modify* command, parameters will be tuned such that the real-space error and the kspace error are equal. This will result in simulations that are either inaccurate or slow, both of which is not desirable. For selecting parameters for the *pppm/disp* that provide fast and accurate simulations, there are two approaches, which both have their up- and downsides.

The first approach is to set desired real-space and kspace accuracies via the *kspace\_modify force/disp/real* and *kspace\_modify force/disp/kspace* commands. Note that the accuracies have to be specified in force units and are thus

dependent on the chosen unit settings. For real units, 0.0001 and 0.002 seem to provide reasonable accurate and efficient computations for the real-space and kspace accuracies. 0.002 and 0.05 work well for most systems using lj units. PPPM parameters will be generated based on the desired accuracies. The upside of this approach is that it usually provides a good set of parameters and will work for both the *kspace\_modify diff ad* and *kspace\_modify diff ik* options. The downside of the method is that setting the PPPM parameters will take some time during the initialization of the simulation.

The second approach is to set the parameters for the *pppm/disp* explicitly using the *kspace\_modify mesh/disp*, *kspace\_modify order/disp*, and *kspace\_modify gewald/disp* commands. This approach requires a more experienced user who understands well the impact of the choice of parameters on the simulation accuracy and performance. This approach provides a fast initialization of the simulation. However, it is sensitive to errors: A combination of parameters that will perform well for one system might result in far-from-optimal conditions for other simulations. For example, parameters that provide accurate and fast computations for all-atomistic force fields can provide insufficient accuracy or united-atomistic force fields (which is related to that the latter typically have larger dispersion coefficients).

To avoid inaccurate or inefficient simulations, the *pppm/disp* stops simulations with an error message if no action is taken to control the PPPM parameters. If the automatic parameter generation is desired and real-space and kspace accuracies are desired to be equal, this error message can be suppressed using the *kspace\_modify disp/auto yes* command.

A reasonable approach that combines the upsides of both methods is to make the first run using the *kspace\_modify force/disp/real* and *kspace\_modify force/disp/kspace* commands, write down the PPPM parameters from the output, and specify these parameters using the second approach in subsequent runs (which have the same composition, force field, and approximately the same volume).

Concerning the performance of the *pppm/disp* there are two more things to consider. The first is that when using the *pppm/disp*, the cutoff parameter does no longer affect the accuracy of the simulation (subject to that *gewald/disp* is adjusted when changing the cutoff). The performance can thus be increased by examining different values for the cutoff parameter. A lower bound for the cutoff is only set by the truncation error of the repulsive term of pair potentials.

The second is that the mixing rule of the pair style has an impact on the computation time when using the *pppm/disp*. Fastest computations are achieved when using the geometric mixing rule. Using the arithmetic mixing rule substantially increases the computational cost. The computational overhead can be reduced using the *kspace\_modify mix/disp geom* and *kspace\_modify splittol* commands. The first command simply enforces geometric mixing of the dispersion coefficients in kspace computations. This introduces some error in the computations but will also significantly speed-up the simulations. The second keyword sets the accuracy with which the dispersion coefficients are approximated using a matrix factorization approach. This may result in better accuracy than using the first command, but will usually also not provide an equally good increase of efficiency.

Finally, *pppm/disp* can also be used when no mixing rules apply. This can be achieved using the *kspace\_modify mix/disp none* command. Note that the code does not check automatically whether any mixing rule is fulfilled. If mixing rules do not apply, the user will have to specify this command explicitly.

## 8.3 Analysis howto

### 8.3.1 Output from LAMMPS (thermo, dumps, computes, fixes, variables)

There are four basic kinds of LAMMPS output:

- *Thermodynamic output*, which is a list of quantities printed every few timesteps to the screen and logfile.
- *Dump files*, which contain snapshots of atoms and various per-atom values and are written at a specified frequency.
- Certain fixes can output user-specified quantities to files: *fix ave/time* for time averaging, *fix ave/chunk* for spatial or other averaging, and *fix print* for single-line output of *variables*. Fix print can also output to the screen.
- *Restart files*.

A simulation prints one set of thermodynamic output and (optionally) restart files. It can generate any number of dump files and fix output files, depending on what *dump* and *fix* commands you specify.

As discussed below, LAMMPS gives you a variety of ways to determine what quantities are computed and printed when the thermodynamics, dump, or fix commands listed above perform output. Throughout this discussion, note that users can also *add their own computes and fixes to LAMMPS* which can then generate values that can then be output with these commands.

The following sub-sections discuss different LAMMPS command related to output and the kind of data they operate on and produce:

- *Global/per-atom/local data*
- *Scalar/vector/array data*
- *Thermodynamic output*
- *Dump file output*
- *Fixes that write output files*
- *Computes that process output quantities*
- *Fixes that process output quantities*
- *Computes that generate values to output*
- *Fixes that generate values to output*
- *Variables that generate values to output*
- *Summary table of output options and data flow between commands*

### Global/per-atom/local data

Various output-related commands work with three different styles of data: global, per-atom, or local. A global datum is one or more system-wide values, e.g. the temperature of the system. A per-atom datum is one or more values per atom, e.g. the kinetic energy of each atom. Local datums are calculated by each processor based on the atoms it owns, but there may be zero or more per atom, e.g. a list of bond distances.

### Scalar/vector/array data

Global, per-atom, and local datums can each come in three kinds: a single scalar value, a vector of values, or a 2d array of values. The doc page for a “compute” or “fix” or “variable” that generates data will specify both the style and kind of data it produces, e.g. a per-atom vector.

When a quantity is accessed, as in many of the output commands discussed below, it can be referenced via the following bracket notation, where ID in this case is the ID of a compute. The leading “c\_” would be replaced by “f\_” for a fix, or “v\_” for a variable:

c_ID	entire scalar, vector, or array
c_ID[I]	one element of vector, one column of array
c_ID[I][J]	one element of array

In other words, using one bracket reduces the dimension of the data once (vector -> scalar, array -> vector). Using two brackets reduces the dimension twice (array -> scalar). Thus a command that uses scalar values as input can typically also process elements of a vector or array.

## Disambiguation

Some computes and fixes produce data in multiple styles, e.g. a global scalar and a per-atom vector. Usually the context in which the input script references the data determines which style is meant. Example: if a compute provides both a global scalar and a per-atom vector, the former will be accessed by using `c_ID` in an equal-style variable, while the latter will be accessed by using `c_ID` in an atom-style variable. Note that atom-style variable formulas can also access global scalars, but in this case it is not possible to do directly because of the ambiguity. Instead, an equal-style variable can be defined which accesses the global scalar, and that variable used in the atom-style variable formula in place of `c_ID`.

## Thermodynamic output

The frequency and format of thermodynamic output is set by the *thermo*, *thermo\_style*, and *thermo\_modify* commands. The *thermo\_style* command also specifies what values are calculated and written out. Pre-defined keywords can be specified (e.g. `press`, `etotal`, etc). Three additional kinds of keywords can also be specified (`c_ID`, `f_ID`, `v_name`), where a *compute* or *fix* or *variable* provides the value to be output. In each case, the compute, fix, or variable must generate global values for input to the *thermo\_style custom* command.

Note that thermodynamic output values can be “extensive” or “intensive”. The former scale with the number of atoms in the system (e.g. total energy), the latter do not (e.g. temperature). The setting for *thermo\_modify norm* determines whether extensive quantities are normalized or not. Computes and fixes produce either extensive or intensive values; see their individual doc pages for details. *Equal-style variables* produce only intensive values; you can include a division by “natoms” in the formula if desired, to make an extensive calculation produce an intensive result.

## Dump file output

Dump file output is specified by the *dump* and *dump\_modify* commands. There are several pre-defined formats (`dump atom`, `dump xtc`, etc).

There is also a *dump custom* format where the user specifies what values are output with each atom. Pre-defined atom attributes can be specified (`id`, `x`, `fx`, etc). Three additional kinds of keywords can also be specified (`c_ID`, `f_ID`, `v_name`), where a *compute* or *fix* or *variable* provides the values to be output. In each case, the compute, fix, or variable must generate per-atom values for input to the *dump custom* command.

There is also a *dump local* format where the user specifies what local values to output. A pre-defined index keyword can be specified to enumerate the local values. Two additional kinds of keywords can also be specified (`c_ID`, `f_ID`), where a *compute* or *fix* or *variable* provides the values to be output. In each case, the compute or fix must generate local values for input to the *dump local* command.

## Fixes that write output files

Several fixes take various quantities as input and can write output files: *fix ave/time*, *fix ave/chunk*, *fix ave/histo*, *fix ave/correlate*, and *fix print*.

The *fix ave/time* command enables direct output to a file and/or time-averaging of global scalars or vectors. The user specifies one or more quantities as input. These can be global *compute* values, global *fix* values, or *variables* of any style except the atom style which produces per-atom values. Since a variable can refer to keywords used by the *thermo\_style custom* command (like `temp` or `press`) and individual per-atom values, a wide variety of quantities can be time averaged and/or output in this way. If the inputs are one or more scalar values, then the fix generate a global scalar or vector of output. If the inputs are one or more vector values, then the fix generates a global vector or array of output. The time-averaged output of this fix can also be used as input to other output commands.

The *fix ave/chunk* command enables direct output to a file of chunk-averaged per-atom quantities like those output in dump files. Chunks can represent spatial bins or other collections of atoms, e.g. individual molecules. The per-atom

quantities can be atom density (mass or number) or atom attributes such as position, velocity, force. They can also be per-atom quantities calculated by a *compute*, by a *fix*, or by an atom-style *variable*. The chunk-averaged output of this *fix* can also be used as input to other output commands.

The *fix ave/histo* command enables direct output to a file of histogrammed quantities, which can be global or per-atom or local quantities. The histogram output of this *fix* can also be used as input to other output commands.

The *fix ave/correlate* command enables direct output to a file of time-correlated quantities, which can be global values. The correlation matrix output of this *fix* can also be used as input to other output commands.

The *fix print* command can generate a line of output written to the screen and log file or to a separate file, periodically during a running simulation. The line can contain one or more *variable* values for any style variable except the vector or atom styles). As explained above, variables themselves can contain references to global values generated by *thermodynamic keywords*, *computes*, *fixes*, or other *variables*, or to per-atom values for a specific atom. Thus the *fix print* command is a means to output a wide variety of quantities separate from normal thermodynamic or dump file output.

### Computes that process output quantities

The *compute reduce* and *compute reduce/region* commands take one or more per-atom or local vector quantities as inputs and “reduce” them (sum, min, max, ave) to scalar quantities. These are produced as output values which can be used as input to other output commands.

The *compute slice* command take one or more global vector or array quantities as inputs and extracts a subset of their values to create a new vector or array. These are produced as output values which can be used as input to other output commands.

The *compute property/atom* command takes a list of one or more pre-defined atom attributes (id, x, fx, etc) and stores the values in a per-atom vector or array. These are produced as output values which can be used as input to other output commands. The list of atom attributes is the same as for the *dump custom* command.

The *compute property/local* command takes a list of one or more pre-defined local attributes (bond info, angle info, etc) and stores the values in a local vector or array. These are produced as output values which can be used as input to other output commands.

### Fixes that process output quantities

The *fix vector* command can create global vectors as output from global scalars as input, accumulating them one element at a time.

The *fix ave/atom* command performs time-averaging of per-atom vectors. The per-atom quantities can be atom attributes such as position, velocity, force. They can also be per-atom quantities calculated by a *compute*, by a *fix*, or by an atom-style *variable*. The time-averaged per-atom output of this *fix* can be used as input to other output commands.

The *fix store/state* command can archive one or more per-atom attributes at a particular time, so that the old values can be used in a future calculation or output. The list of atom attributes is the same as for the *dump custom* command, including per-atom quantities calculated by a *compute*, by a *fix*, or by an atom-style *variable*. The output of this *fix* can be used as input to other output commands.

## Computes that generate values to output

Every *compute* in LAMMPS produces either global or per-atom or local values. The values can be scalars or vectors or arrays of data. These values can be output using the other commands described in this section. The page for each compute command describes what it produces. Computes that produce per-atom or local values have the word “atom” or “local” in their style name. Computes without the word “atom” or “local” produce global values.

## Fixes that generate values to output

Some *fixes* in LAMMPS produces either global or per-atom or local values which can be accessed by other commands. The values can be scalars or vectors or arrays of data. These values can be output using the other commands described in this section. The page for each fix command tells whether it produces any output quantities and describes them.

## Variables that generate values to output

*Variables* defined in an input script can store one or more strings. But equal-style, vector-style, and atom-style or atomfile-style variables generate a global scalar value, global vector or values, or a per-atom vector, respectively, when accessed. The formulas used to define these variables can contain references to the thermodynamic keywords and to global and per-atom data generated by computes, fixes, and other variables. The values generated by variables can be used as input to and thus output by the other commands described in this section.

## Summary table of output options and data flow between commands

This table summarizes the various commands that can be used for generating output from LAMMPS. Each command produces output data of some kind and/or writes data to a file. Most of the commands can take data from other commands as input. Thus you can link many of these commands together in pipeline form, where data produced by one command is used as input to another command and eventually written to the screen or to a file. Note that to hook two commands together the output and input data types must match, e.g. global/per-atom/local data and scalar/vector/array data.

Also note that, as described above, when a command takes a scalar as input, that could be an element of a vector or array. Likewise a vector input could be a column of an array.

Command	Input	Output
<i>thermo_style custom</i>	global scalars	screen, log file
<i>dump custom</i>	per-atom vectors	dump file
<i>dump local</i>	local vectors	dump file
<i>fix print</i>	global scalar from variable	screen, file
<i>print</i>	global scalar from variable	screen
<i>computes</i>	N/A	global/per-atom/local scalar/vector/array
<i>fixes</i>	N/A	global/per-atom/local scalar/vector/array
<i>variables</i>	global scalars and vectors, per-atom vectors	global scalar and vector, per-atom vector
<i>compute reduce</i>	per-atom/local vectors	global scalar/vector
<i>compute slice</i>	global vectors/arrays	global vector/array
<i>compute property/atom</i>	per-atom vectors	per-atom vector/array
<i>compute property/local</i>	local vectors	local vector/array
<i>fix vector</i>	global scalars	global vector
<i>fix ave/atom</i>	per-atom vectors	per-atom vector/array
<i>fix ave/time</i>	global scalars/vectors	global scalar/vector/array, file
<i>fix ave/chunk</i>	per-atom vectors	global array, file
<i>fix ave/histo</i>	global/per-atom/local scalars and vectors	global array, file
<i>fix ave/correlate</i>	global scalars	global array, file
<i>fix store/state</i>	per-atom vectors	per-atom vector/array

### 8.3.2 Use chunks to calculate system properties

In LAMMPS, “chunks” are collections of atoms, as defined by the *compute chunk/atom* command, which assigns each atom to a chunk ID (or to no chunk at all). The number of chunks and the assignment of chunk IDs to atoms can be static or change over time. Examples of “chunks” are molecules or spatial bins or atoms with similar values (e.g. coordination number or potential energy).

The per-atom chunk IDs can be used as input to two other kinds of commands, to calculate various properties of a system:

- *fix ave/chunk*
- any of the *compute \*/chunk* commands

Here a brief overview for each of the 4 kinds of chunk-related commands is provided. Then some examples are given of how to compute different properties with chunk commands.

### Compute chunk/atom command:

This compute can assign atoms to chunks of various styles. Only atoms in the specified group and optional specified region are assigned to a chunk. Here are some possible chunk definitions:

atoms in same molecule	chunk ID = molecule ID
atoms of same atom type	chunk ID = atom type
all atoms with same atom property (charge, radius, etc)	chunk ID = output of compute property/atom
atoms in same cluster	chunk ID = output of <i>compute cluster/atom</i> command
atoms in same spatial bin	chunk ID = bin ID
atoms in same rigid body	chunk ID = molecule ID used to define rigid bodies
atoms with similar potential energy	chunk ID = output of <i>compute pe/atom</i>
atoms with same local defect structure	chunk ID = output of <i>compute centro/atom</i> or <i>compute coord/atom</i> command

Note that chunk IDs are integer values, so for atom properties or computes that produce a floating point value, they will be truncated to an integer. You could also use the compute in a variable that scales the floating point value to spread it across multiple integers.

Spatial bins can be of various kinds, e.g. 1d bins = slabs, 2d bins = pencils, 3d bins = boxes, spherical bins, cylindrical bins.

This compute also calculates the number of chunks *Nchunk*, which is used by other commands to tally per-chunk data. *Nchunk* can be a static value or change over time (e.g. the number of clusters). The chunk ID for an individual atom can also be static (e.g. a molecule ID), or dynamic (e.g. what spatial bin an atom is in as it moves).

Note that this compute allows the per-atom output of other *computes*, *fixes*, and *variables* to be used to define chunk IDs for each atom. This means you can write your own compute or fix to output a per-atom quantity to use as chunk ID. See the *Modify* doc pages for info on how to do this. You can also define a *per-atom variable* in the input script that uses a formula to generate a chunk ID for each atom.

### Fix ave/chunk command:

This fix takes the ID of a *compute chunk/atom* command as input. For each chunk, it then sums one or more specified per-atom values over the atoms in each chunk. The per-atom values can be any atom property, such as velocity, force, charge, potential energy, kinetic energy, stress, etc. Additional keywords are defined for per-chunk properties like density and temperature. More generally any per-atom value generated by other *computes*, *fixes*, and *per-atom variables*, can be summed over atoms in each chunk.

Similar to other averaging fixes, this fix allows the summed per-chunk values to be time-averaged in various ways, and output to a file. The fix produces a global array as output with one row of values per chunk.

### Compute \*/chunk commands:

The following computes operate on chunks of atoms to produce per-chunk values. Any compute whose style name ends in “/chunk” is in this category:

- *compute com/chunk*
- *compute gyration/chunk*
- *compute inertia/chunk*
- *compute msd/chunk*



- *compute property/chunk*
- *compute temp/chunk*
- *compute torque/chunk*
- *compute vcm/chunk*

They each take the ID of a *compute chunk/atom* command as input. As their names indicate, they calculate the center-of-mass, radius of gyration, moments of inertia, mean-squared displacement, temperature, torque, and velocity of center-of-mass for each chunk of atoms. The *compute property/chunk* command can tally the count of atoms in each chunk and extract other per-chunk properties.

The reason these various calculations are not part of the *fix ave/chunk command*, is that each requires a more complicated operation than simply summing and averaging over per-atom values in each chunk. For example, many of them require calculation of a center of mass, which requires summing mass\*position over the atoms and then dividing by summed mass.

All of these computes produce a global vector or global array as output, with one or more values per chunk. The output can be used in various ways:

- As input to the *fix ave/time* command, which can write the values to a file and optionally time average them.
- As input to the *fix ave/histo* command to histogram values across chunks. E.g. a histogram of cluster sizes or molecule diffusion rates.
- As input to special functions of *equal-style variables*, like `sum()` and `max()` and `ave()`. E.g. to find the largest cluster or fastest diffusing molecule or average radius-of-gyration of a set of molecules (chunks).

#### Other chunk commands:

- *compute chunk/spread/atom*
- *compute reduce/chunk*

The *compute chunk/spread/atom* command spreads per-chunk values to each atom in the chunk, producing per-atom values as its output. This can be useful for outputting per-chunk values to a per-atom *dump file*. Or for using an atom's associated chunk value in an *atom-style variable*. Or as input to the *fix ave/chunk* command to spatially average per-chunk values calculated by a per-chunk compute.

The *compute reduce/chunk* command reduces a peratom value across the atoms in each chunk to produce a value per chunk. When used with the *compute chunk/spread/atom* command it can create peratom values that induce a new set of chunks with a second *compute chunk/atom* command.

#### Example calculations with chunks

Here are examples using chunk commands to calculate various properties:

- (1) Average velocity in each of 1000 2d spatial bins:

```
compute cc1 all chunk/atom bin/2d x 0.0 0.1 y lower 0.01 units reduced
fix 1 all ave/chunk 100 10 1000 cc1 vx vy file tmp.out
```

- (2) Temperature in each spatial bin, after subtracting a flow velocity:

```
compute cc1 all chunk/atom bin/2d x 0.0 0.1 y lower 0.1 units reduced
compute vbias all temp/profile 1 0 0 y 10
fix 1 all ave/chunk 100 10 1000 cc1 temp bias vbias file tmp.out
```

(3) Center of mass of each molecule:

```
compute cc1 all chunk/atom molecule
compute myChunk all com/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk[*] file tmp.out mode vector
```

(4) Total force on each molecule and ave/max across all molecules:

```
compute cc1 all chunk/atom molecule
fix 1 all ave/chunk 1000 1 1000 cc1 fx fy fz file tmp.out
variable xave equal ave(f_1[2])
variable xmax equal max(f_1[2])
thermo 1000
thermo_style custom step temp v_xave v_xmax
```

(5) Histogram of cluster sizes:

```
compute cluster all cluster/atom 1.0
compute cc1 all chunk/atom c_cluster compress yes
compute size all property/chunk cc1 count
fix 1 all ave/histo 100 1 100 0 20 20 c_size mode vector ave running beyond ignore file_
↪tmp.histo
```

(6) An example for using a per-chunk value to apply per-atom forces to compress individual polymer chains (molecules) in a mixture, is explained on the [compute chunk/spread/atom](#) command doc page.

(7) An example for using one set of per-chunk values for molecule chunks, to create a second set of micelle-scale chunks (clustered molecules, due to hydrophobicity), is explained on the [compute reduce/chunk](#) command doc page.

(8) An example for using one set of per-chunk values (dipole moment vectors) for molecule chunks, spreading the values to each atom in each chunk, then defining a second set of chunks as spatial bins, and using the [fix ave/chunk](#) command to calculate an average dipole moment vector for each bin. This example is explained on the [compute chunk/spread/atom](#) command doc page.

### 8.3.3 Calculate temperature

Temperature is computed as kinetic energy divided by some number of degrees of freedom (and the Boltzmann constant). Since kinetic energy is a function of particle velocity, there is often a need to distinguish between a particle's advection velocity (due to some aggregate motion of particles) and its thermal velocity. The sum of the two is the particle's total velocity, but the latter is often what is wanted to compute a temperature.

LAMMPS has several options for computing temperatures, any of which can be used in *thermostatting* and *barostatting*. These [compute commands](#) calculate temperature:

- [compute temp](#)
- [compute temp/sphere](#)
- [compute temp/asphere](#)
- [compute temp/com](#)
- [compute temp/deform](#)
- [compute temp/partial](#)
- [compute temp/profile](#)
- [compute temp/ramp](#)

- *compute temp/region*

All but the first 3 calculate velocity biases directly (e.g. advection velocities) that are removed when computing the thermal temperature. *Compute temp/sphere* and *compute temp/asphere* compute kinetic energy for finite-size particles that includes rotational degrees of freedom. They both allow for velocity biases indirectly, via an optional extra argument which is another temperature compute that subtracts a velocity bias. This allows the translational velocity of spherical or aspherical particles to be adjusted in prescribed ways.

### 8.3.4 Calculate elastic constants

Elastic constants characterize the stiffness of a material. The formal definition is provided by the linear relation that holds between the stress and strain tensors in the limit of infinitesimal deformation. In tensor notation, this is expressed as  $s_{ij} = C_{ijkl} * e_{kl}$ , where the repeated indices imply summation.  $s_{ij}$  are the elements of the symmetric stress tensor.  $e_{kl}$  are the elements of the symmetric strain tensor.  $C_{ijkl}$  are the elements of the fourth rank tensor of elastic constants. In three dimensions, this tensor has  $3^4=81$  elements. Using Voigt notation, the tensor can be written as a 6x6 matrix, where  $C_{ij}$  is now the derivative of  $s_i$  w.r.t.  $e_j$ . Because  $s_i$  is itself a derivative w.r.t.  $e_i$ , it follows that  $C_{ij}$  is also symmetric, with at most  $7*6/2 = 21$  distinct elements.

At zero temperature, it is easy to estimate these derivatives by deforming the simulation box in one of the six directions using the *change\_box* command and measuring the change in the stress tensor. A general-purpose script that does this is given in the examples/ELASTIC directory described on the *Examples* doc page.

Calculating elastic constants at finite temperature is more challenging, because it is necessary to run a simulation that performs time averages of differential properties. There are at least 3 ways to do this in LAMMPS. The most reliable way to do this is by exploiting the relationship between elastic constants, stress fluctuations, and the Born matrix, the second derivatives of energy w.r.t. strain (*Ray*). The Born matrix calculation has been enabled by the *compute born/matrix* command, which works for any bonded or non-bonded potential in LAMMPS. The most expensive part of the calculation is the sampling of the stress fluctuations. Several examples of this method are provided in the examples/ELASTIC\_T/BORN\_MATRIX directory described on the *Examples* doc page.

A second way is to measure the change in average stress tensor in an NVT simulations when the cell volume undergoes a finite deformation. In order to balance the systematic and statistical errors in this method, the magnitude of the deformation must be chosen judiciously, and care must be taken to fully equilibrate the deformed cell before sampling the stress tensor. An example of this method is provided in the examples/ELASTIC\_T/DEFORMATION directory described on the *Examples* doc page.

Another approach is to sample the triclinic cell fluctuations that occur in an NPT simulation. This method can also be slow to converge and requires careful post-processing (*Shinoda*). We do not provide an example of this method.

A nice review of the advantages and disadvantages of all of these methods is provided in the paper by Clavier et al. (*Clavier*).

---

**(Ray)** J. R. Ray and A. Rahman, J Chem Phys, 80, 4423 (1984).

**(Shinoda)** Shinoda, Shiga, and Mikami, Phys Rev B, 69, 134103 (2004).

**(Clavier)** G. Clavier, N. Desbiens, E. Bourasseau, V. Lachet, N. Brusselle-Dupend and B. Rousseau, Mol Sim, 43, 1413 (2017).

### 8.3.5 Calculate thermal conductivity

The thermal conductivity  $\kappa$  of a material can be measured in at least 4 ways using various options in LAMMPS. See the examples/KAPPA directory for scripts that implement the 4 methods discussed here for a simple Lennard-Jones fluid model. Also, see the [Howto viscosity](#) page for an analogous discussion for viscosity.

The thermal conductivity tensor  $\kappa$  is a measure of the propensity of a material to transmit heat energy in a diffusive manner as given by Fourier's law

$$J = -\kappa \text{grad}(T)$$

where  $J$  is the heat flux in units of energy per area per time and  $\text{grad}(T)$  is the spatial gradient of temperature. The thermal conductivity thus has units of energy per distance per time per degree K and is often approximated as an isotropic quantity, i.e. as a scalar.

The first method is to setup two thermostatted regions at opposite ends of a simulation box, or one in the middle and one at the end of a periodic box. By holding the two regions at different temperatures with a *thermostatting fix*, the energy added to the hot region should equal the energy subtracted from the cold region and be proportional to the heat flux moving between the regions. See the papers by *Ikeshoji and Hafskjold* and *Wirnsberger et al* for details of this idea. Note that thermostatting fixes such as *fix nvt*, *fix langevin*, and *fix temp/rescale* store the cumulative energy they add/subtract.

Alternatively, as a second method, the *fix heat* or *fix ehex* commands can be used in place of thermostats on each of two regions to add/subtract specified amounts of energy to both regions. In both cases, the resulting temperatures of the two regions can be monitored with the “compute temp/region” command and the temperature profile of the intermediate region can be monitored with the *fix ave/chunk* and *compute ke/atom* commands.

The third method is to perform a reverse non-equilibrium MD simulation using the *fix thermal/conductivity* command which implements the rNEMD algorithm of Muller-Plathe. Kinetic energy is swapped between atoms in two different layers of the simulation box. This induces a temperature gradient between the two layers which can be monitored with the *fix ave/chunk* and *compute ke/atom* commands. The *fix* tallies the cumulative energy transfer that it performs. See the *fix thermal/conductivity* command for details.

The fourth method is based on the Green-Kubo (GK) formula which relates the ensemble average of the auto-correlation of the heat flux to  $\kappa$ . The heat flux can be calculated from the fluctuations of per-atom potential and kinetic energies and per-atom stress tensor in a steady-state equilibrated simulation. This is in contrast to the two preceding non-equilibrium methods, where energy flows continuously between hot and cold regions of the simulation box.

The *compute heat/flux* command can calculate the needed heat flux and describes how to implement the Green\_Kubo formalism using additional LAMMPS commands, such as the *fix ave/correlate* command to calculate the needed auto-correlation. See the page for the *compute heat/flux* command for an example input script that calculates the thermal conductivity of solid Ar via the GK formalism.

---

(Ikeshoji) Ikeshoji and Hafskjold, *Molecular Physics*, 81, 251-261 (1994).

(Wirnsberger) Wirnsberger, Frenkel, and Dellago, *J Chem Phys*, 143, 124104 (2015).

### 8.3.6 Calculate viscosity

The shear viscosity  $\eta$  of a fluid can be measured in at least 6 ways using various options in LAMMPS. See the examples/VISCOSITY directory for scripts that implement the 5 methods discussed here for a simple Lennard-Jones fluid model and 1 method for SPC/E water model. Also, see the [page on calculating thermal conductivity](#) for an analogous discussion for thermal conductivity.

$\eta$  is a measure of the propensity of a fluid to transmit momentum in a direction perpendicular to the direction of velocity or momentum flow. Alternatively it is the resistance the fluid has to being sheared. It is given by

$$J = -\eta \text{grad}(V_{\text{stream}})$$

where  $J$  is the momentum flux in units of momentum per area per time. and  $\text{grad}(V_{\text{stream}})$  is the spatial gradient of the velocity of the fluid moving in another direction, normal to the area through which the momentum flows. Viscosity thus has units of pressure-time.

The first method is to perform a non-equilibrium MD (NEMD) simulation by shearing the simulation box via the *fix deform* command, and using the *fix nvt/sllod* command to thermostat the fluid via the SLLOD equations of motion. Alternatively, as a second method, one or more moving walls can be used to shear the fluid in between them, again with some kind of thermostat that modifies only the thermal (non-shearing) components of velocity to prevent the fluid from heating up.

---

**Note:** A recent (2017) book by (*Daivis and Todd*) discusses use of the SLLOD method and non-equilibrium MD (NEMD) thermostating generally, for both simple and complex fluids, e.g. molecular systems. The latter can be tricky to do correctly.

---

In both cases, the velocity profile setup in the fluid by this procedure can be monitored by the *fix ave/chunk* command, which determines  $\text{grad}(V_{\text{stream}})$  in the equation above. E.g. the derivative in the y-direction of the  $V_x$  component of fluid motion or  $\text{grad}(V_{\text{stream}}) = dV_x/dy$ . The  $P_{xy}$  off-diagonal component of the pressure or stress tensor, as calculated by the *compute pressure* command, can also be monitored, which is the  $J$  term in the equation above. See the *Howto nemd* page for details on NEMD simulations.

The third method is to perform a reverse non-equilibrium MD simulation using the *fix viscosity* command which implements the rNEMD algorithm of Muller-Plathe. Momentum in one dimension is swapped between atoms in two different layers of the simulation box in a different dimension. This induces a velocity gradient which can be monitored with the *fix ave/chunk* command. The *fix* tallies the cumulative momentum transfer that it performs. See the *fix viscosity* command for details.

The fourth method is based on the Green-Kubo (GK) formula which relates the ensemble average of the auto-correlation of the stress/pressure tensor to  $\eta$ . This can be done in a fully equilibrated simulation which is in contrast to the two preceding non-equilibrium methods, where momentum flows continuously through the simulation box.

Here is an example input script that calculates the viscosity of liquid Ar via the GK formalism:

```
# Sample LAMMPS input script for viscosity of liquid Ar

units      real
variable   T equal 200.0      # run temperature
variable   Tinit equal 250.0  # equilibration temperature
variable   V equal vol
variable   dt equal 4.0
variable   p equal 400        # correlation length
variable   s equal 5          # sample interval
variable   d equal $p*$s      # dump interval

# convert from LAMMPS real units to SI

variable   kB equal 1.3806504e-23 # [J/K] Boltzmann
variable   atm2Pa equal 101325.0
variable   A2m equal 1.0e-10
variable   fs2s equal 1.0e-15
variable   convert equal ${atm2Pa}*${atm2Pa}*${fs2s}*${A2m}*${A2m}*${A2m}

# setup problem
```

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```

dimension      3
boundary       p p p
lattice        fcc 5.376 orient x 1 0 0 orient y 0 1 0 orient z 0 0 1
region         box block 0 4 0 4 0 4
create_box     1 box
create_atoms   1 box
mass           1 39.948
pair_style     lj/cut 13.0
pair_coeff     * * 0.2381 3.405
timestep       ${dt}
thermo         $d

# equilibration and thermalization

velocity       all create ${Tinit} 102486 mom yes rot yes dist gaussian
fix            NVT all nvt temp ${Tinit} ${Tinit} 10 drag 0.2
run            8000

# viscosity calculation, switch to NVE if desired

velocity       all create $T 102486 mom yes rot yes dist gaussian
fix            NVT all nvt temp $T $T 10 drag 0.2
#unfix        NVT
#fix           NVE all nve

reset_timestep 0
variable       pxy equal pxy
variable       pxz equal pxz
variable       pyz equal pyz
fix            SS all ave/correlate $s $p $d &
              v_pxy v_pxz v_pyz type auto file S0St.dat ave running
variable       scale equal ${convert}/(${kB}*$T)*$V*$s*${dt}
variable       v11 equal trap(f_SS[3])*${scale}
variable       v22 equal trap(f_SS[4])*${scale}
variable       v33 equal trap(f_SS[5])*${scale}
thermo_style   custom step temp press v_pxy v_pxz v_pyz v_v11 v_v22 v_v33
run            100000
variable       v equal (v_v11+v_v22+v_v33)/3.0
variable       ndens equal count(all)/vol
print          "average viscosity: $v [Pa.s] @ $T K, ${ndens} atoms/A^3"

```

The fifth method is related to the above Green-Kubo method, but uses the Einstein formulation, analogous to the Einstein mean-square-displacement formulation for self-diffusivity. The time-integrated momentum fluxes play the role of Cartesian coordinates, whose mean-square displacement increases linearly with time at sufficiently long times.

The sixth is the periodic perturbation method, which is also a non-equilibrium MD method. However, instead of measuring the momentum flux in response to an applied velocity gradient, it measures the velocity profile in response to applied stress. A cosine-shaped periodic acceleration is added to the system via the *fix accelerate/cos* command, and the *compute viscosity/cos* command is used to monitor the generated velocity profile and remove the velocity bias before thermostating.

---

**Note:** An article by (*Hess*) discussed the accuracy and efficiency of these methods.

---

**(Daivis and Todd)** Daivis and Todd, Nonequilibrium Molecular Dynamics (book), Cambridge University Press, <https://doi.org/10.1017/9781139017848>, (2017).

**(Hess)** Hess, B. The Journal of Chemical Physics 2002, 116 (1), 209-217.

### 8.3.7 Calculate diffusion coefficients

The diffusion coefficient  $D$  of a material can be measured in at least 2 ways using various options in LAMMPS. See the examples/DIFFUSE directory for scripts that implement the 2 methods discussed here for a simple Lennard-Jones fluid model.

The first method is to measure the mean-squared displacement (MSD) of the system, via the `compute msd` command. The slope of the MSD versus time is proportional to the diffusion coefficient. The instantaneous MSD values can be accumulated in a vector via the `fix vector` command, and a line fit to the vector to compute its slope via the `variable slope` function, and thus extract  $D$ .

The second method is to measure the velocity auto-correlation function (VACF) of the system, via the `compute vacf` command. The time-integral of the VACF is proportional to the diffusion coefficient. The instantaneous VACF values can be accumulated in a vector via the `fix vector` command, and time integrated via the `variable trap` function, and thus extract  $D$ .

### 8.3.8 Output structured data from LAMMPS

LAMMPS can output structured data with the `print` and `fix print` command. This gives you flexibility since you can build custom data formats that contain system properties, thermo data, and variables values. This output can be directed to the screen and/or to a file for post processing.

#### Writing the current system state, thermo data, variable values

Use the `print` command to output the current system state, which can include system properties, thermo data and variable values.

#### YAML

```
print ""---
timestep: $(step)
pe: $(pe)
ke: $(ke)
..."" file current_state.yaml screen no
```

Listing 1: current\_state.yaml

```
---
timestep: 250
pe: -4.7774327356321810711
ke: 2.4962152903997174569
```

## JSON

```
print """{
  "timestep": $(step),
  "pe": $(pe),
  "ke": $(ke)
}""" file current_state.json screen no
```

Listing 2: current\_state.json

```
{
  "timestep": 250,
  "pe": -4.7774327356321810711,
  "ke": 2.4962152903997174569
}
```

## YAML format thermo\_style output

## Extracting data from log file

New in version 24Mar2022.

LAMMPS supports the thermo style “yaml” and for “custom” style thermodynamic output the format can be changed to YAML with *thermo\_modify line yaml*. This will produce a block of output in a compact YAML format - one “document” per run - of the following style:

```
---
keywords: ['Step', 'Temp', 'E_pair', 'E_mol', 'TotEng', 'Press', ]
data:
- [100, 0.757453103239935, -5.7585054860159, 0, -4.62236133677021, 0.207261053624721, ]
- [110, 0.759322359337036, -5.7614668389562, 0, -4.62251889318624, 0.194314975399602, ]
- [120, 0.759372342462676, -5.76149365656489, 0, -4.62247073844943, 0.191600048851267, ]
↵]
- [130, 0.756833027516501, -5.75777334823494, 0, -4.62255928350835, 0.208792327853067, ]
↵]
...

```

This data can be extracted and parsed from a log file using python with:

```
import re, yaml
try:
    from yaml import CSafeLoader as Loader
except ImportError:
    from yaml import SafeLoader as Loader

docs = ""
with open("log.lammps") as f:
    for line in f:
        m = re.search(r"^(keywords:.*$|data:$|---$|\.\.\.$| - \[.*\]$)", line)
        if m: docs += m.group(0) + '\n'

thermo = list(yaml.load_all(docs, Loader=Loader))
```

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```
print("Number of runs: ", len(thermo))
print(thermo[1]['keywords'][4], ' = ', thermo[1]['data'][2][4])
```

After loading the YAML data, *thermo* is a list containing a dictionary for each “run” where the tag “keywords” maps to the list of thermo header strings and the tag “data” has a list of lists where the outer list represents the lines of output and the inner list the values of the columns matching the header keywords for that step. The second print() command for example will print the header string for the fifth keyword of the second run and the corresponding value for the third output line of that run:

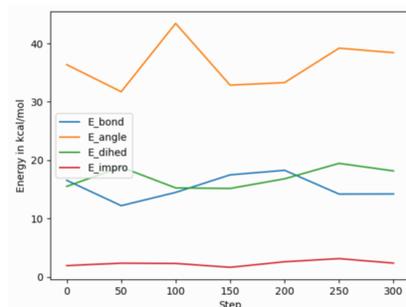
```
Number of runs: 2
TotEng = -4.62140097780047
```

New in version 4May2022.

YAML format output has been added to multiple commands in LAMMPS, for example *dump yaml* or *fix ave/time*. Depending on the kind of data being written, organization of the data or the specific syntax used may change, but the principles are very similar and all files should be readable with a suitable YAML parser.

### Processing scalar data with Python

After reading and parsing the YAML format data, it can be easily imported for further processing and visualization with the *pandas* and *matplotlib* Python modules. Because of the organization of the data in the YAML format thermo output, it needs to be told to process only the ‘data’ part of the imported data to create a pandas data frame, and one needs to set the column names from the ‘keywords’ entry. The following Python script code example demonstrates this, and creates the image shown on the right of a simple plot of various bonded energy contributions versus the timestep from a run of the ‘peptide’ example input after changing the *thermo style* to ‘yaml’. The properties to be used for x and y values can be conveniently selected through the keywords. Please note that those keywords can be changed to custom strings with the *thermo\_modify colname* command.



```
import re, yaml
import pandas as pd
import matplotlib.pyplot as plt

try:
    from yaml import CSafeLoader as Loader
except ImportError:
    from yaml import SafeLoader as Loader

docs = ""
with open("log.lammps") as f:
    for line in f:
        m = re.search(r"^(keywords:.*$|data:$|---$|\.\.\.\$| - \[.*\])", line)
        if m: docs += m.group(0) + '\n'

thermo = list(yaml.load_all(docs, Loader=Loader))

df = pd.DataFrame(data=thermo[0]['data'], columns=thermo[0]['keywords'])
```

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```
fig = df.plot(x='Step', y=['E_bond', 'E_angle', 'E_dihed', 'E_impro'], ylabel='Energy in_
↳kcal/mol')
plt.savefig('thermo_bondeng.png')
```

## Processing vector data with Python

Global *vector* data as produced by *fix ave/time* uses a slightly different organization of the data. You still have the dictionary keys ‘keywords’ and ‘data’ for the column headers and the data. But the data is a dictionary indexed by the time step and for each step there are multiple rows of values each with a list of the averaged properties. This requires a slightly different processing, since the entire data cannot be directly imported into a single pandas DataFrame class instance. The following Python script example demonstrates how to read such data. The result will combine the data for the different steps into one large “multi-index” table. The pandas IndexSlice class can then be used to select data from this combined data frame.

```
import yaml
import pandas as pd

try:
    from yaml import CSafeLoader as Loader
except ImportError:
    from yaml import SafeLoader as Loader

with open("ave.yaml") as f:
    ave = yaml.load(f, Loader=Loader)

keys = ave['keywords']
df = {}
for k in ave['data'].keys():
    df[k] = pd.DataFrame(data=ave['data'][k], columns=keys)

# create multi-index data frame
df = pd.concat(df)

# output only the first 3 value for steps 200 to 300 of the column Pressure
idx = pd.IndexSlice
print(df['Pressure'].loc[idx[200:300, 0:2]])
```

## Processing scalar data with Perl

The ease of processing YAML data is not limited to Python. Here is an example for extracting and processing a LAMMPS log file with Perl instead.

```
use YAML::XS;

open(LOG, "log.lammps") or die("could not open log.lammps: $!");
my $file = "";
while(my $line = <LOG>) {
    if ($line =~ /^(keywords:.*$|data:$|---$|\.\.\.$| - \[.*\]$)/) {
        $file .= $line;
    }
}
```

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```

    }
  }
close(LOG);

# convert YAML to perl as nested hash and array references
my $thermo = Load $file;

# convert references to real arrays
my @keywords = @{$thermo->{'keywords'}};
my @data = @{$thermo->{'data'}};

# print first two columns
print("$keywords[0] $keywords[1]\n");
foreach (@data) {
    print("${$_}[0] ${$_}[1]\n");
}

```

### Writing continuous data during a simulation

The *fix print* command allows you to output an arbitrary string at defined times during a simulation run.

### YAML

```

fix extra all print 50 ""
- timestep: $(step)
  pe: $(pe)
  ke: $(ke)"" file output.yaml screen no

```

Listing 3: output.yaml

```

# Fix print output for fix extra
- timestep: 0
  pe: -6.77336805325924729
  ke: 4.49887500000000026219

- timestep: 50
  pe: -4.8082494418323200591
  ke: 2.5257981827119797558

- timestep: 100
  pe: -4.7875608875581505686
  ke: 2.5062598821985102582

- timestep: 150
  pe: -4.7471033686005483787
  ke: 2.466095925545450207

- timestep: 200
  pe: -4.7509052858544134068

```

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```

ke: 2.4701136792591693592
- timestep: 250
  pe: -4.7774327356321810711
  ke: 2.4962152903997174569

```

Post-processing of YAML files can be easily be done with Python and other scripting languages. In case of Python the *yaml* package allows you to load the data files and obtain a list of dictionaries.

```

import yaml

with open("output.yaml") as f:
    data = yaml.load(f, Loader=yaml.FullLoader)

print(data)

```

```

[{'timestep': 0, 'pe': -6.773368053259247, 'ke': 4.4988750000000003}, {'timestep': 50, 'pe'
↪ ': -4.80824944183232, 'ke': 2.5257981827119798}, {'timestep': 100, 'pe': -4.
↪ 787560887558151, 'ke': 2.5062598821985103}, {'timestep': 150, 'pe': -4.747103368600548,
↪ 'ke': 2.46609592554545}, {'timestep': 200, 'pe': -4.750905285854413, 'ke': 2.
↪ 4701136792591694}, {'timestep': 250, 'pe': -4.777432735632181, 'ke': 2.
↪ 4962152903997175}]

```

## Line Delimited JSON (LD-JSON)

The JSON format itself is very strict when it comes to delimiters. For continuous output/streaming data it is beneficial use the *line delimited JSON* format. Each line represents one JSON object.

```

fix extra all print 50 """"{"timestep": $(step), "pe": $(pe), "ke": $(ke)}"""" title ""_
↪file output.json screen no

```

Listing 4: output.json

```

{"timestep": 0, "pe": -6.77336805325924729, "ke": 4.49887500000000026219}
{"timestep": 50, "pe": -4.8082494418323200591, "ke": 2.5257981827119797558}
{"timestep": 100, "pe": -4.7875608875581505686, "ke": 2.5062598821985102582}
{"timestep": 150, "pe": -4.7471033686005483787, "ke": 2.466095925545450207}
{"timestep": 200, "pe": -4.7509052858544134068, "ke": 2.4701136792591693592}
{"timestep": 250, "pe": -4.7774327356321810711, "ke": 2.4962152903997174569}

```

One simple way to load this data into a Python script is to use the *pandas* package. It can directly load these files into a data frame:

```

import pandas as pd

data = pd.read_json('output.json', lines=True)
print(data)

```

	timestep	pe	ke
0	0	-6.773368	4.498875

(continues on next page)

(continued from previous page)

1	50	-4.808249	2.525798
2	100	-4.787561	2.506260
3	150	-4.747103	2.466096
4	200	-4.750905	2.470114
5	250	-4.777433	2.496215

## 8.4 Force fields howto

### 8.4.1 CHARMM, AMBER, COMPASS, and DREIDING force fields

A force field has 2 parts: the formulas that define it and the coefficients used for a particular system. Here we only discuss formulas implemented in LAMMPS that correspond to formulas commonly used in the CHARMM, AMBER, COMPASS, and DREIDING force fields. Setting coefficients is done either from special sections in an input data file via the *read\_data* command or in the input script with commands like *pair\_coeff* or *bond\_coeff* and so on. See the *Tools* doc page for additional tools that can use CHARMM, AMBER, or Materials Studio generated files to assign force field coefficients and convert their output into LAMMPS input.

See (*MacKerell*) for a description of the CHARMM force field. See (*Cornell*) for a description of the AMBER force field. See (*Sun*) for a description of the COMPASS force field.

The interaction styles listed below compute force field formulas that are consistent with common options in CHARMM or AMBER. See each command's documentation for the formula it computes.

- *bond\_style* harmonic
- *angle\_style* charmm
- *dihedral\_style* charmmfsh
- *dihedral\_style* charmm
- *pair\_style* lj/charmmfsw/coul/charmmfsh
- *pair\_style* lj/charmmfsw/coul/long
- *pair\_style* lj/charmm/coul/charmm
- *pair\_style* lj/charmm/coul/charmm/implicit
- *pair\_style* lj/charmm/coul/long
- *special\_bonds* charmm
- *special\_bonds* amber

---

**Note:** For CHARMM, newer *charmmfsw* or *charmmfsh* styles were released in March 2017. We recommend they be used instead of the older *charmm* styles. See discussion of the differences on the *pair charmm* and *dihedral charmm* doc pages.

---

COMPASS is a general force field for atomistic simulation of common organic molecules, inorganic small molecules, and polymers which was developed using ab initio and empirical parameterization techniques. See the *Tools* page for the *msi2lmp* tool for creating LAMMPS template input and data files from BIOVIA's Materials Studio files. Please note that the *msi2lmp* tool is very old and largely unmaintained, so it does not support all features of Materials Studio provided force field files, especially additions during the last decade. You should watch the output carefully and compare results, where possible. See (*Sun*) for a description of the COMPASS force field.

These interaction styles listed below compute force field formulas that are consistent with the COMPASS force field. See each command's documentation for the formula it computes.

- *bond\_style* class2
- *angle\_style* class2
- *dihedral\_style* class2
- *improper\_style* class2
- *pair\_style* lj/class2
- *pair\_style* lj/class2/coul/cut
- *pair\_style* lj/class2/coul/long
- *special\_bonds* lj/coul 0 0 1

DREIDING is a generic force field developed by the [Goddard group](#) at Caltech and is useful for predicting structures and dynamics of organic, biological and main-group inorganic molecules. The philosophy in DREIDING is to use general force constants and geometry parameters based on simple hybridization considerations, rather than individual force constants and geometric parameters that depend on the particular combinations of atoms involved in the bond, angle, or torsion terms. DREIDING has an *explicit hydrogen bond term* to describe interactions involving a hydrogen atom on very electronegative atoms (N, O, F).

See (*Mayo*) for a description of the DREIDING force field

The interaction styles listed below compute force field formulas that are consistent with the DREIDING force field. See each command's documentation for the formula it computes.

- *bond\_style* harmonic
- *bond\_style* morse
- *angle\_style* cosine/squared
- *angle\_style* harmonic
- *angle\_style* cosine
- *angle\_style* cosine/periodic
- *dihedral\_style* charmm
- *improper\_style* umbrella
- *pair\_style* buck
- *pair\_style* buck/coul/cut
- *pair\_style* buck/coul/long
- *pair\_style* lj/cut
- *pair\_style* lj/cut/coul/cut
- *pair\_style* lj/cut/coul/long
- *pair\_style* hbond/dreiding/lj
- *pair\_style* hbond/dreiding/morse
- *special\_bonds* dreiding

---

(**MacKerell**) MacKerell, Bashford, Bellott, Dunbrack, Evanseck, Field, Fischer, Gao, Guo, Ha, et al, J Phys Chem, 102, 3586 (1998).

(**Cornell**) Cornell, Cieplak, Bayly, Gould, Merz, Ferguson, Spellmeyer, Fox, Caldwell, Kollman, JACS 117, 5179-5197 (1995).

(**Sun**) Sun, J. Phys. Chem. B, 102, 7338-7364 (1998).

(**Mayo**) Mayo, Olfason, Goddard III, J Phys Chem, 94, 8897-8909 (1990).

## 8.4.2 AMOEBA and HIPPO force fields

The AMOEBA and HIPPO polarizable force fields were developed by Jay Ponder’s group at the U Washington at St Louis. The LAMMPS implementation is based on Fortran 90 code provided by the Ponder group in their [Tinker MD software](#).

The current implementation (July 2022) of AMOEBA in LAMMPS matches the version discussed in (*Ponder*), (*Ren*), and (*Shi*). Likewise the current implementation of HIPPO in LAMMPS matches the version discussed in (*Rackers*).

These force fields can be used when polarization effects are desired in simulations of water, organic molecules, and biomolecules including proteins, provided that parameterizations (Tinker PRM force field files) are available for the systems you are interested in. Files in the LAMMPS potentials directory with a “amoeba” or “hippo” suffix can be used. The Tinker distribution and website have additional force field files as well: <https://github.com/TinkerTools/tinker/tree/release/params>.

Note that currently, HIPPO can only be used for water systems, but HIPPO files for a variety of small organic and biomolecules are in preparation by the Ponder group. Those force field files will be included in the LAMMPS distribution when available.

To use the AMOEBA or HIPPO force fields, a simulation must be 3d, and fully periodic or fully non-periodic, and use an orthogonal (not triclinic) simulation box.

The AMOEBA and HIPPO force fields contain the following terms in their energy ( $U$ ) computation. Further details for AMOEBA equations are in (*Ponder*), further details for the HIPPO equations are in (*Rackers*).

$$\begin{aligned}
 U &= U_{intermolecular} + U_{intramolecular} \\
 U_{intermolecular} &= U_{hal} + U_{repulsion} + U_{dispersion} + U_{multipole} + U_{polar} + U_{q\dot{x}fer} \\
 U_{intramolecular} &= U_{bond} + U_{angle} + U_{torsion} + U_{oop} + U_{b\theta} + U_{UB} + U_{pitorsion} + U_{bitorsion}
 \end{aligned}$$

For intermolecular terms, the AMOEBA force field includes only the  $U_{hal}$ ,  $U_{multipole}$ ,  $U_{polar}$  terms. The HIPPO force field includes all but the  $U_{hal}$  term. In LAMMPS, these are all computed by the *pair\_style amoeba or hippo* command. Note that the  $U_{multipole}$  and  $U_{polar}$  terms in this formula are not the same for the AMOEBA and HIPPO force fields.

For intramolecular terms, the  $U_{bond}$ ,  $U_{angle}$ ,  $U_{torsion}$ ,  $U_{oop}$  terms are computed by the *bond\_style class2 angle\_style amoeba, dihedral\_style fourier*, and *improper\_style amoeba* commands respectively. The *angle\_style amoeba* command includes the  $U_{b\theta}$  bond-angle cross term, and the  $U_{UB}$  term for a Urey-Bradley bond contribution between the I,K atoms in the IJK angle.

The  $U_{pitorsion}$  term is computed by the *fix amoeba/pitorsion* command. It computes 6-body interaction between a pair of bonded atoms which each have 2 additional bond partners.

The  $U_{bitorsion}$  term is computed by the *fix amoeba/bitorsion* command. It computes 5-body interaction between two 4-body torsions (dihedrals) which overlap, having 3 atoms in common.

These command doc pages have additional details on the terms they compute:

- *pair\_style amoeba or hippo*
- *bond\_style class2*
- *angle\_style amoeba*

- *dihedral\_style fourier*
- *improper\_style amoeba*
- *fix amoeba/pitortion*
- *fix amoeba/bitortion*

To use the AMOEBA or HIPPO force fields in LAMMPS, use commands like the following appropriately in your input script. The only change needed for AMOEBA vs HIPPO simulation is for the *pair\_style* and *pair\_coeff* commands, as shown below. See examples/amoeba for example input scripts for both AMOEBA and HIPPO.

```

units                real                                # required
atom_style           amoeba
bond_style           class2                            # CLASS2 package
angle_style          amoeba
dihedral_style       fourier                          # EXTRA-MOLECULE package
improper_style       amoeba

# required per-atom data
fix                  amtype all property/atom i_amtype ghost yes
fix                  extra all property/atom &
                    i_amgroup i_ired i_xaxis i_yaxis i_zaxis d_pval ghost yes
fix                  polaxe all property/atom i_polaxe

fix                  pit all amoeba/pitortion          # PiTorsion terms in FF
fix_modify           pit energy yes

# Bitorsion terms in FF
fix                  bit all amoeba/bitortion bitorsion.ubiquitin.data
fix_modify           bit energy yes

read_data            data.ubiquitin fix amtype NULL "Tinker Types" &
                    fix pit "pitortion types" "PiTorsion Coeffs" &
                    fix pit pitortions PiTortions &
                    fix bit bitortions BiTortions

pair_style           amoeba                            # AMOEBA FF
pair_coeff           * * amoeba_ubiquitin.prm amoeba_ubiquitin.key

pair_style           hippo                             # HIPPO FF
pair_coeff           * * hippo_water.prm hippo_water.key

special_bonds        lj/coul 0.5 0.5 0.5 one/five yes # 1-5 neighbors

```

The data file read by the *read\_data* command should be created by the tools/tinker/tinker2lmp.py conversion program described below. It will create a section in the data file with the header “Tinker Types”. A *fix property/atom* command for the data must be specified before the *read\_data* command. In the example above the fix ID is *amtype*.

Similarly, if the system you are simulating defines AMOEBA/HIPPO pitortion or bitortion interactions, there will be entries in the data file for those interactions. They require a *fix amoeba/pitortion* and *fix amoeba/bitortion* command to be defined. In the example above, the IDs for these two fixes are *pit* and *bit*.

Of course, if the system being modeled does not have one or more of the following – bond, angle, dihedral, improper, pitortion, bitortion interactions – then the corresponding style and fix commands above do not need to be used. See the example scripts in examples/amoeba for water systems as examples; they are simpler than what is listed above.



The two *fix property/atom* commands with IDs (in the example above) *extra* and *polaxe* are also needed to define internal per-atom quantities used by the AMOEBA and HIPPO force fields.

The *pair\_coeff* command used for either the AMOEBA or HIPPO force field takes two arguments for Tinker force field files, namely a PRM and KEY file. The keyfile can be specified as NULL and default values for a various settings will be used. Note that these 2 files are meant to allow use of native Tinker files as-is. However LAMMPS does not support all the options which can be included in a Tinker PRM or KEY file. See specifics below.

A *special\_bonds* command with the *one/five* option is required, since the AMOEBA/HIPPO force fields define weighting factors for not only 1-2, 1-3, 1-4 interactions, but also 1-5 interactions. This command will trigger a per-atom list of 1-5 neighbors to be generated. The AMOEBA and HIPPO force fields define their own custom weighting factors for all the 1-2, 1-3, 1-4, 1-5 terms which in the Tinker PRM and KEY files; they can be different for different terms in the force field.

In addition to the list above, these command doc pages have additional details:

- [\*atom\\_style amoeba\*](#)
  - [\*fix property/atom\*](#)
  - [\*special\\_bonds\*](#)
- 

Tinker PRM and KEY files

A Tinker PRM file is composed of sections, each of which has multiple lines. This is the list of PRM sections LAMMPS knows how to parse and use. Any other sections are skipped:

- Angle Bending Parameters
- Atom Type Definitions
- Atomic Multipole Parameters
- Bond Stretching Parameters
- Charge Penetration Parameters
- Charge Transfer Parameters
- Dipole Polarizability Parameters
- Dispersion Parameters
- Force Field Definition
- Literature References
- Out-of-Plane Bend Parameters
- Pauli Repulsion Parameters
- Pi-Torsion Parameters
- Stretch-Bend Parameters
- Torsion-Torsion Parameters
- Torsional Parameters
- Urey-Bradley Parameters
- Van der Waals Pair Parameters
- Van der Waals Parameters

A Tinker KEY file is composed of lines, each of which has a keyword followed by zero or more parameters. This is the list of keywords LAMMPS knows how to parse and use in the same manner Tinker does. Any other keywords are skipped. The value in parenthesis is the default value for the keyword if it is not specified, or if the keyfile in the *pair\_coeff* command is specified as NULL:

- a-axis (0.0)
- b-axis (0.0)
- c-axis (0.0)
- ctrn-cutoff (6.0)
- ctrn-taper (0.9 \* ctrn-cutoff)
- cutoff
- delta-halgren (0.07)
- dewald (no long-range dispersion unless specified)
- dewald-alpha (0.4)
- dewald-cutoff (7.0)
- dispersion-cutoff (9.0)
- dispersion-taper (9.0 \* dispersion-cutoff)
- dpme-grid
- dpme-order (4)
- ewald (no long-range electrostatics unless specified)
- ewald-alpha (0.4)
- ewald-cutoff (7.0)
- gamma-halgren (0.12)
- mpole-cutoff (9.0)
- mpole-taper (0.65 \* mpole-cutoff)
- pcg-guess (enabled by default)
- pcg-noguess (disable pcg-guess if specified)
- pcg-noprecond (disable pcg-precond if specified)
- pcg-peek (1.0)
- pcg-precond (enabled by default)
- pewald-alpha (0.4)
- pme-grid
- pme-order (5)
- polar-eps (1.0e-6)
- polar-iter (100)
- polar-predict (no prediction operation unless specified)
- ppme-order (5)
- repulsion-cutoff (6.0)

- repulsion-taper (0.9 \* repulsion-cutoff)
- taper
- usolve-cutoff (4.5)
- usolve-diag (2.0)
- vdw-cutoff (9.0)
- vdw-taper (0.9 \* vdw-cutoff)

### Tinker2lmp.py tool

This conversion tool is found in the tools/tinker directory. As shown in examples/amoeba/README, these commands produce the data files found in examples/amoeba, and also illustrate all the options available to use with the tinker2lmp.py script:

```
% python tinker2lmp.py -xyz water_dimer.xyz -amoeba amoeba_water.prm -data data.water_
↳dimer.amoeba           # AMOEBA non-periodic system
% python tinker2lmp.py -xyz water_dimer.xyz -hippo hippo_water.prm -data data.water_
↳dimer.hippo           # HIPPO non-periodic system
% python tinker2lmp.py -xyz water_box.xyz -amoeba amoeba_water.prm -data data.water_box.
↳amoeba -pbc 18.643 18.643 18.643 # AMOEBA periodic system
% python tinker2lmp.py -xyz water_box.xyz -hippo hippo_water.prm -data data.water_box.
↳hippo -pbc 18.643 18.643 18.643 # HIPPO periodic system
% python tinker2lmp.py -xyz ubiquitin.xyz -amoeba amoeba_ubiquitin.prm -data data.
↳ubiquitin.new -pbc 54.99 41.91 41.91 -bitorsion bitorsion.ubiquitin.data.new #
↳system with bitorsions
```

Switches and their arguments may be specified in any order.

The -xyz switch is required and specifies an input XYZ file as an argument. The format of this file is an extended XYZ format defined and used by Tinker for its input. Example \*.xyz files are in the examples/amoeba directory. The file lists the atoms in the system. Each atom has the following information: Tinker species name (ignored by LAMMPS), xyz coordinates, Tinker numeric type, and a list of atom IDs the atom is bonded to.

Here is more information about the extended XYZ format defined and used by Tinker, and links to programs that convert standard PDB files to the extended XYZ format:

- [http://openbabel.org/docs/current/FileFormats/Tinker\\_XYZ\\_format.html](http://openbabel.org/docs/current/FileFormats/Tinker_XYZ_format.html)
- <https://github.com/emleddin/pdbxyz-xyzpdb>
- <https://github.com/TinkerTools/tinker/blob/release/source/pdbxyz.f>

The -amoeba or -hippo switch is required. It specifies an input AMOEBA or HIPPO PRM force field file as an argument. This should be the same file used by the *pair\_style* command in the input script.

The -data switch is required. It specifies an output file name for the LAMMPS data file that will be produced.

For periodic systems, the -pbc switch is required. It specifies the periodic box size for each dimension (x,y,z). For a Tinker simulation these are specified in the KEY file.

The -bitorsion switch is only needed if the system contains Tinker bitorsion interactions. The data for each type of bitorsion interaction will be written to the specified file, and read by the *fix amoeba/bitorsion* command. The data includes 2d arrays of values to which splines are fit, and thus is not compatible with the LAMMPS data file format.

---

**(Ponder)** Ponder, Wu, Ren, Pande, Chodera, Schnieders, Haque, Mobley, Lambrecht, DiStasio Jr, M. Head-Gordon, Clark, Johnson, T. Head-Gordon, J Phys Chem B, 114, 2549-2564 (2010).

**(Rackers)** Rackers, Silva, Wang, Ponder, J Chem Theory Comput, 17, 7056-7084 (2021).

**(Ren)** Ren and Ponder, J Phys Chem B, 107, 5933 (2003).

**(Shi)** Shi, Xia, Zhang, Best, Wu, Ponder, Ren, J Chem Theory Comp, 9, 4046, 2013.

### 8.4.3 TIP3P water model

The TIP3P water model as implemented in CHARMM (*MacKerell*) specifies a 3-site rigid water molecule with charges and Lennard-Jones parameters assigned to each of the 3 atoms. In LAMMPS the *fix shake* command can be used to hold the two O-H bonds and the H-O-H angle rigid. A bond style of *harmonic* and an angle style of *harmonic* or *charmm* should also be used.

These are the additional parameters (in real units) to set for O and H atoms and the water molecule to run a rigid TIP3P-CHARMM model with a cutoff. The K values can be used if a flexible TIP3P model (without fix shake) is desired. If the LJ epsilon and sigma for HH and OH are set to 0.0, it corresponds to the original 1983 TIP3P model (*Jorgensen*).

O mass = 15.9994  
H mass = 1.008  
O charge = -0.834  
H charge = 0.417  
LJ  $\epsilon$  of OO = 0.1521  
LJ  $\sigma$  of OO = 3.1507  
LJ  $\epsilon$  of HH = 0.0460  
LJ  $\sigma$  of HH = 0.4000  
LJ  $\epsilon$  of OH = 0.0836  
LJ  $\sigma$  of OH = 1.7753  
K of OH bond = 450  
 $r_0$  of OH bond = 0.9572  
K of HOH angle = 55  
 $\theta$  of HOH angle = 104.52°

These are the parameters to use for TIP3P with a long-range Coulomb solver (e.g. Ewald or PPPM in LAMMPS), see (*Price*) for details:

O mass = 15.9994  
H mass = 1.008  
O charge = -0.830  
H charge = 0.415  
LJ  $\epsilon$  of OO = 0.102  
LJ  $\sigma$  of OO = 3.188  
LJ  $\epsilon, \sigma$  of OH, HH = 0.0  
K of OH bond = 450  
 $r_0$  of OH bond = 0.9572  
K of HOH angle = 55  
 $\theta$  of HOH angle = 104.52°

Wikipedia also has a nice article on [water models](#).

**(MacKerell)** MacKerell, Bashford, Bellott, Dunbrack, Evanseck, Field, Fischer, Gao, Guo, Ha, et al, J Phys Chem, 102, 3586 (1998).

**(Jorgensen)** Jorgensen, Chandrasekhar, Madura, Impey, Klein, J Chem Phys, 79, 926 (1983).

**(Price)** Price and Brooks, J Chem Phys, 121, 10096 (2004).

#### 8.4.4 TIP4P water model

The four-point TIP4P rigid water model extends the traditional three-point TIP3P model by adding an additional site, usually massless, where the charge associated with the oxygen atom is placed. This site M is located at a fixed distance away from the oxygen along the bisector of the HOH bond angle. A bond style of *harmonic* and an angle style of *harmonic* or *charmm* should also be used.

A TIP4P model is run with LAMMPS using either this command for a cutoff model:

- `pair_style lj/cut/tip4p/cut`

or these two commands for a long-range model:

- `pair_style lj/cut/tip4p/long`
- `kspace_style ppm/tip4p`

For both models, the bond lengths and bond angles should be held fixed using the *fix shake* command.

These are the additional parameters (in real units) to set for O and H atoms and the water molecule to run a rigid TIP4P model with a cutoff (*Jorgensen*). Note that the OM distance is specified in the *pair\_style* command, not as part of the pair coefficients.

```
O mass = 15.9994
H mass = 1.008
O charge = -1.040
H charge = 0.520
r0 of OH bond = 0.9572
θ of HOH angle = 104.52°
OM distance = 0.15
LJ ε of O-O = 0.1550
LJ σ of O-O = 3.1536
LJ ε, σ of OH, HH = 0.0
Coulomb cutoff = 8.5
```

For the TIP4/Ice model (J Chem Phys, 122, 234511 (2005); <https://doi.org/10.1063/1.1931662>) these values can be used:

```
O mass = 15.9994
H mass = 1.008
O charge = -1.1794
```

H charge = 0.5897  
 $r_0$  of OH bond = 0.9572  
 $\theta$  of HOH angle = 104.52°  
OM distance = 0.1577  
LJ  $\epsilon$  of O-O = 0.21084  
LJ  $\sigma$  of O-O = 3.1668  
LJ  $\epsilon$ ,  $\sigma$  of OH, HH = 0.0  
Coulomb cutoff = 8.5

For the TIP4P/2005 model (J Chem Phys, 123, 234505 (2005); <https://doi.org/10.1063/1.2121687>), these values can be used:

O mass = 15.9994  
H mass = 1.008  
O charge = -1.1128  
H charge = 0.5564  
 $r_0$  of OH bond = 0.9572  
 $\theta$  of HOH angle = 104.52°  
OM distance = 0.1546  
LJ  $\epsilon$  of O-O = 0.1852  
LJ  $\sigma$  of O-O = 3.1589  
LJ  $\epsilon$ ,  $\sigma$  of OH, HH = 0.0  
Coulomb cutoff = 8.5

These are the parameters to use for TIP4P with a long-range Coulombic solver (e.g. Ewald or PPPM in LAMMPS):

O mass = 15.9994  
H mass = 1.008  
O charge = -1.0484  
H charge = 0.5242  
 $r_0$  of OH bond = 0.9572  
 $\theta$  of HOH angle = 104.52°  
OM distance = 0.1250  
LJ  $\epsilon$  of O-O = 0.16275  
LJ  $\sigma$  of O-O = 3.16435  
LJ  $\epsilon$ ,  $\sigma$  of OH, HH = 0.0

Note that when using the TIP4P pair style, the neighbor list cutoff for Coulomb interactions is effectively extended by a distance  $2 * (\text{OM distance})$ , to account for the offset distance of the fictitious charges on O atoms in water molecules. Thus it is typically best in an efficiency sense to use a LJ cutoff  $\geq$  Coulomb cutoff +  $2 * (\text{OM distance})$ , to shrink the size of the neighbor list. This leads to slightly larger cost for the long-range calculation, so you can test the trade-off for your model. The OM distance and the LJ and Coulombic cutoffs are set in the *pair\_style lj/cut/tip4p/long* command.

Wikipedia also has a nice article on [water models](#).

---

(**Jorgensen**) Jorgensen, Chandrasekhar, Madura, Impey, Klein, J Chem Phys, 79, 926 (1983).

### 8.4.5 SPC water model

The SPC water model specifies a 3-site rigid water molecule with charges and Lennard-Jones parameters assigned to each of the 3 atoms. In LAMMPS the *fix shake* command can be used to hold the two O-H bonds and the H-O-H angle rigid. A bond style of *harmonic* and an angle style of *harmonic* or *charmm* should also be used.

These are the additional parameters (in real units) to set for O and H atoms and the water molecule to run a rigid SPC model.

O mass = 15.9994  
H mass = 1.008  
O charge = -0.820  
H charge = 0.410  
LJ  $\epsilon$  of OO = 0.1553  
LJ  $\sigma$  of OO = 3.166  
LJ  $\epsilon$ ,  $\sigma$  of OH, HH = 0.0  
 $r_0$  of OH bond = 1.0  
 $\theta_0$  of HOH angle = 109.47°

Note that as originally proposed, the SPC model was run with a 9 Angstrom cutoff for both LJ and Coulomb terms. It can also be used with long-range electrostatic solvers (e.g. Ewald or PPPM in LAMMPS) without changing any of the parameters above, although it becomes a different model in that mode of usage.

The SPC/E (extended) water model is the same, except the partial charge assignments change:

O charge = -0.8476  
H charge = 0.4238

See the (*Berendsen*) reference for more details on both the SPC and SPC/E models.

Wikipedia also has a nice article on [water models](#).

---

(**Berendsen**) Berendsen, Grigera, Straatsma, J Phys Chem, 91, 6269-6271 (1987).

## 8.5 Packages howto

### 8.5.1 Finite-size spherical and aspherical particles

Typical MD models treat atoms or particles as point masses. Sometimes it is desirable to have a model with finite-size particles such as spheroids or ellipsoids or generalized aspherical bodies. The difference is that such particles have a moment of inertia, rotational energy, and angular momentum. Rotation is induced by torque coming from interactions with other particles.

LAMMPS has several options for running simulations with these kinds of particles. The following aspects are discussed in turn:

- atom styles
- pair potentials
- time integration
- computes, thermodynamics, and dump output
- rigid bodies composed of finite-size particles

Example input scripts for these kinds of models are in the `body`, `colloid`, `dipole`, `ellipse`, `line`, `peri`, `pour`, and `tri` directories of the *examples directory* in the LAMMPS distribution.

#### Atom styles

There are several *atom styles* that allow for definition of finite-size particles: `sphere`, `dipole`, `ellipsoid`, `line`, `tri`, `peri`, and `body`.

The `sphere` style defines particles that are spheroids and each particle can have a unique diameter and mass (or density). These particles store an angular velocity ( $\omega$ ) and can be acted upon by torque. The “set” command can be used to modify the diameter and mass of individual particles, after then are created.

The `dipole` style does not actually define finite-size particles, but is often used in conjunction with spherical particles, via a command like

```
atom_style hybrid sphere dipole
```

This is because when dipoles interact with each other, they induce torques, and a particle must be finite-size (i.e. have a moment of inertia) in order to respond and rotate. See the *atom\_style dipole* command for details. The “set” command can be used to modify the orientation and length of the dipole moment of individual particles, after then are created.

The `ellipsoid` style defines particles that are ellipsoids and thus can be aspherical. Each particle has a shape, specified by 3 diameters, and mass (or density). These particles store an angular momentum and their orientation (quaternion), and can be acted upon by torque. They do not store an angular velocity ( $\omega$ ), which can be in a different direction than angular momentum, rather they compute it as needed. The “set” command can be used to modify the diameter, orientation, and mass of individual particles, after then are created. It also has a brief explanation of what quaternions are.

The `line` style defines line segment particles with two end points and a mass (or density). They can be used in 2d simulations, and they can be joined together to form rigid bodies which represent arbitrary polygons.

The `tri` style defines triangular particles with three corner points and a mass (or density). They can be used in 3d simulations, and they can be joined together to form rigid bodies which represent arbitrary particles with a triangulated surface.

The `peri` style is used with *Peridynamic models* and defines particles as having a volume, that is used internally in the *pair\_style peri* potentials.



The body style allows for definition of particles which can represent complex entities, such as surface meshes of discrete points, collections of sub-particles, deformable objects, etc. The body style is discussed in more detail on the *Howto body* doc page.

Note that if one of these atom styles is used (or multiple styles via the *atom\_style hybrid* command), not all particles in the system are required to be finite-size or aspherical.

For example, in the ellipsoid style, if the 3 shape parameters are set to the same value, the particle will be a sphere rather than an ellipsoid. If the 3 shape parameters are all set to 0.0 or if the diameter is set to 0.0, it will be a point particle. In the line or tri style, if the lineflag or triflag is specified as 0, then it will be a point particle.

Some of the pair styles used to compute pairwise interactions between finite-size particles also compute the correct interaction with point particles as well, e.g. the interaction between a point particle and a finite-size particle or between two point particles. If necessary, *pair\_style hybrid* can be used to insure the correct interactions are computed for the appropriate style of interactions. Likewise, using groups to partition particles (ellipsoids versus spheres versus point particles) will allow you to use the appropriate time integrators and temperature computations for each class of particles. See the doc pages for various commands for details.

Also note that for *2d simulations*, atom styles sphere and ellipsoid still use 3d particles, rather than as circular disks or ellipses. This means they have the same moment of inertia as the 3d object. When temperature is computed, the correct degrees of freedom are used for rotation in a 2d versus 3d system.

## Pair potentials

When a system with finite-size particles is defined, the particles will only rotate and experience torque if the force field computes such interactions. These are the various *pair styles* that generate torque:

- *pair\_style gran/history*
- *pair\_style gran/hertz*
- *pair\_style gran/no\_history*
- *pair\_style dipole/cut*
- *pair\_style gayberne*
- *pair\_style resquared*
- *pair\_style brownian*
- *pair\_style lubricate*
- *pair\_style line/lj*
- *pair\_style tri/lj*
- *pair\_style body/nparticle*

The granular pair styles are used with spherical particles. The dipole pair style is used with the dipole atom style, which could be applied to spherical or ellipsoidal particles. The GayBerne and REsquared potentials require ellipsoidal particles, though they will also work if the 3 shape parameters are the same (a sphere). The Brownian and lubrication potentials are used with spherical particles. The line, tri, and body potentials are used with line segment, triangular, and body particles respectively.

## Time integration

There are several fixes that perform time integration on finite-size spherical particles, meaning the integrators update the rotational orientation and angular velocity or angular momentum of the particles:

- *fix nve/sphere*
- *fix nvt/sphere*
- *fix npt/sphere*

Likewise, there are 3 fixes that perform time integration on ellipsoidal particles:

- *fix nve/asphere*
- *fix nvt/asphere*
- *fix npt/asphere*

The advantage of these fixes is that those which thermostat the particles include the rotational degrees of freedom in the temperature calculation and thermostatting. The *fix langevin* command can also be used with its *omega* or *angmom* options to thermostat the rotational degrees of freedom for spherical or ellipsoidal particles. Other thermostatting fixes only operate on the translational kinetic energy of finite-size particles.

These fixes perform constant NVE time integration on line segment, triangular, and body particles:

- *fix nve/line*
- *fix nve/tri*
- *fix nve/body*

Note that for mixtures of point and finite-size particles, these integration fixes can only be used with *groups* which contain finite-size particles.

## Computes, thermodynamics, and dump output

There are several computes that calculate the temperature or rotational energy of spherical or ellipsoidal particles:

- *compute temp/sphere*
- *compute temp/asphere*
- *compute erotate/sphere*
- *compute erotate/asphere*

These include rotational degrees of freedom in their computation. If you wish the thermodynamic output of temperature or pressure to use one of these computes (e.g. for a system entirely composed of finite-size particles), then the compute can be defined and the *thermo\_modify* command used. Note that by default thermodynamic quantities will be calculated with a temperature that only includes translational degrees of freedom. See the *thermo\_style* command for details.

These commands can be used to output various attributes of finite-size particles:

- *dump custom*
- *compute property/atom*
- *dump local*
- *compute body/local*

Attributes include the dipole moment, the angular velocity, the angular momentum, the quaternion, the torque, the end-point and corner-point coordinates (for line and tri particles), and sub-particle attributes of body particles.

## Rigid bodies composed of finite-size particles

The *fix rigid* command treats a collection of particles as a rigid body, computes its inertia tensor, sums the total force and torque on the rigid body each timestep due to forces on its constituent particles, and integrates the motion of the rigid body.

If any of the constituent particles of a rigid body are finite-size particles (spheres or ellipsoids or line segments or triangles), then their contribution to the inertia tensor of the body is different than if they were point particles. This means the rotational dynamics of the rigid body will be different. Thus a model of a dimer is different if the dimer consists of two point masses versus two spheroids, even if the two particles have the same mass. Finite-size particles that experience torque due to their interaction with other particles will also impart that torque to a rigid body they are part of.

See the “fix rigid” command for example of complex rigid-body models it is possible to define in LAMMPS.

Note that the *fix shake* command can also be used to treat 2, 3, or 4 particles as a rigid body, but it always assumes the particles are point masses.

Also note that body particles cannot be modeled with the *fix rigid* command. Body particles are treated by LAMMPS as single particles, though they can store internal state, such as a list of sub-particles. Individual body particles are typically treated as rigid bodies, and their motion integrated with a command like *fix nve/body*. Interactions between pairs of body particles are computed via a command like *pair\_style body/nparticle*.

### 8.5.2 Granular models

Granular system are composed of spherical particles with a diameter, as opposed to point particles. This means they have an angular velocity and torque can be imparted to them to cause them to rotate.

To run a simulation of a granular model, you will want to use the following commands:

- *atom\_style sphere*
- *fix nve/sphere*
- *fix gravity*

This compute

- *compute erotate/sphere*

calculates rotational kinetic energy which can be *output with thermodynamic info*. The compute

- *compute fabric*

calculates various versions of the fabric tensor for granular and non-granular pair styles.

Use one of these 4 pair potentials, which compute forces and torques between interacting pairs of particles:

- *pair\_style gran/history*
- *pair\_style gran/no\_history*
- *pair\_style gran/hertzian*
- *pair\_style granular*

These commands implement fix options specific to granular systems:

- *fix freeze*
- *fix pour*
- *fix viscous*

- *fix wall/gran*
- *fix wall/gran/region*

The fix style *freeze* zeroes both the force and torque of frozen atoms, and should be used for granular system instead of the fix style *setforce*.

For computational efficiency, you can eliminate needless pairwise computations between frozen atoms by using this command:

- *neigh\_modify* exclude

---

**Note:** By default, for 2d systems, granular particles are still modeled as 3d spheres, not 2d discs (circles), meaning their moment of inertia will be the same as in 3d. If you wish to model granular particles in 2d as 2d discs, see the note on this topic on the [Howto 2d](#) doc page, where 2d simulations are discussed.

---

### 8.5.3 Body particles

#### Overview:

In LAMMPS, body particles are generalized finite-size particles. Individual body particles can represent complex entities, such as surface meshes of discrete points, collections of sub-particles, deformable objects, etc. Note that other kinds of finite-size spherical and aspherical particles are also supported by LAMMPS, such as spheres, ellipsoids, line segments, and triangles, but they are simpler entities than body particles. See the [Howto spherical](#) page for a general overview of all these particle types.

Body particles are used via the *atom\_style body* command. It takes a body style as an argument. The current body styles supported by LAMMPS are as follows. The name in the first column is used as the *bstyle* argument for the *atom\_style body* command.

<i>nparticle</i>	rigid body with N sub-particles
<i>rounded/polygon</i>	2d polygons with N vertices
<i>rounded/polyhedron</i>	3d polyhedra with N vertices, E edges and F faces

The body style determines what attributes are stored for each body and thus how they can be used to compute pairwise body/body or bond/non-body (point particle) interactions. More details of each style are described below.

More styles may be added in the future. See the [page on creating new body styles](#) for details on how to add a new body style to the code.

---

#### When to use body particles:

You should not use body particles to model a rigid body made of simpler particles (e.g. point, sphere, ellipsoid, line segment, triangular particles), if the interaction between pairs of rigid bodies is just the summation of pairwise interactions between the simpler particles. LAMMPS already supports this kind of model via the *fix rigid* command. Any of the numerous pair styles that compute interactions between simpler particles can be used. The *fix rigid* command time integrates the motion of the rigid bodies. All of the standard LAMMPS commands for thermostating, adding constraints, performing output, etc will operate as expected on the simple particles.

By contrast, when body particles are used, LAMMPS treats an entire body as a single particle for purposes of computing pairwise interactions, building neighbor lists, migrating particles between processors, output of particles to a dump file, etc. This means that interactions between pairs of bodies or between a body and non-body (point) particle need to be encoded in an appropriate pair style. If such a pair style were to mimic the *fix rigid* model, it would need to loop over the entire collection of interactions between pairs of simple particles within the two bodies, each time a single body/body interaction was computed.

Thus it only makes sense to use body particles and develop such a pair style, when particle/particle interactions are more complex than what the *fix rigid* command can already calculate. For example, consider particles with one or more of the following attributes:

- represented by a surface mesh
- represented by a collection of geometric entities (e.g. planes + spheres)
- deformable
- internal stress that induces fragmentation

For these models, the interaction between pairs of particles is likely to be more complex than the summation of simple pairwise interactions. An example is contact or frictional forces between particles with planar surfaces that interpenetrate. Likewise, the body particle may store internal state, such as a stress tensor used to compute a fracture criterion.

These are additional LAMMPS commands that can be used with body particles of different styles

<i>fix nve/body</i>	integrate motion of a body particle in NVE ensemble
<i>fix nvt/body</i>	ditto for NVT ensemble
<i>fix npt/body</i>	ditto for NPT ensemble
<i>fix nph/body</i>	ditto for NPH ensemble
<i>compute body/local</i>	store sub-particle attributes of a body particle
<i>compute temp/body</i>	compute temperature of body particles
<i>dump local</i>	output sub-particle attributes of a body particle
<i>dump image</i>	output body particle attributes as an image

The pair styles defined for use with specific body styles are listed in the sections below.

### Specifics of body style *nparticle*:

The *nparticle* body style represents body particles as a rigid body with a variable number  $N$  of sub-particles. It is provided as a vanilla, prototypical example of a body particle, although as mentioned above, the *fix rigid* command already duplicates its functionality.

The *atom\_style* body command for this body style takes two additional arguments:

```
atom_style body nparticle Nmin Nmax
Nmin = minimum # of sub-particles in any body in the system
Nmax = maximum # of sub-particles in any body in the system
```

The  $Nmin$  and  $Nmax$  arguments are used to bound the size of data structures used internally by each particle.

When the *read\_data* command reads a data file for this body style, the following information must be provided for each entry in the *Bodies* section of the data file:

```
atom-ID 1 M
N
ixx iyy izz ixz iyz
x1 y1 z1
...
xN yN zN
```

where  $M = 6 + 3*N$ , and  $N$  is the number of sub-particles in the body particle.

The integer line has a single value  $N$ . The floating point line(s) list 6 moments of inertia followed by the coordinates of the  $N$  sub-particles ( $x_1$  to  $z_N$ ) as  $3N$  values. These values can be listed on as many lines as you wish; see the [read\\_data](#) command for more details.

The 6 moments of inertia ( $ixx,iyy,izz,ixy,ixz,iyz$ ) should be the values consistent with the current orientation of the rigid body around its center of mass. The values are with respect to the simulation box XYZ axes, not with respect to the principal axes of the rigid body itself. LAMMPS performs the latter calculation internally. The coordinates of each sub-particle are specified as its  $x,y,z$  displacement from the center-of-mass of the body particle. The center-of-mass position of the particle is specified by the  $x,y,z$  values in the *Atoms* section of the data file, as is the total mass of the body particle.

The [pair\\_style body/nparticle](#) command can be used with this body style to compute body/body and body/non-body interactions.

---

### Specifics of body style rounded/polygon:

The *rounded/polygon* body style represents body particles as a 2d polygon with a variable number of  $N$  vertices. This style can only be used for 2d models; see the [boundary](#) command. See the “[pair\\_style body/rounded/polygon](#)” page for a diagram of two squares with rounded circles at the vertices. Special cases for  $N = 1$  (circle) and  $N = 2$  (rod with rounded ends) can also be specified.

One use of this body style is for 2d discrete element models, as described in [Fraige](#).

Similar to body style *nparticle*, the `atom_style` body command for this body style takes two additional arguments:

```
atom_style body rounded/polygon Nmin Nmax
Nmin = minimum # of vertices in any body in the system
Nmax = maximum # of vertices in any body in the system
```

The  $Nmin$  and  $Nmax$  arguments are used to bound the size of data structures used internally by each particle.

When the [read\\_data](#) command reads a data file for this body style, the following information must be provided for each body in the *Bodies* section of the data file:

```
atom-ID 1 M
N
ixx iyy izz ixy ixz iyz
x1 y1 z1
...
xN yN zN
diameter
```

where  $M = 6 + 3*N + 1$ , and  $N$  is the number of vertices in the body particle.

The integer line has a single value  $N$ . The floating point line(s) list 6 moments of inertia, followed by the coordinates of the  $N$  vertices ( $x_1$  to  $z_N$ ) as  $3N$  values (with  $z = 0.0$  for each), followed by a diameter value = the rounded diameter of the circle that surrounds each vertex. The diameter value can be different for each body particle. These floating-point values can be listed on as many lines as you wish; see the [read\\_data](#) command for more details.

---

**Note:** It is important that the vertices for each polygonal body particle be listed in order around its perimeter, so that edges can be inferred. LAMMPS does not check that this is the case.

---

The 6 moments of inertia ( $ixx,iyy,izz,ixy,ixz,iyz$ ) should be the values consistent with the current orientation of the rigid body around its center of mass. The values are with respect to the simulation box XYZ axes, not with respect to the principal axes of the rigid body itself. LAMMPS performs the latter calculation internally. The coordinates of each

vertex are specified as its x,y,z displacement from the center-of-mass of the body particle. The center-of-mass position of the particle is specified by the x,y,z values in the *Atoms* section of the data file.

For example, the following information would specify a square particle whose edge length is  $\sqrt{2}$  and rounded diameter is 1.0. The orientation of the square is aligned with the xy coordinate axes which is consistent with the 6 moments of inertia:  $ixx\ iyy\ izz\ ixy\ ixz\ iyz = 1\ 1\ 4\ 0\ 0\ 0$ . Note that only *Izz* matters in 2D simulations.

```
3 1 19
4
1 1 4 0 0 0
-0.7071 -0.7071 0
-0.7071 0.7071 0
0.7071 0.7071 0
0.7071 -0.7071 0
1.0
```

A rod in 2D, whose length is 4.0, mass 1.0, rounded at two ends by circles of diameter 0.5, is specified as follows:

```
1 1 13
2
1 1 1.33333 0 0 0
-2 0 0
2 0 0
0.5
```

A disk, whose diameter is 3.0, mass 1.0, is specified as follows:

```
1 1 10
1
1 1 4.5 0 0 0
0 0 0
3.0
```

The *pair\_style body/rounded/polygon* command can be used with this body style to compute body/body interactions. The *fix wall/body/polygon* command can be used with this body style to compute the interaction of body particles with a wall.

### Specifics of body style rounded/polyhedron:

The *rounded/polyhedron* body style represents body particles as a 3d polyhedron with a variable number of *N* vertices, *E* edges and *F* faces. This style can only be used for 3d models; see the *boundary* command. See the “pair\_style body/rounded/polygon” page for a diagram of a two 2d squares with rounded circles at the vertices. A 3d cube with rounded spheres at the 8 vertices and 12 rounded edges would be similar. Special cases for *N* = 1 (sphere) and *N* = 2 (rod with rounded ends) can also be specified.

This body style is for 3d discrete element models, as described in *Wang*.

Similar to body style *rounded/polygon*, the *atom\_style body* command for this body style takes two additional arguments:

```
atom_style body rounded/polyhedron Nmin Nmax
Nmin = minimum # of vertices in any body in the system
Nmax = maximum # of vertices in any body in the system
```

The *Nmin* and *Nmax* arguments are used to bound the size of data structures used internally by each particle.

When the `read_data` command reads a data file for this body style, the following information must be provided for each entry in the *Bodies* section of the data file:

```
atom-ID 3 M
N E F
ixx iyy izz ixy ixz iyz
x1 y1 z1
...
xN yN zN
0 1
1 2
2 3
...
0 1 2 -1
0 2 3 -1
...
1 2 3 4
diameter
```

where  $M = 6 + 3*N + 2*E + 4*F + 1$ , and  $N$  is the number of vertices in the body particle,  $E$  = number of edges,  $F$  = number of faces. For  $N = 1$  or  $2$ , the format is simpler.  $E$  and  $F$  are ignored and no edges or faces are listed, so that  $M = 6 + 3*N + 1$ .

The integer line has three values: number of vertices ( $N$ ), number of edges ( $E$ ) and number of faces ( $F$ ). The floating point line(s) list 6 moments of inertia followed by the coordinates of the  $N$  vertices ( $x1$  to  $zN$ ) as  $3N$  values, followed by  $2N$  vertex indices corresponding to the end points of the  $E$  edges, then  $4*F$  vertex indices defining  $F$  faces. The last value is the diameter value = the rounded diameter of the sphere that surrounds each vertex. The diameter value can be different for each body particle. These floating-point values can be listed on as many lines as you wish; see the `read_data` command for more details.

Note that vertices are numbered from 0 to  $N-1$  inclusive. The order of the 2 vertices in each edge does not matter. Faces can be triangles or quadrilaterals. In both cases 4 vertices must be specified. For a triangle the 4th vertex is -1. The 4 vertices within each triangle or quadrilateral face should be ordered by the right-hand rule so that the normal vector of the face points outwards from the center of mass. For polyhedron with faces with more than 4 vertices, you should split the complex face into multiple simple faces, each of which is a triangle or quadrilateral.

**Note:** If a face is a quadrilateral then its 4 vertices must be co-planar. LAMMPS does not check that this is the case. If you have a quad-face of a polyhedron that is not planar (e.g. a cube whose vertices have been randomly displaced), then you should represent the single quad face as two triangle faces instead.

The 6 moments of inertia ( $ixx,iyy,izz,ixy,ixz,iyz$ ) should be the values consistent with the current orientation of the rigid body around its center of mass. The values are with respect to the simulation box  $XYZ$  axes, not with respect to the principal axes of the rigid body itself. LAMMPS performs the latter calculation internally. The coordinates of each vertex are specified as its  $x,y,z$  displacement from the center-of-mass of the body particle. The center-of-mass position of the particle is specified by the  $x,y,z$  values in the *Atoms* section of the data file.

For example, the following information would specify a cubic particle whose edge length is 2.0 and rounded diameter is 0.5. The orientation of the cube is aligned with the  $xyz$  coordinate axes which is consistent with the 6 moments of inertia:  $ixx iyy izz ixy ixz iyz = 0.667 0.667 0.667 0 0 0$ .

```
1 3 79
8 12 6
0.667 0.667 0.667 0 0 0
1 1 1
```

(continues on next page)



(continued from previous page)

```

1 -1 1
-1 -1 1
-1 1 1
1 1 -1
1 -1 -1
-1 -1 -1
-1 1 -1
0 1
1 2
2 3
3 0
4 5
5 6
6 7
7 4
0 4
1 5
2 6
3 7
0 1 2 3
4 5 6 7
0 1 5 4
1 2 6 5
2 3 7 6
3 0 4 7
0.5

```

A rod in 3D, whose length is 4.0, mass 1.0 and rounded at two ends by circles of diameter 0.5, is specified as follows:

```

1 3 13
2 1 1
0 1.33333 1.33333 0 0 0
-2 0 0
2 0 0
0.5

```

A sphere whose diameter is 3.0 and mass 1.0, is specified as follows:

```

1 3 10
1 1 1
0.9 0.9 0.9 0 0 0
0 0 0
3.0

```

The number of edges and faces for a rod or sphere must be listed, but is ignored.

The *pair\_style body/rounded/polhedron* command can be used with this body style to compute body/body interactions. The *fix wall/body/polyhedron* command can be used with this body style to compute the interaction of body particles with a wall.

### Output specifics for all body styles:

For the *compute body/local* and *dump local* commands, all 3 of the body styles described on this page produces one datum for each of the N vertices (of sub-particles) in a body particle. The datum has 3 values:

1 = x position of vertex (or sub-particle) 2 = y position of vertex 3 = z position of vertex
--

These values are the current position of the vertex within the simulation domain, not a displacement from the center-of-mass (COM) of the body particle itself. These values are calculated using the current COM and orientation of the body particle.

The *dump image* command and its *body* keyword can be used to render body particles.

For the *nparticle* body style, each body is drawn as a collection of spheres, one for each sub-particle. The size of each sphere is determined by the *bflag1* parameter for the *body* keyword. The *bflag2* argument is ignored.

For the *rounded/polygon* body style, each body is drawn as a polygon with N line segments. For the *rounded/polyhedron* body style, each face of each body is drawn as a polygon with N line segments. The drawn diameter of each line segment is determined by the *bflag1* parameter for the *body* keyword. The *bflag2* argument is ignored.

Note that for both the *rounded/polygon* and *rounded/polyhedron* styles, line segments are drawn between the pairs of vertices. Depending on the diameters of the line segments this may be slightly different than the physical extent of the body as calculated by the *pair\_style rounded/polygon* or *pair\_style rounded/polyhedron* commands. Conceptually, the pair styles define the surface of a 2d or 3d body by lines or planes that are tangent to the finite-size spheres of specified diameter which are placed on each vertex position.

---

**(Fraige)** F. Y. Fraige, P. A. Langston, A. J. Matchett, J. Dodds, *Particuology*, 6, 455 (2008).

**(Wang)** J. Wang, H. S. Yu, P. A. Langston, F. Y. Fraige, *Granular Matter*, 13, 1 (2011).

## 8.5.4 Polarizable models

In polarizable force fields the charge distributions in molecules and materials respond to their electrostatic environments. Polarizable systems can be simulated in LAMMPS using three methods:

- the fluctuating charge method, implemented in the *QEQ* package,
- the adiabatic core-shell method, implemented in the *CORESHELL* package,
- the thermalized Drude dipole method, implemented in the *DRUDE* package.

The fluctuating charge method calculates instantaneous charges on interacting atoms based on the electronegativity equalization principle. It is implemented in the *fix qeq* which is available in several variants. It is a relatively efficient technique since no additional particles are introduced. This method allows for charge transfer between molecules or atom groups. However, because the charges are located at the interaction sites, off-plane components of polarization cannot be represented in planar molecules or atom groups.

The two other methods share the same basic idea: polarizable atoms are split into one core atom and one satellite particle (called shell or Drude particle) attached to it by a harmonic spring. Both atoms bear a charge and they represent collectively an induced electric dipole. These techniques are computationally more expensive than the QEq method because of additional particles and bonds. These two charge-on-spring methods differ in certain features, with the core-shell model being normally used for ionic/crystalline materials, whereas the so-called Drude model is normally used for molecular systems and fluid states.

The core-shell model is applicable to crystalline materials where the high symmetry around each site leads to stable trajectories of the core-shell pairs. However, bonded atoms in molecules can be so close that a core would interact too strongly or even capture the Drude particle of a neighbor. The Drude dipole model is relatively more complex in order to remedy this and other issues. Specifically, the Drude model includes specific thermostating of the core-Drude pairs and short-range damping of the induced dipoles.

The three polarization methods can be implemented through a self-consistent calculation of charges or induced dipoles at each timestep. In the fluctuating charge scheme this is done by the matrix inversion method in *fix qeq/point*, but for core-shell or Drude-dipoles the relaxed-dipoles technique would require an slow iterative procedure. These self-consistent solutions yield accurate trajectories since the additional degrees of freedom representing polarization are massless. An alternative is to attribute a mass to the additional degrees of freedom and perform time integration using an extended Lagrangian technique. For the fluctuating charge scheme this is done by *fix qeq/dynamic*, and for the charge-on-spring models by the methods outlined in the next two sections. The assignment of masses to the additional degrees of freedom can lead to unphysical trajectories if care is not exerted in choosing the parameters of the polarizable models and the simulation conditions.

In the core-shell model the vibration of the shells is kept faster than the ionic vibrations to mimic the fast response of the polarizable electrons. But in molecular systems thermalizing the core-Drude pairs at temperatures comparable to the rest of the simulation leads to several problems (kinetic energy transfer, too short a timestep, etc.) In order to avoid these problems the relative motion of the Drude particles with respect to their cores is kept “cold” so the vibration of the core-Drude pairs is very slow, approaching the self-consistent regime. In both models the temperature is regulated using the velocities of the center of mass of core+shell (or Drude) pairs, but in the Drude model the actual relative core-Drude particle motion is thermostatted separately as well.

### 8.5.5 Adiabatic core/shell model

The adiabatic core-shell model by *Mitchell and Fincham* is a simple method for adding polarizability to a system. In order to mimic the electron shell of an ion, a satellite particle is attached to it. This way the ions are split into a core and a shell where the latter is meant to react to the electrostatic environment inducing polarizability. See the *Howto polarizable* page for a discussion of all the polarizable models available in LAMMPS.

Technically, shells are attached to the cores by a spring force  $f = k \cdot r$  where  $k$  is a parameterized spring constant and  $r$  is the distance between the core and the shell. The charges of the core and the shell add up to the ion charge, thus  $q(\text{ion}) = q(\text{core}) + q(\text{shell})$ . This setup introduces the ion polarizability ( $\alpha$ ) given by  $\alpha = q(\text{shell})^2 / k$ . In a similar fashion the mass of the ion is distributed on the core and the shell with the core having the larger mass.

To run this model in LAMMPS, *atom\_style full* can be used since atom charge and bonds are needed. Each kind of core/shell pair requires two atom types and a bond type. The core and shell of a core/shell pair should be bonded to each other with a harmonic bond that provides the spring force. For example, a data file for NaCl, as found in *examples/coreshell*, has this format:

```
432 atoms # core and shell atoms
216 bonds # number of core/shell springs

4 atom types # 2 cores and 2 shells for Na and Cl
2 bond types

0.0 24.09597 xlo xhi
0.0 24.09597 ylo yhi
0.0 24.09597 zlo zhi

Masses # core/shell mass ratio = 0.1

1 20.690784 # Na core
2 31.90500 # Cl core
3 2.298976 # Na shell
4 3.54500 # Cl shell

Atoms
```

(continues on next page)

(continued from previous page)

```

1  1  2  1.5005  0.00000000  0.00000000  0.00000000 # core of core/shell pair 1
2  1  4 -2.5005  0.00000000  0.00000000  0.00000000 # shell of core/shell pair 1
3  2  1  1.5056  4.01599500  4.01599500  4.01599500 # core of core/shell pair 2
4  2  3 -0.5056  4.01599500  4.01599500  4.01599500 # shell of core/shell pair 2
(...)

Bonds # Bond topology for spring forces

1  2  1  2 # spring for core/shell pair 1
2  2  3  4 # spring for core/shell pair 2
(...)
```

Non-Coulombic (e.g. Lennard-Jones) pairwise interactions are only defined between the shells. Coulombic interactions are defined between all cores and shells. If desired, additional bonds can be specified between cores.

The *special\_bonds* command should be used to turn-off the Coulombic interaction within core/shell pairs, since that interaction is set by the bond spring. This is done using the *special\_bonds* command with a 1-2 weight = 0.0, which is the default value. It needs to be considered whether one has to adjust the *special\_bonds* weighting according to the molecular topology since the interactions of the shells are bypassed over an extra bond.

Note that this core/shell implementation does not require all ions to be polarized. One can mix core/shell pairs and ions without a satellite particle if desired.

Since the core/shell model permits distances of  $r = 0.0$  between the core and shell, a pair style with a “cs” suffix needs to be used to implement a valid long-range Coulombic correction. Several such pair styles are provided in the CORESHELL package. See [this page](#) for details. All of the core/shell enabled pair styles require the use of a long-range Coulombic solver, as specified by the *kspcse\_style* command. Either the PPPM or Ewald solvers can be used.

For the NaCl example problem, these pair style and bond style settings are used:

```

pair_style      born/coul/long/cs 20.0 20.0
pair_coeff      * *          0.0 1.000  0.00 0.00  0.00
pair_coeff      3 3          487.0 0.23768 0.00 1.05  0.50 #Na-Na
pair_coeff      3 4          145134.0 0.23768 0.00 6.99  8.70 #Na-Cl
pair_coeff      4 4          405774.0 0.23768 0.00 72.40 145.40 #Cl-Cl

bond_style      harmonic
bond_coeff      1 63.014 0.0
bond_coeff      2 25.724 0.0
```

When running dynamics with the adiabatic core/shell model, the following issues should be considered. The relative motion of the core and shell particles corresponds to the polarization, hereby an instantaneous relaxation of the shells is approximated and a fast core/shell spring frequency ensures a nearly constant internal kinetic energy during the simulation. Thermostats can alter this polarization behavior, by scaling the internal kinetic energy, meaning the shell will not react freely to its electrostatic environment. Therefore it is typically desirable to decouple the relative motion of the core/shell pair, which is an imaginary degree of freedom, from the real physical system. To do that, the *compute temp/cs* command can be used, in conjunction with any of the thermostat fixes, such as *fix nvt* or *fix langevin*. This compute uses the center-of-mass velocity of the core/shell pairs to calculate a temperature, and insures that velocity is what is rescaled for thermostatting purposes. This compute also works for a system with both core/shell pairs and non-polarized ions (ions without an attached satellite particle). The *compute temp/cs* command requires input of two groups, one for the core atoms, another for the shell atoms. Non-polarized ions which might also be included in the treated system should not be included into either of these groups, they are taken into account by the *group-ID* (second argument) of the compute. The groups can be defined using the *group \*type\** command. Note that to perform thermostatting using this definition of temperature, the *fix modify temp* command should be used to assign the compute to the thermostat fix. Likewise the *thermo\_modify temp* command can be used to make this temperature be output for

the overall system.

For the NaCl example, this can be done as follows:

```
group cores type 1 2
group shells type 3 4
compute CSequ all temp/cs cores shells
fix thermostoberendsen all temp/berendsen 1427 1427 0.4 # thermostat for the true_
↪physical system
fix thermostatequ all nve # integrator as needed for the_
↪berendsen thermostat
fix_modify thermostoberendsen temp CSequ
thermo_modify temp CSequ # output of center-of-mass_
↪derived temperature
```

The pressure for the core/shell system is computed via the regular LAMMPS convention by *treating the cores and shells as individual particles*. For the thermo output of the pressure as well as for the application of a barostat, it is necessary to use an additional *pressure* compute based on the default *temperature* and specifying it as a second argument in *fix modify* and *thermo\_modify* resulting in:

```
(...)
compute CSequ all temp/cs cores shells
compute thermo_press_lmp all pressure thermo_temp # pressure for individual_
↪particles
thermo_modify temp CSequ press thermo_press_lmp # modify thermo to regular_
↪pressure
fix press_bar all npt temp 300 300 0.04 iso 0 0 0.4
fix_modify press_bar temp CSequ press thermo_press_lmp # pressure modification for_
↪correct kinetic scalar
```

If *compute temp/cs* is used, the decoupled relative motion of the core and the shell should in theory be stable. However numerical fluctuation can introduce a small momentum to the system, which is noticeable over long trajectories. Therefore it is recommendable to use the *fix momentum* command in combination with *compute temp/cs* when equilibrating the system to prevent any drift.

When initializing the velocities of a system with core/shell pairs, it is also desirable to not introduce energy into the relative motion of the core/shell particles, but only assign a center-of-mass velocity to the pairs. This can be done by using the *bias* keyword of the *velocity create* command and assigning the *compute temp/cs* command to the *temp* keyword of the *velocity* command, e.g.

```
velocity all create 1427 134 bias yes temp CSequ
velocity all scale 1427 temp CSequ
```

To maintain the correct polarizability of the core/shell pairs, the kinetic energy of the internal motion shall remain nearly constant. Therefore the choice of spring force and mass ratio need to ensure much faster relative motion of the 2 atoms within the core/shell pair than their center-of-mass velocity. This allows the shells to effectively react instantaneously to the electrostatic environment and limits energy transfer to or from the core/shell oscillators. This fast movement also dictates the timestep that can be used.

The primary literature of the adiabatic core/shell model suggests that the fast relative motion of the core/shell pairs only allows negligible energy transfer to the environment. The mentioned energy transfer will typically lead to a small drift in total energy over time. This internal energy can be monitored using the *compute chunk/atom* and *compute temp/chunk* commands. The internal kinetic energies of each core/shell pair can then be summed using the *sum()* special function of the *variable* command. Or they can be time/averaged and output using the *fix ave/time* command. To use these commands, each core/shell pair must be defined as a “chunk”. If each core/shell pair is defined as its own molecule, the molecule ID can be used to define the chunks. If cores are bonded to each other to form larger molecules,

the chunks can be identified by the *fix property/atom* via assigning a core/shell ID to each atom using a special field in the data file read by the *read\_data* command. This field can then be accessed by the *compute property/atom* command, to use as input to the *compute chunk/atom* command to define the core/shell pairs as chunks.

For example if core/shell pairs are the only molecules:

```
read_data NaCl_CS_x0.1_prop.data
compute prop all property/atom molecule
compute cs_chunk all chunk/atom c_prop
compute csthern all temp/chunk cs_chunk temp internal com yes cdof 3.0 # note the
↪chosen degrees of freedom for the core/shell pairs
fix ave_chunk all ave/time 10 1 10 c_csthern file chunk.dump mode vector
```

For example if core/shell pairs and other molecules are present:

```
fix csinfo all property/atom i_CSID # property/atom command
read_data NaCl_CS_x0.1_prop.data fix csinfo NULL CS-Info # atom property added in the
↪data-file
compute prop all property/atom i_CSID
(...)
```

The additional section in the data file would be formatted like this:

```
CS-Info # header of additional section

1 1 # column 1 = atom ID, column 2 = core/shell ID
2 1
3 2
4 2
5 3
6 3
7 4
8 4
(...)
```

(**Mitchell and Fincham**) Mitchell, Fincham, J Phys Condensed Matter, 5, 1031-1038 (1993).

(**Fincham**) Fincham, Mackrodt and Mitchell, J Phys Condensed Matter, 6, 393-404 (1994).

### 8.5.6 Drude induced dipoles

The thermalized Drude model represents induced dipoles by a pair of charges (the core atom and the Drude particle) connected by a harmonic spring. See the *Howto polarizable* doc page for a discussion of all the polarizable models available in LAMMPS.

The Drude model has a number of features aimed at its use in molecular systems (*Lamoureux and Roux*):

- Thermostatting of the additional degrees of freedom associated with the induced dipoles at very low temperature, in terms of the reduced coordinates of the Drude particles with respect to their cores. This makes the trajectory close to that of relaxed induced dipoles.
- Consistent definition of 1-2 to 1-4 neighbors. A core-Drude particle pair represents a single (polarizable) atom, so the special screening factors in a covalent structure should be the same for the core and the Drude particle. Drude particles have to inherit the 1-2, 1-3, 1-4 special neighbor relations from their respective cores.

- Stabilization of the interactions between induced dipoles. Drude dipoles on covalently bonded atoms interact too strongly due to the short distances, so an atom may capture the Drude particle of a neighbor, or the induced dipoles within the same molecule may align too much. To avoid this, damping at short range can be done by Thole functions (for which there are physical grounds). This Thole damping is applied to the point charges composing the induced dipole (the charge of the Drude particle and the opposite charge on the core, not to the total charge of the core atom).

A detailed tutorial covering the usage of Drude induced dipoles in LAMMPS is on the [here](#).

As with the core-shell model, the cores and Drude particles should appear in the data file as standard atoms. The same holds for the springs between them, which are described by standard harmonic bonds. The nature of the atoms (core, Drude particle or non-polarizable) is specified via the `fix drude` command. The special list of neighbors is automatically refactored to account for the equivalence of core and Drude particles as regards special 1-2 to 1-4 screening. It may be necessary to use the `extra/special/per/atom` keyword of the `read_data` command. If using `fix shake`, make sure no Drude particle is in this fix group.

There are three ways to thermostat the Drude particles at a low temperature: use either `fix langevin/drude` for a Langevin thermostat, or `fix drude/transform/*` for a Nose-Hoover thermostat, or `fix tgnvt/drude` for a temperature-grouped Nose-Hoover thermostat. The first and third require use of the command `comm_modify vel yes`. The second requires two separate integration fixes like `nvt` or `npt`. The correct temperatures of the reduced degrees of freedom can be calculated using the `compute temp/drude`. This requires also to use the command `comm_modify vel yes`.

Short-range damping of the induced dipole interactions can be achieved using Thole functions through the `pair style thole` in `pair_style hybrid/overlay` with a Coulomb pair style. It may be useful to use `coul/long/cs` or similar from the CORESHELL package if the core and Drude particle come too close, which can cause numerical issues.

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(Lamoureux and Roux) G. Lamoureux, B. Roux, J. Chem. Phys 119, 3025 (2003)

## 8.5.7 Tutorial for Thermalized Drude oscillators in LAMMPS

This tutorial explains how to use Drude oscillators in LAMMPS to simulate polarizable systems using the DRUDE package. As an illustration, the input files for a simulation of 250 phenol molecules are documented. First of all, LAMMPS has to be compiled with the DRUDE package activated. Then, the data file and input scripts have to be modified to include the Drude dipoles and how to handle them.

Example input scripts available: `examples/PACKAGES/drude`

---

### Overview of Drude induced dipoles

Polarizable atoms acquire an induced electric dipole moment under the action of an external electric field, for example the electric field created by the surrounding particles. Drude oscillators represent these dipoles by two fixed charges: the core (DC) and the Drude particle (DP) bound by a harmonic potential. The Drude particle can be thought of as the electron cloud whose center can be displaced from the position of the corresponding nucleus.

The sum of the masses of a core-Drude pair should be the mass of the initial (unsplit) atom,  $m_C + m_D = m$ . The sum of their charges should be the charge of the initial (unsplit) atom,  $q_C + q_D = q$ . A harmonic potential between the core and Drude partners should be present, with force constant  $k_D$  and an equilibrium distance of zero. The (half-)stiffness of the *harmonic bond*  $K_D = k_D/2$  and the Drude charge  $q_D$  are related to the atom polarizability  $\alpha$  by

$$K_D = \frac{1}{2} \frac{q_D^2}{\alpha}$$

Ideally, the mass of the Drude particle should be small, and the stiffness of the harmonic bond should be large, so that the Drude particle remains close to the core. The values of Drude mass, Drude charge, and force constant can be chosen following different strategies, as in the following examples of polarizable force fields:

- *Lamoureux and Roux* suggest adopting a global half-stiffness,  $K_D = 500$  kcal/(mol Ang<sup>2</sup>) - which corresponds to a force constant  $k_D = 4184$  kJ/(mol Ang<sup>2</sup>) - for all types of core-Drude bond, a global mass  $m_D = 0.4$  g/mol (or u) for all types of Drude particles, and to calculate the Drude charges for individual atom types from the atom polarizabilities using equation (1). This choice is followed in the polarizable CHARMM force field.
- Alternately *Schroeder and Steinhauser* suggest adopting a global charge  $q_D = -1.0e$  and a global mass  $m_D = 0.1$  g/mol (or u) for all Drude particles, and to calculate the force constant for each type of core-Drude bond from equation (1). The timesteps used by these authors are between 0.5 and 2 fs, with the degrees of freedom of the Drude oscillators kept cold at 1 K.
- In both these force fields hydrogen atoms are treated as non-polarizable.

The motion of the Drude particles can be calculated by minimizing the energy of the induced dipoles at each timestep, by an iterative, self-consistent procedure. The Drude particles can be massless and therefore do not contribute to the kinetic energy. However, the relaxed method is computational slow. An extended-lagrangian method can be used to calculate the positions of the Drude particles, but this requires them to have mass. It is important in this case to decouple the degrees of freedom associated with the Drude oscillators from those of the normal atoms. Thermalizing the Drude dipoles at temperatures comparable to the rest of the simulation leads to several problems (kinetic energy transfer, very short timestep, etc.), which can be remedied by the “cold Drude” technique (*Lamoureux and Roux*).

Two closely related models are used to represent polarization through “charges on a spring”: the core-shell model and the Drude model. Although the basic idea is the same, the core-shell model is normally used for ionic/crystalline materials, whereas the Drude model is normally used for molecular systems and fluid states. In ionic crystals the symmetry around each ion and the distance between them are such that the core-shell model is sufficiently stable. But to be applicable to molecular/covalent systems the Drude model includes two important features:

1. The possibility to thermostat the additional degrees of freedom associated with the induced dipoles at very low temperature, in terms of the reduced coordinates of the Drude particles with respect to their cores. This makes the trajectory close to that of relaxed induced dipoles.
2. The Drude dipoles on covalently bonded atoms interact too strongly due to the short distances, so an atom may capture the Drude particle (shell) of a neighbor, or the induced dipoles within the same molecule may align too much. To avoid this, damping at short of the interactions between the point charges composing the induced dipole can be done by *Thole* functions.

### Preparation of the data file

The data file is similar to a standard LAMMPS data file for *atom\_style full*. The DPs and the *harmonic bonds* connecting them to their DC should appear in the data file as normal atoms and bonds.

You can use the *polarizer* tool (Python script distributed with the DRUDE package) to convert a non-polarizable data file (here *data.102494.lmp*) to a polarizable data file (*data-p.lmp*)

```
polarizer -q -f phenol.dff data.102494.lmp data-p.lmp
```

This will automatically insert the new atoms and bonds. The masses and charges of DCs and DPs are computed from *phenol.dff*, as well as the DC-DP bond constants. The file *phenol.dff* contains the polarizabilities of the atom types and the mass of the Drude particles, for instance:

```
# units: kJ/mol, A, deg
# kforce is in the form k/2 r_D^2
# type m_D/u q_D/e k_D alpha/A3 thole
OH 0.4 -1.0 4184.0 0.63 0.67
CA 0.4 -1.0 4184.0 1.36 2.51
CAI 0.4 -1.0 4184.0 1.09 2.51
```

The hydrogen atoms are absent from this file, so they will be treated as non-polarizable atoms. In the non-polarizable data file *data.102494.lmp*, atom names corresponding to the atom type numbers have to be specified as comments at the end of lines of the *Masses* section. You probably need to edit it to add these names. It should look like



## Masses

```
1 12.011 # CAI
2 12.011 # CA
3 15.999 # OH
4 1.008 # HA
5 1.008 # HO
```

## Basic input file

The atom style should be set to (or derive from) *full*, so that you can define atomic charges and molecular bonds, angles, dihedrals...

The *polarizer* tool also outputs certain lines related to the input script (the use of these lines will be explained below). In order for LAMMPS to recognize that you are using Drude oscillators, you should use the fix *drude*. The command is

```
fix DRUDE all drude C C C N N D D D
```

The N, C, D following the *drude* keyword have the following meaning: There is one tag for each atom type. This tag is C for DCs, D for DPs and N for non-polarizable atoms. Here the atom types 1 to 3 (C and O atoms) are DC, atom types 4 and 5 (H atoms) are non-polarizable and the atom types 6 to 8 are the newly created DPs.

By recognizing the fix *drude*, LAMMPS will find and store matching DC-DP pairs and will treat DP as equivalent to their DC in the *special bonds* relations. It may be necessary to extend the space for storing such special relations. In this case extra space should be reserved by using the *extra/special/per/atom* keyword of either the *read\_data* or *create\_box* command. With our phenol, there is 1 more special neighbor for which space is required. Otherwise LAMMPS crashes and gives the required value.

```
read_data data-p.lmp extra/special/per/atom 1
```

Let us assume we want to run a simple NVT simulation at 300 K. Note that Drude oscillators need to be thermalized at a low temperature in order to approximate a self-consistent field (SCF), therefore it is not possible to simulate an NVE ensemble with this package. Since dipoles are approximated by a charged DC-DP pair, the *pair\_style* must include Coulomb interactions, for instance *lj/cut/coul/long* with *k-space\_style ppm*. For example, with a cutoff of 10. and a precision 1.e-4:

```
pair_style lj/cut/coul/long 10.0
k-space_style ppm 1.0e-4
```

As compared to the non-polarizable input file, *pair\_coeff* lines need to be added for the DPs. Since the DPs have no Lennard-Jones interactions, their  $\epsilon$  is 0. so the only *pair\_coeff* line that needs to be added is

```
pair_coeff * 6* 0.0 0.0 # All-DPs
```

Now for the thermalization, the simplest choice is to use the fix *langevin/drude*.

```
fix LANG all langevin/drude 300. 100 12345 1. 20 13977
```

This applies a Langevin thermostat at temperature 300. to the centers of mass of the DC-DP pairs, with relaxation time 100 and with random seed 12345. This fix applies also a Langevin thermostat at temperature 1. to the relative motion of the DPs around their DCs, with relaxation time 20 and random seed 13977. Only the DCs and non-polarizable atoms need to be in this fix's group. LAMMPS will thermostat the DPs together with their DC. For this, ghost atoms need to know their velocities. Thus you need to add the following command:

```
comm_modify vel yes
```

In order to avoid that the center of mass of the whole system drifts due to the random forces of the Langevin thermostat on DCs, you can add the *zero yes* option at the end of the fix line.

If the fix *shake* is used to constrain the C-H bonds, it should be invoked after the fix *langevin/drude* for more accuracy.

```
fix SHAKE ATOMS shake 0.0001 20 0 t 4 5
```

---

**Note:** The group of the fix *shake* must not include the DPs. If the group *ATOMS* is defined by non-DPs atom types, you could use

---

Since the fix *langevin/drude* does not perform time integration (just modification of forces but no position/velocity updates), the fix *nve* should be used in conjunction.

```
fix NVE all nve
```

To avoid the flying ice cube artifact, where the atoms progressively freeze and the center of mass of the whole system drifts faster and faster, the fix *momentum* can be used. For instance:

```
fix MOMENTUM all momentum 100 linear 1 1 1
```

Finally, do not forget to update the atom type elements if you use them in a *dump\_modify ... element ...* command, by adding the element type of the DPs. Here for instance

```
dump DUMP all custom 10 dump.lammpstrj id mol type element x y z ix iy iz
dump_modify DUMP element C C O H H D D D
```

The input file should now be ready for use!

You will notice that the global temperature *thermo\_temp* computed by LAMMPS is not 300. K as wanted. This is because LAMMPS treats DPs as standard atoms in his default compute. If you want to output the temperatures of the DC-DP pair centers of mass and of the DPs relative to their DCs, you should use the *compute temp\_drude*

```
compute TDRUDE all temp/drude
```

And then output the correct temperatures of the Drude oscillators using *thermo\_style custom* with respectively *c\_TDRUDE[1]* and *c\_TDRUDE[2]*. These should be close to 300.0 and 1.0 on average.

```
thermo_style custom step temp c_TDRUDE[1] c_TDRUDE[2]
```

---

### Thole screening

Dipolar interactions represented by point charges on springs may not be stable, for example if the atomic polarizability is too high for instance, a DP can escape from its DC and be captured by another DC, which makes the force and energy diverge and the simulation crash. Even without reaching this extreme case, the correlation between nearby dipoles on the same molecule may be exaggerated. Often, special bond relations prevent bonded neighboring atoms to see the charge of each other's DP, so that the problem does not always appear. It is possible to use screened dipole-dipole interactions by using the *\*pair\_style thole\**. This is implemented as a correction to the Coulomb pair\_styles, which dampens at short distance the interactions between the charges representing the induced dipoles. It is to be used as *hybrid/overlay* with any standard *coul* pair style. In our example, we would use

```
pair_style hybrid/overlay lj/cut/coul/long 10.0 thole 2.6 10.0
```

This tells LAMMPS that we are using two `pair_styles`. The first one is as above (`lj/cut/coul/long 10.0`). The second one is a `thole` `pair_style` with default screening factor 2.6 (*Noskov*) and cutoff 10.0.

Since `hybrid/overlay` does not support mixing rules, the interaction coefficients of all the pairs of atom types with  $i < j$  should be explicitly defined. The output of the `polarizer` script can be used to complete the `pair_coeff` section of the input file. In our example, this will look like:

```
pair_coeff 1 1 lj/cut/coul/long 0.0700 3.550
pair_coeff 1 2 lj/cut/coul/long 0.0700 3.550
pair_coeff 1 3 lj/cut/coul/long 0.1091 3.310
pair_coeff 1 4 lj/cut/coul/long 0.0458 2.985
pair_coeff 2 2 lj/cut/coul/long 0.0700 3.550
pair_coeff 2 3 lj/cut/coul/long 0.1091 3.310
pair_coeff 2 4 lj/cut/coul/long 0.0458 2.985
pair_coeff 3 3 lj/cut/coul/long 0.1700 3.070
pair_coeff 3 4 lj/cut/coul/long 0.0714 2.745
pair_coeff 4 4 lj/cut/coul/long 0.0300 2.420
pair_coeff * 5 lj/cut/coul/long 0.0000 0.000
pair_coeff * 6* lj/cut/coul/long 0.0000 0.000
pair_coeff 1 1 thole 1.090 2.510
pair_coeff 1 2 thole 1.218 2.510
pair_coeff 1 3 thole 0.829 1.590
pair_coeff 1 6 thole 1.090 2.510
pair_coeff 1 7 thole 1.218 2.510
pair_coeff 1 8 thole 0.829 1.590
pair_coeff 2 2 thole 1.360 2.510
pair_coeff 2 3 thole 0.926 1.590
pair_coeff 2 6 thole 1.218 2.510
pair_coeff 2 7 thole 1.360 2.510
pair_coeff 2 8 thole 0.926 1.590
pair_coeff 3 3 thole 0.630 0.670
pair_coeff 3 6 thole 0.829 1.590
pair_coeff 3 7 thole 0.926 1.590
pair_coeff 3 8 thole 0.630 0.670
pair_coeff 6 6 thole 1.090 2.510
pair_coeff 6 7 thole 1.218 2.510
pair_coeff 6 8 thole 0.829 1.590
pair_coeff 7 7 thole 1.360 2.510
pair_coeff 7 8 thole 0.926 1.590
pair_coeff 8 8 thole 0.630 0.670
```

For the `thole` pair style the coefficients are

1. the atom polarizability in units of cubic length
2. the screening factor of the Thole function (optional, default value specified by the `pair_style` command)
3. the cutoff (optional, default value defined by the `pair_style` command)

The special neighbors have charge-charge and charge-dipole interactions screened by the `coul` factors of the `special_bonds` command (0.0, 0.0, and 0.5 in the example above). Without using the `pair_style thole`, dipole-dipole interactions are screened by the same factor. By using the `pair_style thole`, dipole-dipole interactions are screened by Thole's function, whatever their special relationship (except within each DC-DP pair of course). Consider for example 1-2 neighbors: using the `pair_style thole`, their dipoles will see each other (despite the `coul` factor being 0.) and the

interactions between these dipoles will be damped by Thole's function.

---

### Thermostats and barostats

Using a Nose-Hoover barostat with the *langevin/drude* thermostat is straightforward using *fix nph* instead of *nve*. For example:

```
fix NPH all nph iso 1. 1. 500
```

It is also possible to use a Nose-Hoover instead of a Langevin thermostat. This requires to use *\*fix drude/transform\** just before and after the time integration fixes. The *fix drude/transform/direct* converts the atomic masses, positions, velocities and forces into a reduced representation, where the DCs transform into the centers of mass of the DC-DP pairs and the DPs transform into their relative position with respect to their DC. The *fix drude/transform/inverse* performs the reverse transformation. For a NVT simulation, with the DCs and atoms at 300 K and the DPs at 1 K relative to their DC one would use

```
fix DIRECT all drude/transform/direct
fix NVT1 ATOMS nvt temp 300. 300. 100
fix NVT2 DRUDES nvt temp 1. 1. 20
fix INVERSE all drude/transform/inverse
```

For our phenol example, the groups would be defined as

```
group ATOMS type 1 2 3 4 5 # DCs and non-polarizable atoms
group CORES type 1 2 3 # DCs
group DRUDES type 6 7 8 # DPs
```

Note that with the fixes *drude/transform*, it is not required to specify *comm\_modify vel yes* because the fixes do it anyway (several times and for the forces also).

It is a bit more tricky to run a NPT simulation with Nose-Hoover barostat and thermostat. First, the volume should be integrated only once. So the fix for DCs and atoms should be *npt* while the fix for DPs should be *nvt* (or vice versa). Second, the *fix npt* computes a global pressure and thus a global temperature whatever the fix group. We do want the pressure to correspond to the whole system, but we want the temperature to correspond to the fix group only. We must then use the *fix\_modify* command for this. In the end, the block of instructions for thermostating and barostatting will look like

```
compute TATOMS ATOMS temp
fix DIRECT all drude/transform/direct
fix NPT ATOMS npt temp 300. 300. 100 iso 1. 1. 500
fix_modify NPT temp TATOMS press thermo_press
fix NVT DRUDES nvt temp 1. 1. 20
fix INVERSE all drude/transform/inverse
```

Another option for thermalizing the Drude model is to use the temperature-grouped Nose-Hoover (TGNH) thermostat proposed by (Son). This is implemented as *fix tgnvt/drude* and *fix tgnpt/drude*. It separates the kinetic energy into three contributions: the molecular center of mass (COM) motion, the motion of atoms or atom-Drude pairs relative to molecular COMs, and the relative motion of atom-Drude pairs. An independent Nose-Hoover chain is applied to each type of motion. When TGNH is used, the temperatures of molecular, atomic and Drude motion can be printed out with *thermo\_style command* command.

NVT simulation with TGNH thermostat

```
comm_modify vel yes
fix TGNVT all tgnvt/drude temp 300. 300. 100 1. 20
thermo_style custom f_TGNVT[1] f_TGNVT[2] f_TGNVT[3]
```

NPT simulation with TGNH thermostat

```
comm_modify vel yes
fix TGNPT all tgnpt/drude temp 300. 300. 100 1. 20 iso 1. 1. 500
thermo_style custom f_TGNPT[1] f_TGNPT[2] f_TGNPT[3]
```

## Rigid bodies

You may want to simulate molecules as rigid bodies (but polarizable). Common cases are water models such as *SWM4-NDP*, which is a kind of polarizable TIP4P water. The rigid bodies and the DPs should be integrated separately, even with the Langevin thermostat. Let us review the different thermostats and ensemble combinations.

NVT ensemble using Langevin thermostat:

```
comm_modify vel yes
fix LANG all langevin/drude 300. 100 12435 1. 20 13977
fix RIGID ATOMS rigid/nve/small molecule
fix NVE DRUDES nve
```

NVT ensemble using Nose-Hoover thermostat:

```
fix DIRECT all drude/transform/direct
fix RIGID ATOMS rigid/nvt/small molecule temp 300. 300. 100
fix NVT DRUDES nvt temp 1. 1. 20
fix INVERSE all drude/transform/inverse
```

NPT ensemble with Langevin thermostat:

```
comm_modify vel yes
fix LANG all langevin/drude 300. 100 12435 1. 20 13977
fix RIGID ATOMS rigid/nph/small molecule iso 1. 1. 500
fix NVE DRUDES nve
```

NPT ensemble using Nose-Hoover thermostat:

```
compute TATOM ATOMS temp
fix DIRECT all drude/transform/direct
fix RIGID ATOMS rigid/npt/small molecule temp 300. 300. 100 iso 1. 1. 500
fix_modify RIGID temp TATOM press thermo_press
fix NVT DRUDES nvt temp 1. 1. 20
fix INVERSE all drude/transform/inverse
```

(**Lamoureux and Roux**) Lamoureux and Roux, J Chem Phys, 119, 3025-3039 (2003)

(**Schroeder**) Schroeder and Steinhauser, J Chem Phys, 133, 154511 (2010).

(**Thole**) Chem Phys, 59, 341 (1981).

(**Noskov**) Noskov, Lamoureux and Roux, J Phys Chem B, 109, 6705 (2005).

(**SWM4-NDP**) Lamoureux, Harder, Vorobyov, Roux, MacKerell, Chem Phys Let, 418, 245-249 (2006)

(Son) Son, McDaniel, Cui and Yethiraj, J Phys Chem Lett, 10, 7523 (2019).

## 8.5.8 Manifolds (surfaces)

### Overview:

This page is not about a LAMMPS input script command, but about manifolds, which are generalized surfaces, as defined and used by the MANIFOLD package, to track particle motion on the manifolds. See the src/MANIFOLD/README file for more details about the package and its commands.

Below is a list of currently supported manifolds by the MANIFOLD package, their parameters and a short description of them. The parameters listed here are in the same order as they should be passed to the relevant fixes.

<i>manifold</i>	<i>parameters</i>	<i>equation</i>	<i>description</i>
cylinder	R	$x^2 + y^2 - R^2 = 0$	Cylinder along z-axis, axis going through (0,0,0)
cylinder_dent	R l a	$x^2 + y^2 - r(z)^2 = 0$ , $r(x) = R$ if $ z  > l$ , $r(z) = R - a*(1 + \cos(z/l))/2$ otherwise	A cylinder with a dent around $z = 0$
dumbbell	a A B c	$-(x^2 + y^2) + (a^2 - z^2/c^2) * (1 + (A*\sin(B*z^2))^4) = 0$	A dumbbell
ellipsoid	a b c	$(x/a)^2 + (y/b)^2 + (z/c)^2 = 0$	An ellipsoid
gaussian_bump	A l rc1 rc2	if $(x < rc1) -z + A * \exp(-x^2 / (2 l^2))$ ; else if $(x < rc2) -z + a + b*x + c*x^2 + d*x^3$ ; else $z$	A Gaussian bump at $x = y = 0$ , smoothly tapered to a flat plane $z = 0$ .
plane	a b c x0 y0 z0	$a*(x-x0) + b*(y-y0) + c*(z-z0) = 0$	A plane with normal (a,b,c) going through point (x0,y0,z0)
plane_wiggle	w	$z - a*\sin(w*x) = 0$	A plane with a sinusoidal modulation on z along x.
sphere	R	$x^2 + y^2 + z^2 - R^2 = 0$	A sphere of radius R
supersphere	R q	$ x ^q +  y ^q +  z ^q - R^q = 0$	A supersphere of hyperradius R
spine	a, A, B, B2, c	$-(x^2 + y^2) + (a^2 - z^2/f(z)^2)*(1 + (A*\sin(g(z)*z^2))^4)$ , $f(z) = c$ if $z > 0$ , 1 otherwise; $g(z) = B$ if $z > 0$ , B2 otherwise	An approximation to a dendritic spine
spine_two	a, A, B, B2, c	$-(x^2 + y^2) + (a^2 - z^2/f(z)^2)*(1 + (A*\sin(g(z)*z^2))^2)$ , $f(z) = c$ if $z > 0$ , 1 otherwise; $g(z) = B$ if $z > 0$ , B2 otherwise	Another approximation to a dendritic spine
thylakoid	wB LB IB	Various, see ( <i>Paquay</i> )	A model grana thylakoid consisting of two block-like compartments connected by a bridge of width wB, length LB and taper length IB
torus	R r	$(R - \sqrt{x^2 + y^2})^2 + z^2 - r^2 = 0$	A torus with large radius R and small radius r, centered on (0,0,0)

(Paquay) Paquay and Kusters, Biophys. J., 110, 6, (2016). preprint available at [arXiv:1411.3019](https://arxiv.org/abs/1411.3019).

## 8.5.9 Magnetic spins

The magnetic spin simulations are enabled by the SPIN package, whose implementation is detailed in *Tranchida*.

The model represents the simulation of atomic magnetic spins coupled to lattice vibrations. The dynamics of those magnetic spins can be used to simulate a broad range a phenomena related to magneto-elasticity, or or to study the influence of defects on the magnetic properties of materials.

The magnetic spins are interacting with each others and with the lattice via pair interactions. Typically, the magnetic exchange interaction can be defined using the *pair/spin/exchange* command. This exchange applies a magnetic torque to a given spin, considering the orientation of its neighboring spins and their relative distances. It also applies a force on the atoms as a function of the spin orientations and their associated inter-atomic distances.

The command *fix precession/spin* allows to apply a constant magnetic torque on all the spins in the system. This torque can be an external magnetic field (Zeeman interaction), and an uniaxial or cubic magnetic anisotropy.

A Langevin thermostat can be applied to those magnetic spins using *fix langevin/spin*. Typically, this thermostat can be coupled to another Langevin thermostat applied to the atoms using *fix langevin* in order to simulate thermostatted spin-lattice systems.

The magnetic Gilbert damping can also be applied using *fix langevin/spin*. It allows to either dissipate the thermal energy of the Langevin thermostat, or to perform a relaxation of the magnetic configuration toward an equilibrium state.

The command *fix setforce/spin* allows to set the components of the magnetic precession vectors (while erasing and replacing the previously computed magnetic precession vectors on the atom). This command can be used to freeze the magnetic moment of certain atoms in the simulation by zeroing their precession vector.

The command *fix nve/spin* can be used to perform a symplectic integration of the combined dynamics of spins and atomic motions.

The minimization style *min/spin* can be applied to the spins to perform a minimization of the spin configuration.

All the computed magnetic properties can be output by two main commands. The first one is *compute spin*, that enables to evaluate magnetic averaged quantities, such as the total magnetization of the system along x, y, or z, the spin temperature, or the magnetic energy. The second command is *compute property/atom*. It enables to output all the per atom magnetic quantities. Typically, the orientation of a given magnetic spin, or the magnetic force acting on this spin.

---

(**Tranchida**) Tranchida, Plimpton, Thibaudau and Thompson, Journal of Computational Physics, 372, 406-425, (2018).

## 8.6 Tutorials howto

### 8.6.1 Using CMake with LAMMPS tutorial

The support for building LAMMPS with CMake is a recent addition to LAMMPS thanks to the efforts of Christoph Junghans (LANL) and Richard Berger (Temple U). One of the key strengths of CMake is that it is not tied to a specific platform or build system and thus generate the files necessary to build and develop for different build systems and on different platforms. Note, that this applies to the build system itself not the LAMMPS code. In other words, without additional porting effort, it is not possible - for example - to compile LAMMPS with Visual C++ on Windows. The build system output can also include support files necessary to program LAMMPS as a project in integrated development environments (IDE) like Eclipse, Visual Studio, QtCreator, Xcode, CodeBlocks, Kate and others.

A second important feature of CMake is, that it can detect and validate available libraries, optimal settings, available support tools and so on, so that by default LAMMPS will take advantage of available tools without requiring to provide the details about how to enable/integrate them.

The downside of this approach is, that there is some complexity associated with running CMake itself and how to customize the building of LAMMPS. This tutorial will show how to manage this through some selected examples. Please see the chapter about *building LAMMPS* for descriptions of specific flags and options for LAMMPS in general and for specific packages.

CMake can be used through either the command-line interface (CLI) program `cmake` (or `cmake3`), a text mode interactive user interface (TUI) program `ccmake` (or `ccmake3`), or a graphical user interface (GUI) program `cmake-gui`. All of them are portable software available on all supported platforms and can be used interchangeably. The minimum supported CMake version is 3.10 (3.12 or later is recommended).

All details about features and settings for CMake are in the [CMake online documentation](#). We focus below on the most important aspects with respect to compiling LAMMPS.

### Prerequisites

This tutorial assumes that you are operating in a command-line environment using a shell like Bash.

- Linux: any Terminal window will work
- MacOS X: launch the Terminal application.
- Windows 10: install and run the *Windows Subsystem for Linux*

We also assume that you have downloaded and unpacked a recent LAMMPS source code package or used Git to create a clone of the LAMMPS sources on your compilation machine.

You should change into the top level directory of the LAMMPS source tree all paths mentioned in the tutorial are relative to that. Immediately after downloading it should look like this:

```
$ ls
bench  doc      lib      potentials  README  tools
cmake  examples LICENSE  python      src
```

### Build versus source directory

When using CMake the build procedure is separated into multiple distinct phases:

1. **Configuration:** detect or define which features and settings should be enable and used and how LAMMPS should be compiled
2. **Compilation:** generate and compile all necessary source files and build libraries and executables.
3. **Installation:** copy selected files from the compilation into your file system, so they can be used without having to keep the source and build tree around.

The configuration and compilation of LAMMPS has to happen in a dedicated *build directory* which must be different from the source directory. Also the source directory (`src`) must remain pristine, so it is not allowed to “install” packages using the traditional make process and after an compilation attempt all created source files must be removed. This can be achieved with `make no-all purge`.

You can pick **any** folder outside the source tree. We recommend to create a folder `build` in the top-level directory, or multiple folders in case you want to have separate builds of LAMMPS with different options (`build-parallel`, `build-serial`) or with different compilers (`build-gnu`, `build-clang`, `build-intel`) and so on. All the auxiliary files created by one build process (executable, object files, log files, etc) are stored in this directory or sub-directories within it that CMake creates.



## Running CMake

### CLI version

In the (empty) build directory, we now run the command `cmake ../cmake`, which will start the configuration phase and you will see the progress of the configuration printed to the screen followed by a summary of the enabled features, options and compiler settings. A typical summary screen will look like this:

```
$ cmake ../cmake/
-- The CXX compiler identification is GNU 8.2.0
-- Check for working CXX compiler: /opt/tools/gcc-8.2.0/bin/c++
-- Check for working CXX compiler: /opt/tools/gcc-8.2.0/bin/c++ - works
-- Detecting CXX compiler ABI info
-- Detecting CXX compiler ABI info - done
-- Detecting CXX compile features
-- Detecting CXX compile features - done
-- Found Git: /usr/bin/git (found version "2.25.2")
-- Running check for auto-generated files from make-based build system
-- Found MPI_CXX: /usr/lib64/mpich/lib/libmpicxx.so (found version "3.1")
-- Found MPI: TRUE (found version "3.1")
-- Looking for C++ include omp.h
-- Looking for C++ include omp.h - found
-- Found OpenMP_CXX: -fopenmp (found version "4.5")
-- Found OpenMP: TRUE (found version "4.5")
-- Found JPEG: /usr/lib64/libjpeg.so (found version "62")
-- Found PNG: /usr/lib64/libpng.so (found version "1.6.37")
-- Found ZLIB: /usr/lib64/libz.so (found version "1.2.11")
-- Found GZIP: /usr/bin/gzip
-- Found FFMPEG: /usr/bin/ffmpeg
-- Performing Test COMPILER_SUPPORTS_ffast-math
-- Performing Test COMPILER_SUPPORTS_ffast-math - Success
-- Performing Test COMPILER_SUPPORTS_march=native
-- Performing Test COMPILER_SUPPORTS_march=native - Success
-- Looking for C++ include cmath
-- Looking for C++ include cmath - found
-- Generating style_angle.h...
[...]
-- Generating lmpinstalledpkgs.h...
-- The following tools and libraries have been found and configured:
* Git
* MPI
* OpenMP
* JPEG
* PNG
* ZLIB

-- <<< Build configuration >>>
  Build type:      RelWithDebInfo
  Install path:   /home/akohlmey/.local
  Generator:      Unix Makefiles using /usr/bin/gmake
-- <<< Compilers and Flags: >>>
-- C++ Compiler:  /opt/tools/gcc-8.2.0/bin/c++
  Type:          GNU
```

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```

Version:      8.2.0
C++ Flags:   -O2 -g -DNDEBUG
Defines:     LAMMPS_SMALLBIG;LAMMPS_MEMALIGN=64;LAMMPS_JPEG;LAMMPS_PNG;LAMMPS_
↪GZIP;LAMMPS_FFMPEG
Options:     -ffast-math;-march=native
-- <<< Linker flags: >>>
-- Executable name: lmp
-- Static library flags:
-- <<< MPI flags >>>
-- MPI includes: /usr/include/mpich-x86_64
-- MPI libraries: /usr/lib64/mpich/lib/libmpicxx.so;/usr/lib64/mpich/lib/libmpi.so;
-- Configuring done
-- Generating done
-- Build files have been written to: /home/akohlmeier/compile/lammps/build

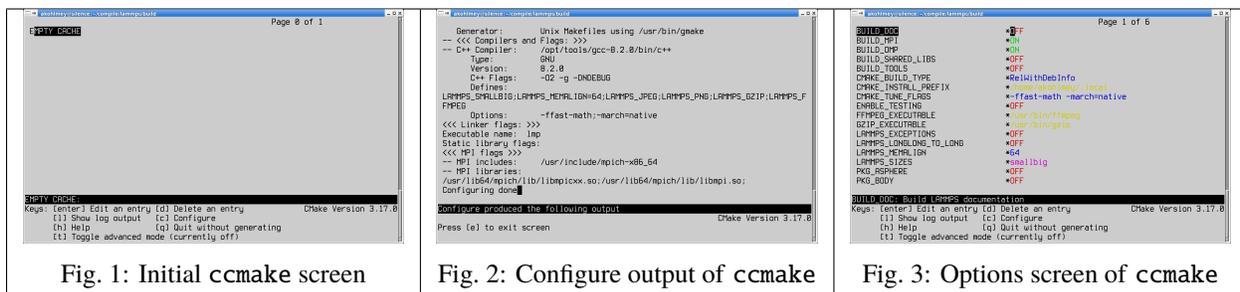
```

The `cmake` command has one mandatory argument, and that is a folder with either the file `CMakeLists.txt` or `CMakeCache.txt`. The `CMakeCache.txt` file is created during the CMake configuration run and contains all active settings, thus after a first run of CMake all future runs in the build folder can use the folder `.` and CMake will know where to find the CMake scripts and reload the settings from the previous step. This means, that one can modify an existing configuration by re-running CMake, but only needs to provide flags indicating the desired change, everything else will be retained. One can also mix compilation and configuration, i.e. start with a minimal configuration and then, if needed, enable additional features and recompile.

The steps above **will NOT compile the code**. The compilation can be started in a portable fashion with `cmake --build .`, or you use the selected built tool, e.g. `make`.

## TUI version

For the text mode UI CMake program the basic principle is the same. You start the command `ccmake ../cmake` in the build folder.



This will show you the initial screen (left image) with the empty configuration cache. Now you type the ‘c’ key to run the configuration step. That will do a first configuration run and show the summary (center image). You exit the summary screen with ‘e’ and see now the main screen with detected options and settings. You can now make changes by moving and down with the arrow-g of the keyboard and modify entries. For on/off settings, the enter key will toggle the state. For others, hitting enter will allow you to modify the value and you commit the change by hitting the enter key again or cancel using the escape key. All “new” settings will be marked with a star ‘\*’ and for as long as one setting is marked like this, you have to re-run the configuration by hitting the ‘c’ key again, sometimes multiple times unless the TUI shows the word “generate” next to the letter ‘g’ and by hitting the ‘g’ key the build files will be written to the folder and the TUI exits. You can quit without generating build files by hitting ‘q’.

## GUI version

For the graphical CMake program the steps are similar to the TUI version. You can type the command `cmake-gui . ./cmake` in the `build` folder. In this case the path to the CMake script folder is not required, it can also be entered from the GUI.

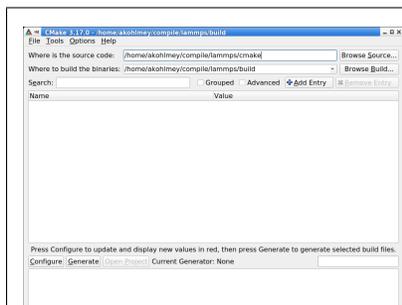


Fig. 4: Initial `cmake-gui` screen

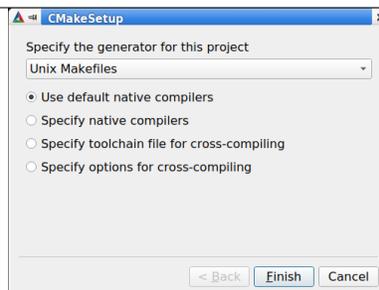


Fig. 5: Generator selection in `cmake-gui`

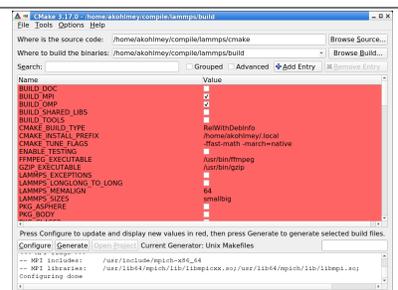


Fig. 6: Options screen of `cmake-gui`

Again, you start with an empty configuration cache (left image) and need to start the configuration step. For the very first configuration in a folder, you will have a pop-up dialog (center image) asking to select the desired build tool and some configuration settings (stick with the default) and then you get the option screen with all new settings highlighted in red. You can modify them (or not) and click on the “configure” button again until satisfied and click on the “generate” button to write out the build files. You can exit the GUI from the “File” menu or hit “ctrl-q”.

## Setting options

Options that enable, disable or modify settings are modified by setting the value of CMake variables. This is done on the command line with the `-D` flag in the format `-D VARIABLE=value`, e.g. `-D CMAKE_BUILD_TYPE=Release` or `-D BUILD_MPI=on`. There is one quirk: when used before the CMake directory, there may be a space between the `-D` flag and the variable, after it must not be. Such CMake variables can have boolean values (on/off, yes/no, or 1/0 are all valid) or are strings representing a choice, or a path, or are free format. If the string would contain whitespace, it must be put in quotes, for example `-D CMAKE_TUNE_FLAGS="-ftree-vectorize -ffast-math"`.

CMake variables fall into two categories: 1) common CMake variables that are used by default for any CMake configuration setup and 2) project specific variables, i.e. settings that are specific for LAMMPS. Also CMake variables can be flagged as *advanced*, which means they are not shown in the text mode or graphical CMake program in the overview of all settings by default, but only when explicitly requested (by hitting the ‘t’ key or clicking on the ‘Advanced’ check-box).

### Some common CMake variables

Variable	Description
CMAKE_INSTALL_PREFIX	root directory of install location for <code>make install</code> (default: <code>\$HOME/.local</code> )
LAMMPS_INSTALL_RPATH	set or remove runtime path setting from binaries for <code>make install</code> (default: <code>off</code> )
CMAKE_BUILD_TYPE	controls compilation options: one of <code>RelWithDebInfo</code> (default), <code>Release</code> , <code>Debug</code> , <code>MinSizeRel</code>
BUILD_SHARED_LIBS	if set to <code>on</code> build the LAMMPS library as shared library (default: <code>off</code> )
CMAKE_MAKE_PROGRAM	name/path of the compilation command (default depends on <code>-G</code> option, usually <code>make</code> )
CMAKE_VERBOSE_MAKEFILE	if set to <code>on</code> echo commands while executing during build (default: <code>off</code> )
CMAKE_C_COMPILER	C compiler to be used for compilation (default: system specific, <code>gcc</code> on Linux)
CMAKE_CXX_COMPILER	C++ compiler to be used for compilation (default: system specific, <code>g++</code> on Linux)
CMAKE_Fortran_COMPILER	Fortran compiler to be used for compilation (default: system specific, <code>gfortran</code> on Linux)
CXX_COMPILER_LAUNCHER	tool to launch the C++ compiler, e.g. <code>ccache</code> or <code>distcc</code> for faster compilation (default: empty)

### Some common LAMMPS specific variables

Variable	Description
BUILD_MPI	build LAMMPS with MPI support (default: <code>on</code> if a working MPI available, else <code>off</code> )
BUILD_OMP	build LAMMPS with OpenMP support (default: <code>on</code> if compiler supports OpenMP fully, else <code>off</code> )
BUILD_TOOLS	compile some additional executables from the <code>tools</code> folder (default: <code>off</code> )
BUILD_LAMMPS_SHELL	compile the LAMMPS shell from the <code>tools/lammps-shell</code> folder (default: <code>off</code> )
BUILD_DOC	include building the HTML format documentation for packaging/installing (default: <code>off</code> )
CMAKE_TUNE_FLAGS	common compiler flags, for optimization or instrumentation (default:)
LAMMPS_MACHINE	when set to name the LAMMPS executable and library will be called <code>lmp_name</code> and <code>liblammps_name.a</code>
LAMMPS_EXCEPTIONS	when set to <code>on</code> errors will throw a C++ exception instead of aborting (default: <code>off</code> )
FFT	select which FFT library to use: <code>FFTW3</code> , <code>MKL</code> , <code>KISS</code> (default, unless <code>FFTW3</code> is found)
FFT_SINGLE	select whether to use single precision FFTs (default: <code>off</code> )
WITH_JPEG	whether to support JPEG format in <i>dump image</i> (default: <code>on</code> if found)
WITH_PNG	whether to support PNG format in <i>dump image</i> (default: <code>on</code> if found)
WITH_GZIP	whether to support reading and writing compressed files (default: <code>on</code> if found)
WITH_FFmpeg	whether to support generating movies with <i>dump movie</i> (default: <code>on</code> if found)

### Enabling or disabling LAMMPS packages

The LAMMPS software is organized into a common core that is always included and a large number of *add-on packages* that have to be enabled to be included into a LAMMPS executable. Packages are enabled through setting variables of the kind `PKG_<NAME>` to `on` and disabled by setting them to `off` (or using `yes`, `no`, `1`, `0` correspondingly). `<NAME>` has to be replaced by the name of the package, e.g. `MOLECULE` or `EXTRA-PAIR`.

## Using presets

Since LAMMPS has a lot of optional features and packages, specifying them all on the command line can be tedious. Or when selecting a different compiler toolchain, multiple options have to be changed consistently and that is rather error prone. Or when enabling certain packages, they require consistent settings to be operated in a particular mode. For this purpose, we are providing a selection of “preset files” for CMake in the folder `cmake/presets`. They represent a way to pre-load or override the CMake configuration cache by setting or changing CMake variables. Preset files are loaded using the `-C` command line flag. You can combine loading multiple preset files or change some variables later with additional `-D` flags. A few examples:

```
cmake -C ../cmake/presets/basic.cmake -D PKG_MISC=on ../cmake
cmake -C ../cmake/presets/clang.cmake -C ../cmake/presets/most.cmake ../cmake
cmake -C ../cmake/presets/basic.cmake -D BUILD_MPI=off ../cmake
```

The first command will install the packages `KSPACE`, `MANYBODY`, `MOLECULE`, `RIGID` and `MISC`; the first four from the preset file and the fifth from the explicit variable definition. The second command will first switch the compiler toolchain to use the Clang compilers and install a large number of packages that are not depending on any special external libraries or tools and are not very unusual. The third command will enable the first four packages like above and then enforce compiling LAMMPS as a serial program (using the `MPI STUBS` library).

It is also possible to do this incrementally.

```
cmake -C ../cmake/presets/basic.cmake ../cmake
cmake -D PKG_MISC=on .
```

will achieve the same final configuration as in the first example above. In this scenario it is particularly convenient to do the second configuration step using either the text mode or graphical user interface (`ccmake` or `cmake-gui`).

---

**Note:** Using a preset to select a compiler package (`clang.cmake`, `gcc.cmake`, `intel.cmake`, `oneapi.cmake`, or `pgi.cmake`) are an exception to the mechanism of updating the configuration incrementally, as they will trigger a reset of cached internal CMake settings and thus reset settings to their default values.

---

## Compilation and build targets

The actual compilation will be started by running the selected build command (on Linux this is by default `make`, see below how to select alternatives). You can also use the portable command `cmake --build .` which will adapt to whatever the selected build command is. This is particularly convenient, if you have set a custom build command via the `CMAKE_MAKE_PROGRAM` variable.

When calling the build program, you can also select which “target” is to be build through appending the `--target` flag and the name of the target to the build command. When using `make` as build tool, you can just append the target name to the command. Example: `cmake --build . --target all` or `make all`. The following abstract targets are available:

Target	Description
<code>all</code>	build “everything” (default)
<code>lammps</code>	build the LAMMPS library and executable
<code>doc</code>	build the HTML documentation (if configured)
<code>install</code>	install all target files into folders in <code>CMAKE_INSTALL_PREFIX</code>
<code>test</code>	run some tests (if configured with <code>-D ENABLE_TESTING=on</code> )
<code>clean</code>	remove all generated files

## Choosing generators

While CMake usually defaults to creating makefiles to compile software with the make program, it supports multiple alternate build tools (e.g. `ninja-build` which tends to be faster and more efficient in parallelizing builds than `make`) and can generate project files for integrated development environments (IDEs) like VisualStudio, Eclipse or CodeBlocks. This is specific to how the local CMake version was configured and compiled. The list of available options can be seen at the end of the output of `cmake --help`. Example on Fedora 31 this is:

### Generators

The following generators are available on this platform (\* marks default):

```
* Unix Makefiles           = Generates standard UNIX makefiles.
  Green Hills MULTI       = Generates Green Hills MULTI files
                          (experimental, work-in-progress).
  Ninja                   = Generates build.ninja files.
  Ninja Multi-Config      = Generates build-<Config>.ninja files.
  Watcom WMake            = Generates Watcom WMake makefiles.
  CodeBlocks - Ninja      = Generates CodeBlocks project files.
  CodeBlocks - Unix Makefiles = Generates CodeBlocks project files.
  CodeLite - Ninja        = Generates CodeLite project files.
  CodeLite - Unix Makefiles = Generates CodeLite project files.
  Sublime Text 2 - Ninja  = Generates Sublime Text 2 project files.
  Sublime Text 2 - Unix Makefiles
                          = Generates Sublime Text 2 project files.
  Kate - Ninja            = Generates Kate project files.
  Kate - Unix Makefiles   = Generates Kate project files.
  Eclipse CDT4 - Ninja    = Generates Eclipse CDT 4.0 project files.
  Eclipse CDT4 - Unix Makefiles= Generates Eclipse CDT 4.0 project files.
```

Below is a screenshot of using the CodeBlocks IDE with the ninja build tool after running CMake as follows:

```
cmake -G 'CodeBlocks - Ninja' ../cmake/presets/most.cmake ../cmake/
```

The screenshot shows the Code::Blocks IDE with the following components:

- Management Panel:** Shows the project structure for 'lammmps' with subfolders for CMake Files, Sources, and Headers.
- Source Editor:** Displays the `src/lammps.cpp` file, showing the `LAMMPS::LAMMPS` constructor function. Lines 91-113 are visible, including the function signature and the initialization of various pointers like `memory`, `error`, `universe`, `clientserver`, `cslib`, `cscomm`, `screen`, `logfile`, `infile`, `initclock`, and `init_pkg_lists`.
- Output Window:** Shows the execution output for the 'lammmps' process. It includes:
  - Performance metrics: Loop time of 0.408998 on 1 procs for 250 steps with 4000 atoms. Performance: 264059,963 tau/day, 611,250 timesteps/s, 99.9% CPU use with 1 MPI tasks x 1 OpenMP threads.
  - MPI task timing breakdown table:
  - System statistics: Nlocal, Histogram, Nghost, Histogram, Neighs, Histogram.
  - Summary: Total # of neighbors = 151513, Ave neighs/atom = 37.8783, Neighbor list builds = 12, Dangerous builds not checked, Total wall time: 0:00:00.
  - Process termination: Process returned 0 (0x0) execution time : 0,457 s. Press ENTER to continue.
  - Build command: `Run: all in lammmps (compiler: GNU GCC Compiler)`
- Log & others:** Shows the compilation process, including the command `g++ -P /home/akohmey/compile/lammps/... -o lammmps` and the message 'Process terminated with status 0 (0 minutes), 0 error(s), 0 warning(s) (0 minute(s), 0 second(s))'.
- Status Bar:** Shows the current file path, compiler, encoding, cursor position, and mode.

## 8.6.2 LAMMPS GitHub tutorial

written by Stefan Paquay

This document describes the process of how to use GitHub to integrate changes or additions you have made to LAMMPS into the official LAMMPS distribution. It uses the process of updating this very tutorial as an example to describe the individual steps and options. You need to be familiar with git and you may want to have a look at the [git book](#) to familiarize yourself with some of the more advanced git features used below.

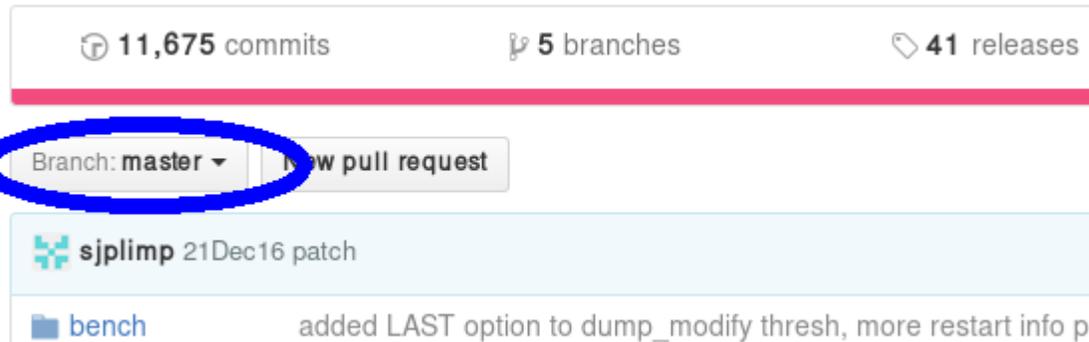
As of fall 2016, submitting contributions to LAMMPS via pull requests on GitHub is the preferred option for integrating contributed features or improvements to LAMMPS, as it significantly reduces the amount of work required by the LAMMPS developers. Consequently, creating a pull request will increase your chances to have your contribution included and will reduce the time until the integration is complete. For more information on the requirements to have your code included into LAMMPS please see [this page](#).

### Making an account

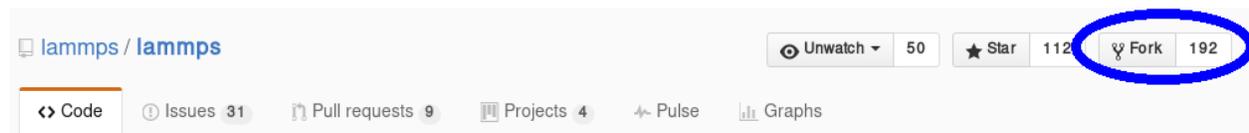
First of all, you need a GitHub account. This is fairly simple, just go to [GitHub](#) and create an account by clicking the “Sign up for GitHub” button. Once your account is created, you can sign in by clicking the button in the top left and filling in your username or e-mail address and password.

## Forking the repository

To get changes into LAMMPS, you need to first fork the *lammps/lammps* repository on GitHub. At the time of writing, *develop* is the preferred target branch. Thus go to [LAMMPS on GitHub](#) and make sure branch is set to “develop”, as shown in the figure below.



If it is not, use the button to change it to *develop*. Once it is, use the fork button to create a fork.



This will create a fork (which is essentially a copy, but uses less resources) of the LAMMPS repository under your own GitHub account. You can make changes in this fork and later file *pull requests* to allow the upstream repository to merge changes from your own fork into the one we just forked from (or others that were forked from the same repository). At the same time, you can set things up, so you can include changes from upstream into your repository and thus keep it in sync with the ongoing LAMMPS development.

## Adding changes to your own fork

Additions to the upstream version of LAMMPS are handled using *feature branches*. For every new feature, a so-called feature branch is created, which contains only those modification relevant to one specific feature. For example, adding a single fix would consist of creating a branch with only the fix header and source file and nothing else. It is explained in more detail here: [feature branch workflow](#).

### Feature branches

First of all, create a clone of your version on GitHub on your local machine via HTTPS:

```
$ git clone https://github.com/<your user name>/lammps.git <some name>
```

or, if you have set up your GitHub account for using SSH keys, via SSH:

```
$ git clone git@github.com:<your user name>/lammps.git
```

You can find the proper URL by clicking the “Clone or download”-button:



Pakketeret2 / lammps  
forked from lammps/lammps

Unwatch 1 Star 0 Fork 192

Code Pull requests 0 Projects 0 Wiki Pulse Graphs Settings

Public/backup repository of the LAMMPS MD software package <http://lammps.sandia.gov> Edit

16,191 commits 8 branches 10 releases 12 contributors GPL-2.0

Branch: master New pull request Create new file Upload files Clone or download

This branch is 6178 commits ahead, 1662 commits behind lammps:master.

Pakketeret2 Merged stuffs.

- bench Merge remote branch 'lammps-ro/master' into lammps-icm
- doc Updated docs.

Clone with SSH Use an SSH key and passphrase from account.  
git@github.com:Pakketeret2/lammps.git Download ZIP

The above command copies (“clones”) the git repository to your local machine to a directory with the name you chose. If none is given, it will default to “lammps”. Typical names are “mylammps” or something similar.

You can use this local clone to make changes and test them without interfering with the repository on GitHub.

To pull changes from upstream into this copy, you can go to the directory and use git pull:

```
$ cd mylammps
$ git checkout develop
$ git pull https://github.com/lammps/lammps develop
```

You can also add this URL as a remote:

```
$ git remote add upstream https://www.github.com/lammps/lammps
```

From then on you can update your upstream branches with:

```
$ git fetch upstream
```

and then refer to the upstream repository branches with *upstream/develop* or *upstream/release* and so on.

At this point, you typically make a feature branch from the updated branch for the feature you want to work on. This tutorial contains the workflow that updated this tutorial, and hence we will call the branch “github-tutorial-update”:

```
$ git fetch upstream
$ git checkout -b github-tutorial-update upstream/develop
```

Now that we have changed branches, we can make our changes to our local repository. Just remember that if you want to start working on another, unrelated feature, you should switch branches!

---

**Note:** Committing changes to the *develop*, *release*, or *stable* branches is strongly discouraged. While it may be convenient initially, it will create more work in the long run. Various texts and tutorials on using git effectively discuss the motivation for using feature branches instead.

---

### After changes are made

After everything is done, add the files to the branch and commit them:

```
$ git add doc/src/Howto_github.txt
$ git add doc/src/JPG/tutorial*.png
```

**Warning:** Do not use `git commit -a` (or `git add -A`). The `-a` flag (or `-A` flag) will automatically include **all** modified **and** new files and that is rarely the behavior you want. It can easily lead to accidentally adding unrelated and unwanted changes into the repository. Instead it is preferable to explicitly use `git add`, `git rm`, `git mv` for adding, removing, renaming individual files, respectively, and then `git commit` to finalize the commit. Carefully check all pending changes with `git status` before committing them. If you find doing this on the command line too tedious, consider using a GUI, for example the one included in git distributions written in Tk, i.e. use `git gui` (on some Linux distributions it may be required to install an additional package to use it).

After adding all files, the change set can be committed with some useful message that explains the change.

```
$ git commit -m 'Finally updated the GitHub tutorial'
```

After the commit, the changes can be pushed to the same branch on GitHub:

```
$ git push
```

Git will ask you for your user name and password on GitHub if you have not configured anything. If your local branch is not present on GitHub yet, it will ask you to add it by running

```
$ git push --set-upstream origin github-tutorial-update
```

If you correctly type your user name and password, the feature branch should be added to your fork on GitHub.

If you want to make really sure you push to the right repository (which is good practice), you can provide it explicitly:

```
$ git push origin
```

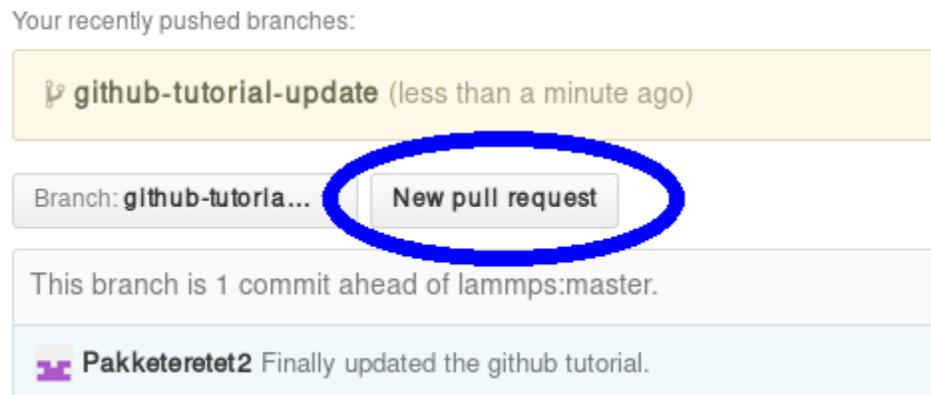
or using an explicit URL:

```
$ git push git@github.com:Pakketeretet2/lammps.git
```

---

### Filing a pull request

Up to this point in the tutorial, all changes were to *your* clones of LAMMPS. Eventually, however, you want this feature to be included into the official LAMMPS version. To do this, you will want to file a pull request by clicking on the “New pull request” button:

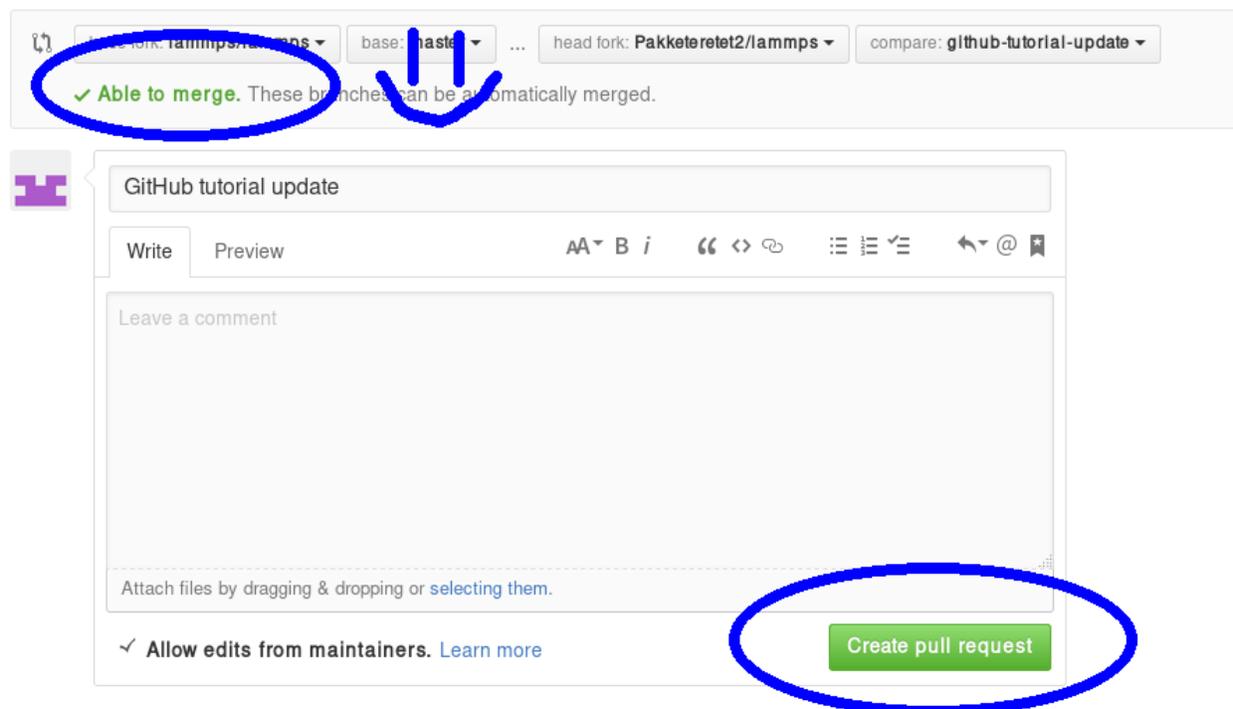


Make sure that the current branch is set to the correct one, which, in this case, is “github-tutorial-update”. If done correctly, the only changes you will see are those that were made on this branch.

This will open up a new window that lists changes made to the repository. If you are just adding new files, there is not much to do, but I suppose merge conflicts are to be resolved here if there are changes in existing files. If all changes can automatically be merged, green text at the top will say so and you can click the “Create pull request” button, see image.

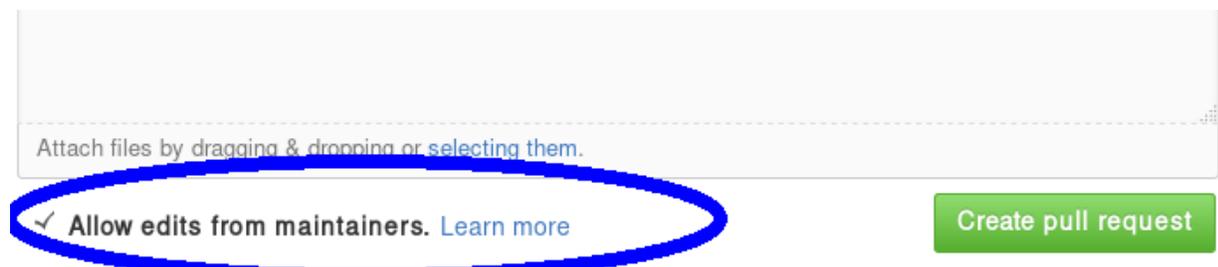
## Open a pull request

Create a new pull request by comparing changes across two branches. If you need to, you can also [compare across forks](#).



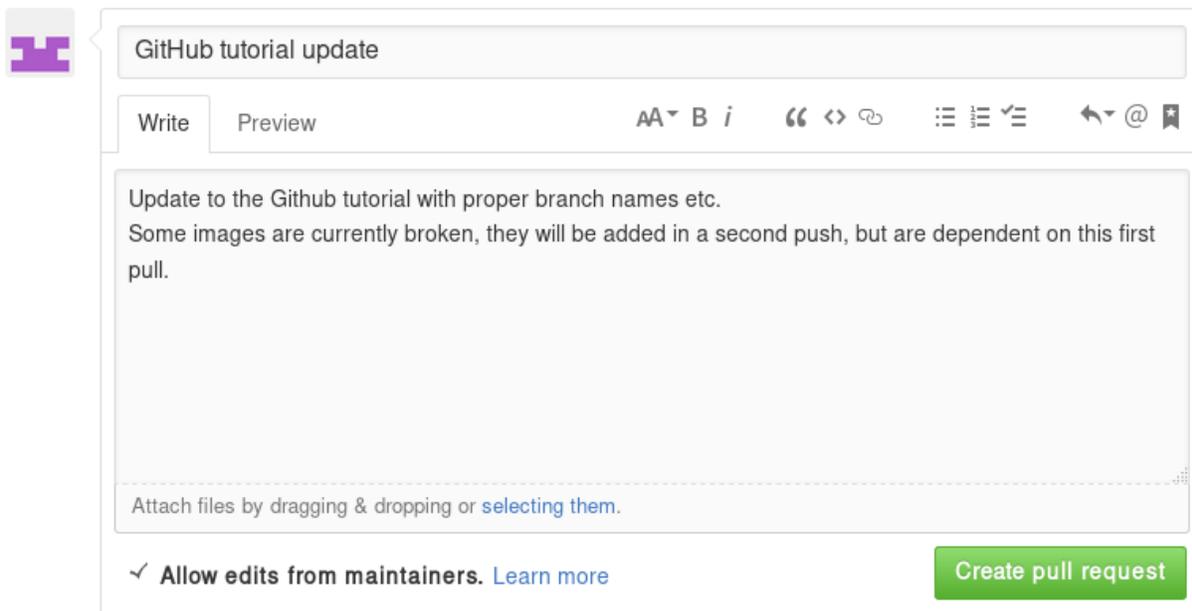
Before creating the pull request, make sure the short title is accurate and add a comment with details about your pull request. Here you write what your modifications do and why they should be incorporated upstream.

Note the checkbox that says “Allow edits from maintainers”. This is checked by default checkbox (although in my version of Firefox, only the checkmark is visible):



If it is checked, maintainers can immediately add their own edits to the pull request. This helps the inclusion of your branch significantly, as simple/trivial changes can be added directly to your pull request branch by the LAMMPS maintainers. The alternative would be that they make changes on their own version of the branch and file a reverse pull request to you. Just leave this box checked unless you have a very good reason not to.

Now just write some nice comments and click on “Create pull request”.



---

### After filing a pull request

**Note:** When you submit a pull request (or ask for a pull request) for the first time, you will receive an invitation to become a LAMMPS project collaborator. Please accept this invite as being a collaborator will simplify certain administrative tasks and will probably speed up the merging of your feature, too.

---

You will notice that after filing the pull request, some checks are performed automatically:

## GitHub tutorial update #315

**Open** Pakketeret2 wants to merge 2 commits into lammps:master from Pakketeret2:github-tutorial-update

Conversation 0 Commits 2 Files changed 5

**Pakketeret2** commented a minute ago Collaborator

Update to the Github tutorial with proper branch names etc.  
Some images are currently broken, they will be added in a second push, but are dependent on this first pull.

**Pakketeret2** added some commits 18 minutes ago

- Finally updated the github tutorial. 391ab76
- Almost done with the tutorial now. 4d98bbd

Add more commits by pushing to the `github-tutorial-update` branch on `Pakketeret2/lammps`.

**Some checks haven't completed yet** Hide all checks  
3 pending checks

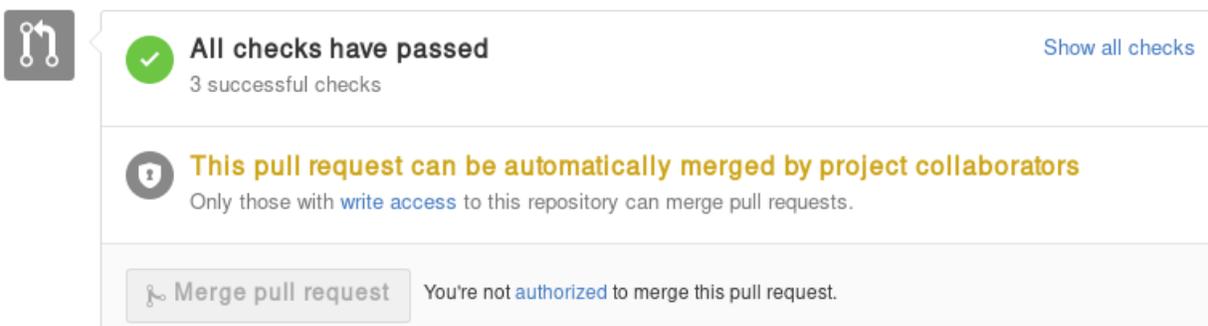
- `lammps/pull-requests/openmpi-pr` — head run star... [Details](#)
- `lammps/pull-requests/serial-pr` — head run started [Details](#)
- `lammps/pull-requests/shlib-pr` — head run started [Details](#)

**This pull request can be automatically merged by project collaborators**  
Only those with [write access](#) to this repository can merge pull requests.

[Merge pull request](#) You're not [authorized](#) to merge this pull request.

If all is fine, you will see this:

Add more commits by pushing to the `github-tutorial-update` branch on `Pakketeret2/lammps`.



**All checks have passed** [Show all checks](#)  
3 successful checks

**This pull request can be automatically merged by project collaborators**  
Only those with [write access](#) to this repository can merge pull requests.

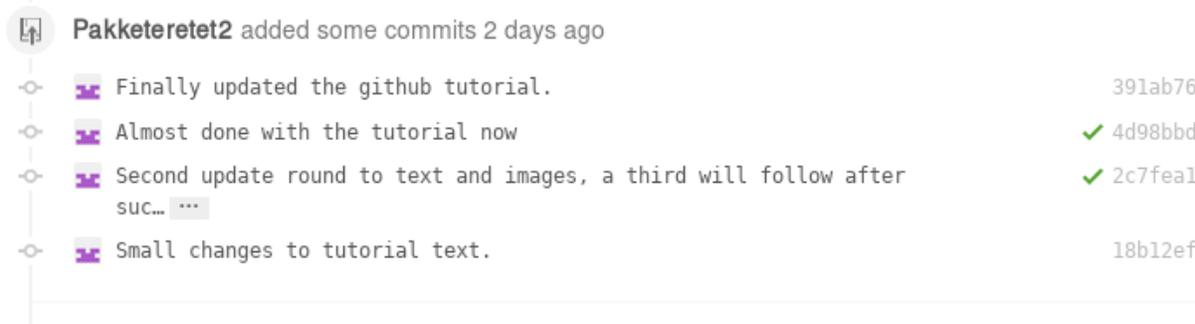
[Merge pull request](#) You're not [authorized](#) to merge this pull request.

If any of the checks are failing, your pull request will not be processed, as your changes may break compilation for certain configurations or may not merge cleanly. It is your responsibility to remove the reason(s) for the failed test(s). If you need help with this, please contact the LAMMPS developers by adding a comment explaining your problems with resolving the failed tests.

A few further interesting things (can) happen to pull requests before they are included.

### Additional changes

First of all, any additional changes you push into your branch in your repository will automatically become part of the pull request:



**Pakketeret2** added some commits 2 days ago

- Finally updated the github tutorial. 391ab76
- Almost done with the tutorial now ✓ 4d98bbd
- Second update round to text and images, a third will follow after suc... ✓ 2c7feal
- Small changes to tutorial text. 18b12ef

This means you can add changes that should be part of the feature after filing the pull request, which is useful in case you have forgotten them, or if a developer has requested that something needs to be changed before the feature can be accepted into the official LAMMPS version. After each push, the automated checks are run again.

### Labels

LAMMPS developers may add labels to your pull request to assign it to categories (mostly for bookkeeping purposes), but a few of them are important: *needs\_work*, *work\_in\_progress*, *run\_tests*, *test\_for\_regression*, and *ready\_for\_merge*. The first two indicate, that your pull request is not considered to be complete. With “needs\_work” the burden is on exclusively on you; while “work\_in\_progress” can also mean, that a LAMMPS developer may want to add changes. Please watch the comments to the pull requests. The two “test” labels are used to trigger extended tests before the code is merged. This is sometimes done by LAMMPS developers, if they suspect that there may be some subtle side effects from your changes. It is not done by default, because those tests are very time consuming. The *ready\_for\_merge* label is usually attached when the LAMMPS developer assigned to the pull request considers this request complete and to trigger a final full test evaluation.

### Reviews

As of Fall 2021, a pull request needs to pass all automatic tests and at least 1 approving review from a LAMMPS developer with write access to the repository before it is eligible for merging. In case your changes touch code that certain developers are associated with, they are auto-requested by the GitHub software. Those associations are set in the file `.github/CODEOWNERS` Thus if you want to be automatically notified to review when anybody changes files or

packages, that **you** have contributed to LAMMPS, you can add suitable patterns to that file, or a LAMMPS developer may add you.

Otherwise, you can also manually request reviews from specific developers, or LAMMPS developers - in their assessment of your pull request - may determine who else should be reviewing your contribution and add that person. Through reviews, LAMMPS developers also may request specific changes from you. If those are not addressed, your pull requests cannot be merged.

### Assignees

There is an assignee property for pull requests. If the request has not been reviewed by any developer yet, it is not assigned to anyone. After revision, a developer can choose to assign it to either a) you, b) a LAMMPS developer (including him/herself) or c) Axel Kohlmeyer (akohlmey).

- Case a) happens if changes are required on your part
- Case b) means that at the moment, it is being tested and reviewed by a LAMMPS developer with the expectation that some changes would be required. After the review, the developer can choose to implement changes directly or suggest them to you.
- Case c) means that the pull request has been assigned to the developer overseeing the merging of pull requests into the *develop* branch.

In this case, Axel assigned the tutorial to Steve:

Update to the Simba tutorial with proper branch names etc.  
Some images are currently broken, they will be added in a second push, but are dependent on this first pull.

**Assignees**  
sjplimp

**Labels**  
documentation

**Projects**  
None yet

**Milestone**  
No milestone

Pakketeret2 added some commits 2 days ago

- Finally updated the github tutorial. 391ab76
- Almost done with the tutorial now ✓ 4d98bbd
- Second update round to text and images, a third will follow after suc... ✓ 2c7feal
- Small changes to tutorial text. 18b12ef

sjplimp was assigned by akohlmey 2 days ago

### Edits from LAMMPS maintainers

If you allowed edits from maintainers (the default), any LAMMPS maintainer can add changes to your pull request. In this case, both Axel and Richard made changes to the tutorial:

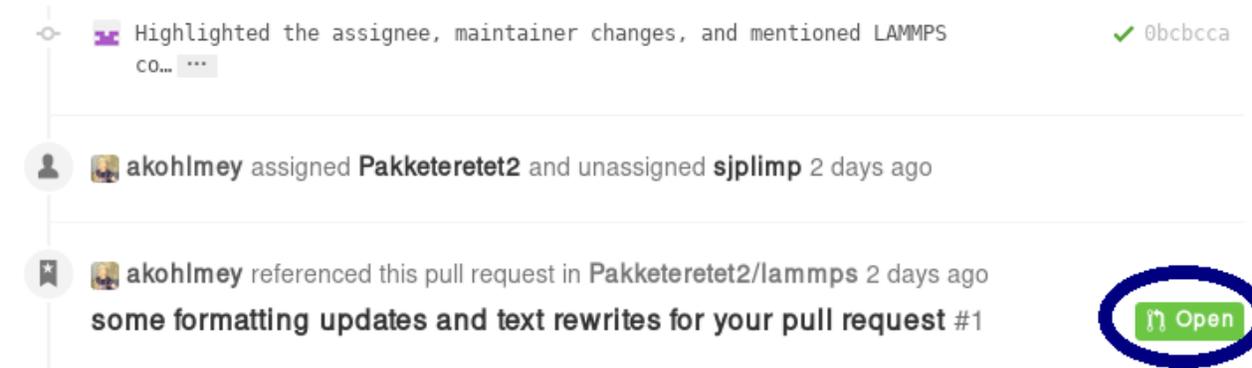
Pakketeret2 and others added some commits 2 days ago

- Updated some inconsistent text. ✓ 4f096db
- make it more explicit, that master needs to be updated and feature br... ✓ 7d057d4
- Add warning formatting ✓ 4f45d39

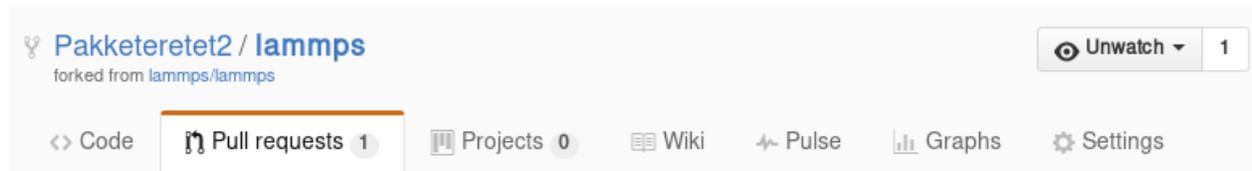
### Reverse pull requests

Sometimes, however, you might not feel comfortable having other people push changes into your own branch, or maybe the maintainers are not sure their idea was the right one. In such a case, they can make changes, reassign you as the assignee, and file a “reverse pull request”, i.e. file a pull request in **your** forked GitHub repository to include changes in the branch, that you have submitted as a pull request yourself. In that case, you can choose to merge their changes

back into your branch, possibly make additional changes or corrections and proceed from there. It looks something like this:



For some reason, the highlighted button did not work in my case, but I can go to my own repository and merge the pull request from there:



## some formatting updates and text rewrites for your pull request #1

[Open](#) akohlmei wants to merge 1 commit into `Pakketeretet2:github-tutorial-update` from `akohlmei:pull-315`

Conversation 0 Commits 1 Files changed 1

akohlmei commented 2 days ago

Here are some additional changes for your lammps/#315 pull request. It updates the "do not use git add -a" paragraph and some formatting improvements.

some formatting updates and text rewrites in the "do not use git add ..."



Be sure to check the changes to see if you agree with them by clicking on the tab button:



## some formatting updates and text rewrites for your pull request #1

 **akohlmey** wants to merge 1 commit into `Pakketeret2:github-tutorial-update` from `akohlmey:pull-315`

Conversation 0   Commits 1   **Files changed 1**

 **akohlmey** commented 2 days ago

Here are some additional changes for your lammps/#315 pull request. It updates the "do not use git add -a" paragraph and some formatting improvements.

 **Pakketeret2** commented 26 seconds ago Owner

Thanks Axel, will merge after a review.

 **This branch has no conflicts with the base branch**  
Merging can be performed automatically.

**Merge pull request** or view [command line instructions](#).

In this case, most of it is changes in the markup and a short rewrite of Axel's explanation of the "git gui" and "git add" commands.

Changes from all commits ▾ 1 file ▾ +37 -31

<pre> 112 \$ git add doc/src/JPG/tutorial_*.png :pre 113 114 -IMPORTANT NOTE: Do not use 'git commit -a'. The -a flag will automatically 115 -include _all_ modified or new files and that is rarely the behavior you want. 116 -It can easily create to accidentally adding unrelated and unwanted changes into 117 -the repository. It is highly preferable to explicitly use 'git add', 'git rm', 118 -'git mv' for adding, removing, renaming files, respectively, and then 'git 119 -commit' to finalize the commit. If you find doing this on the command line too 120 -tedious, consider using a GUI, the one included in git distributions written in 121 -Tk, i.e. use 'git gui'. 122 123 -After adding all files, the change can be committed with some useful message 124 -that explains the change. 125 126 \$ git commit -m 'Finally updated the github tutorial' :pre 127 128 @@ -213,23 +216,26 @@ repository will automatically become part of the pull request: 129 130 :c,image(JPG/tutorial_additional_changes.png) </pre>	<pre> 113 \$ git add doc/src/JPG/tutorial_*.png :pre 114 115 +IMPORTANT NOTE: Do not use 'git commit -a'. The -a flag will 116 +automatically include _all_ modified or new files and that is rarely the 117 +behavior you want. It can easily lead to accidentally adding unrelated 118 +and unwanted changes into the repository. Instead it is preferable to 119 +explicitly use 'git add', 'git rm', 'git mv' for adding, removing, 120 +renaming files, respectively, and then 'git commit' to finalize the 121 +commit. If you find doing this on the command line too tedious, 122 +consider using a GUI, for example the one included in git distributions 123 +written in Tk, i.e. use 'git gui' (on some Linux distributions it may 124 +be required to install an additional package to use it). 125 126 +After adding all files, the change <b>set</b> can be committed with some 127 +<b>useful message</b> that explains the change. 128 129 \$ git commit -m 'Finally updated the github tutorial' :pre 130 131 132 :c,image(JPG/tutorial_additional_changes.png) </pre>
---	---

Because the changes are OK with us, we are going to merge by clicking on "Merge pull request". After a merge it looks like this:

A screenshot of a GitHub interface. At the top, a comment from user 'Pakketeretet2' says 'Thanks Axel, will merge after a review.' Below this, a notification indicates that 'Pakketeretet2' merged commit '1310438' into the branch 'Pakketeretet2:github-tutorial-update'. A 'Revert' button is visible. At the bottom, there is a light blue notification box with a lightbulb icon that says 'Avoid bugs by automatically running your tests.' and provides information about continuous integration.

Now, since in the meantime our local text for the tutorial also changed, we need to pull Axel's change back into our branch, and merge them:

```
$ git add Howto_github.txt
$ git add JPG/tutorial_reverse_pull_request*.png
$ git commit -m "Updated text and images on reverse pull requests"
$ git pull
```

In this case, the merge was painless because git could auto-merge:

```
[ stefan@phys8250 src ] % git pull
Auto-merging doc/src/tutorial_github.txt
Waiting for Emacs...
Merge made by the 'recursive' strategy.
 doc/src/tutorial_github.txt | 68 ++++++-----
 1 file changed, 37 insertions(+), 31 deletions(-)
[ stefan@phys8250 src ] %
```

With Axel's changes merged in and some final text updates, our feature branch is now perfect as far as we are concerned, so we are going to commit and push again:

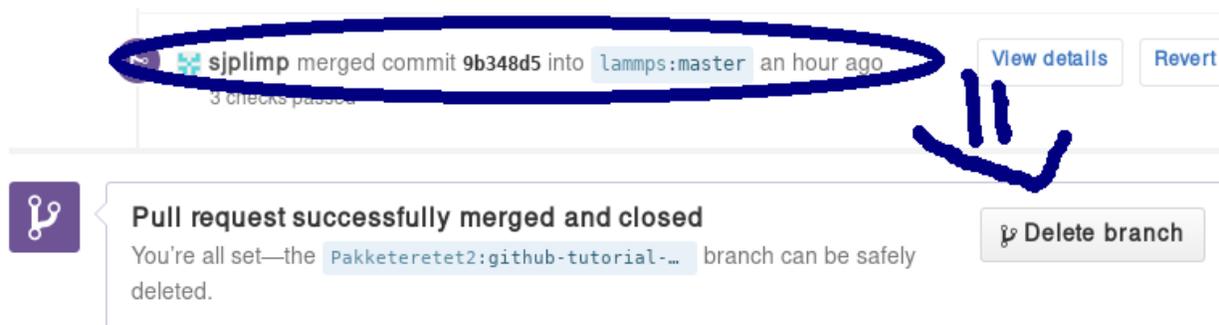
```
$ git add Howto_github.txt
$ git add JPG/tutorial_reverse_pull_request6.png
$ git commit -m "Merged Axel's suggestions and updated text"
$ git push git@github.com:Pakketeretet2/lammps
```

This merge also shows up on the lammps GitHub page:

A screenshot of a GitHub pull request page. The pull request title is 'polish the introduction, some more clarifications, corrections and fo...'. It is marked as 'Merged' with a green checkmark and the commit hash '6af56e6'. A comment from user 'akohlmei' says 'A few more suggested edits to the tutorial from me #2'. Another comment from 'akohlmei' says 'assigned sjplimp and unassigned Pakketeretet2 2 hours ago'.

After a merge

When everything is fine, the feature branch is merged into the *develop* branch:



Now one question remains: What to do with the feature branch that got merged into upstream?

It is in principle safe to delete them from your own fork. This helps keep it a bit more tidy. Note that you first have to switch to another branch!

```
$ git checkout develop
$ git pull https://github.com/lammps/lammps develop
$ git branch -d github-tutorial-update
```

If you do not pull first, it is not really a problem but git will warn you at the next statement that you are deleting a local branch that was not yet fully merged into HEAD. This is because git does not yet know your branch just got merged into LAMMPS upstream. If you first delete and then pull, everything should still be fine. You can display all branches that are fully merged by:

Finally, if you delete the branch locally, you might want to push this to your remote(s) as well:

```
$ git push origin :github-tutorial-update
```

### Recent changes in the workflow

Some changes to the workflow are not captured in this tutorial. For example, in addition to the *develop* branch, to which all new features should be submitted, there is also a *release* and a *stable* branch; these have the same content as *develop*, but are only updated after a patch release or stable release was made. Furthermore, the naming of the patches now follow the pattern “patch\_<Day><Month><Year>” to simplify comparisons between releases. Finally, all patches and submissions are subject to automatic testing and code checks to make sure they at the very least compile.

A discussion of the LAMMPS developer GitHub workflow can be found in the file [doc/github-development-workflow.md](#)

## 8.6.3 PyLammps Tutorial

### Contents

- *PyLammps Tutorial*
  - *Overview*
    - \* *Comparison of lammps and PyLammps interfaces*
      - *lammps.lammps*
      - *lammps.PyLammps*
  - *Quick Start*

- \* *System-wide Installation*
  - *Step 1: Building LAMMPS as a shared library*
  - *Step 2: Installing the LAMMPS Python package*
- \* *Installation inside of a virtualenv*
  - *Benefits of using a virtualenv*
  - *Creating a virtualenv with lammps installed*
- *Creating a new instance of PyLammps*
- *Commands*
- *System state*
- *Working with LAMMPS variables*
- *Retrieving the value of an arbitrary LAMMPS expressions*
- *Accessing atom data*
- *Evaluating thermo data*
- *Error handling with PyLammps*
- *Using PyLammps in IPython notebooks and Jupyter*
- *IPyLammps Examples*
  - \* *Validating a dihedral potential*
  - \* *Running a Monte Carlo relaxation*
- *Using PyLammps and mpi4py (Experimental)*
- *Feedback and Contributing*

## Overview

PyLammps is a Python wrapper class for LAMMPS which can be created on its own or use an existing lammps Python object. It creates a simpler, more “pythonic” interface to common LAMMPS functionality, in contrast to the lammps wrapper for the C-style LAMMPS library interface which is written using [Python ctypes](#). The lammps wrapper is discussed on the [Use Python with LAMMPS](#) doc page.

Unlike the flat ctypes interface, PyLammps exposes a discoverable API. It no longer requires knowledge of the underlying C++ code implementation. Finally, the IPyLammps wrapper builds on top of PyLammps and adds some additional features for IPython integration into Jupyter notebooks, e.g. for embedded visualization output from *dump style image*.

## Comparison of lammps and PyLammps interfaces

### lammps.lammps

- uses ctypes
- direct memory access to native C++ data
- provides functions to send and receive data to LAMMPS
- requires knowledge of how LAMMPS internally works (C pointers, etc)

## lammeps.PyLammps

- higher-level abstraction built on top of original ctypes interface
- manipulation of Python objects
- communication with LAMMPS is hidden from API user
- shorter, more concise Python
- better IPython integration, designed for quick prototyping

## Quick Start

### System-wide Installation

#### Step 1: Building LAMMPS as a shared library

To use LAMMPS inside of Python it has to be compiled as shared library. This library is then loaded by the Python interface. In this example we enable the MOLECULE package and compile LAMMPS with C++ exceptions, PNG, JPEG and FFMPEG output support enabled.

Step 1a: For the CMake based build system, the steps are:

```
mkdir $LAMMPS_DIR/build-shared
cd $LAMMPS_DIR/build-shared

# MPI, PNG, Jpeg, FFMPEG are auto-detected
cmake ../cmake -DPKG_MOLECULE=yes -DLAMMPS_EXCEPTIONS=yes -DBUILD_LIB=yes -DBUILD_SHARED_
↳LIBS=yes
make
```

Step 1b: For the legacy, make based build system, the steps are:

```
cd $LAMMPS_DIR/src

# add packages if necessary
make yes-MOLECULE

# compile shared library using Makefile
make mpi mode=shlib LMP_INC="-DLAMMPS_PNG -DLAMMPS_JPEG -DLAMMPS_FFMPEG -DLAMMPS_
↳EXCEPTIONS" JPG_LIB="-lpng -ljpeg"
```

#### Step 2: Installing the LAMMPS Python package

PyLammps is part of the lammeps Python package. To install it simply install that package into your current Python installation with:

```
make install-python
```

---

**Note:** Recompiling the shared library requires re-installing the Python package

---

## Installation inside of a virtualenv

You can use virtualenv to create a custom Python environment specifically tuned for your workflow.

### Benefits of using a virtualenv

- isolation of your system Python installation from your development installation
- installation can happen in your user directory without root access (useful for HPC clusters)
- installing packages through pip allows you to get newer versions of packages than e.g., through apt-get or yum package managers (and without root access)
- you can even install specific old versions of a package if necessary

### Prerequisite (e.g. on Ubuntu)

```
apt-get install python-virtualenv
```

### Creating a virtualenv with lammmps installed

```
# create virtualenv named 'testing'
virtualenv $HOME/python/testing

# activate 'testing' environment
source $HOME/python/testing/bin/activate
```

Now configure and compile the LAMMPS shared library as outlined above. When using CMake and the shared library has already been build, you need to re-run CMake to update the location of the python executable to the location in the virtual environment with:

```
cmake . -DPYTHON_EXECUTABLE=$(which python)

# install LAMMPS package in virtualenv
(testing) make install-python

# install other useful packages
(testing) pip install matplotlib jupyter mpi4py

...

# return to original shell
(testing) deactivate
```

## Creating a new instance of PyLammps

To create a PyLammps object you need to first import the class from the lammps module. By using the default constructor, a new *lammps* instance is created.

```
from lammps import PyLammps
L = PyLammps()
```

You can also initialize PyLammps on top of this existing *lammps* object:

```
from lammps import lammps, PyLammps
lmp = lammps()
L = PyLammps(ptr=lmp)
```

## Commands

Sending a LAMMPS command with the existing library interfaces is done using the command method of the lammps object instance.

For instance, let's take the following LAMMPS command:

```
region box block 0 10 0 5 -0.5 0.5
```

In the original interface this command can be executed with the following Python code if *L* was a lammps instance:

```
L.command("region box block 0 10 0 5 -0.5 0.5")
```

With the PyLammps interface, any command can be split up into arbitrary parts separated by white-space, passed as individual arguments to a region method.

```
L.region("box block", 0, 10, 0, 5, -0.5, 0.5)
```

Note that each parameter is set as Python literal floating-point number. In the PyLammps interface, each command takes an arbitrary parameter list and transparently merges it to a single command string, separating individual parameters by white-space.

The benefit of this approach is avoiding redundant command calls and easier parameterization. In the original interface parameterization needed to be done manually by creating formatted strings.

```
L.command("region box block %f %f %f %f %f %f" % (xlo, xhi, ylo, yhi, zlo, zhi))
```

In contrast, methods of PyLammps accept parameters directly and will convert them automatically to a final command string.

```
L.region("box block", xlo, xhi, ylo, yhi, zlo, zhi)
```

## System state

In addition to dispatching commands directly through the PyLammps object, it also provides several properties which allow you to query the system state.

### **L.system**

Is a dictionary describing the system such as the bounding box or number of atoms

### **L.system.xlo, L.system.xhi**

bounding box limits along x-axis

### **L.system.ylo, L.system.yhi**

bounding box limits along y-axis

### **L.system.zlo, L.system.zhi**

bounding box limits along z-axis

### **L.communication**

configuration of communication subsystem, such as the number of threads or processors

### **L.communication.nthreads**

number of threads used by each LAMMPS process

### **L.communication.nprocs**

number of MPI processes used by LAMMPS

### **L.fixes**

List of fixes in the current system

### **L.computes**

List of active computes in the current system

### **L.dump**

List of active dumps in the current system

### **L.groups**

List of groups present in the current system

## Working with LAMMPS variables

LAMMPS variables can be both defined and accessed via the PyLammps interface.

To define a variable you can use the *variable* command:

```
L.variable("a index 2")
```

A dictionary of all variables is returned by `L.variables`

you can access an individual variable by retrieving a variable object from the `L.variables` dictionary by name

```
a = L.variables['a']
```

The variable value can then be easily read and written by accessing the `value` property of this object.

```
print(a.value)
a.value = 4
```



## Retrieving the value of an arbitrary LAMMPS expressions

LAMMPS expressions can be immediately evaluated by using the `eval` method. The passed string parameter can be any expression containing global thermo values, variables, compute or fix data.

```
result = L.eval("ke") # kinetic energy
result = L.eval("pe") # potential energy

result = L.eval("v_t/2.0")
```

## Accessing atom data

All atoms in the current simulation can be accessed by using the `L.atoms` list. Each element of this list is an object which exposes its properties (`id`, `type`, `position`, `velocity`, `force`, etc.).

```
# access first atom
L.atoms[0].id
L.atoms[0].type

# access second atom
L.atoms[1].position
L.atoms[1].velocity
L.atoms[1].force
```

Some properties can also be used to set:

```
# set position in 2D simulation
L.atoms[0].position = (1.0, 0.0)

# set position in 3D simulation
L.atoms[0].position = (1.0, 0.0, 1.)
```

## Evaluating thermo data

Each simulation run usually produces thermo output based on system state, computes, fixes or variables. The trajectories of these values can be queried after a run via the `L.runs` list. This list contains a growing list of run data. The first element is the output of the first run, the second element that of the second run.

```
L.run(1000)
L.runs[0] # data of first 1000 time steps

L.run(1000)
L.runs[1] # data of second 1000 time steps
```

Each run contains a dictionary of all trajectories. Each trajectory is accessible through its thermo name:

```
L.runs[0].thermo.Step # list of time steps in first run
L.runs[0].thermo.Ke  # list of kinetic energy values in first run
```

Together with `matplotlib` plotting data out of LAMMPS becomes simple:

```
import matplotlib.pyplot as plt
steps = L.runs[0].thermo.Step
ke     = L.runs[0].thermo.Ke
plt.plot(steps, ke)
```

## Error handling with PyLammps

Compiling the shared library with C++ exception support provides a better error handling experience. Without exceptions the LAMMPS code will terminate the current Python process with an error message. C++ exceptions allow capturing them on the C++ side and rethrowing them on the Python side. This way you can handle LAMMPS errors through the Python exception handling mechanism.

**Warning:** Capturing a LAMMPS exception in Python can still mean that the current LAMMPS process is in an illegal state and must be terminated. It is advised to save your data and terminate the Python instance as quickly as possible.

## Using PyLammps in IPython notebooks and Jupyter

If the LAMMPS Python package is installed for the same Python interpreter as IPython, you can use PyLammps directly inside of an IPython notebook inside of Jupyter. Jupyter is a powerful integrated development environment (IDE) for many dynamic languages like Python, Julia and others, which operates inside of any web browser. Besides auto-completion and syntax highlighting it allows you to create formatted documents using Markup, mathematical formulas, graphics and animations intermixed with executable Python code. It is a great format for tutorials and showcasing your latest research.

To launch an instance of Jupyter simply run the following command inside your Python environment (this assumes you followed the Quick Start instructions):

```
jupyter notebook
```

## IPyLammps Examples

Examples of IPython notebooks can be found in the `python/examples/pylammps` sub-directory. To open these notebooks launch `jupyter notebook` inside this directory and navigate to one of them. If you compiled and installed a LAMMPS shared library with exceptions, PNG, JPEG and FFMPEG support you should be able to rerun all of these notebooks.

## Validating a dihedral potential

This example showcases how an IPython Notebook can be used to compare a simple LAMMPS simulation of a harmonic dihedral potential to its analytical solution. Four atoms are placed in the simulation and the dihedral potential is applied on them using a datafile. Then one of the atoms is rotated along the central axis by setting its position from Python, which changes the dihedral angle.

```
phi = [d * math.pi / 180 for d in range(360)]
pos = [(1.0, math.cos(p), math.sin(p)) for p in phi]
pe = []
```

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```

for p in pos:
    L.atoms[3].position = p
    L.run(0)
    pe.append(L.eval("pe"))

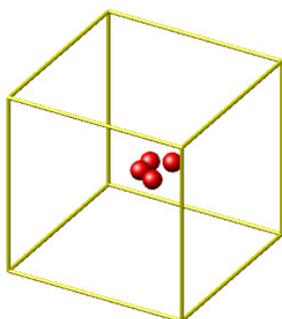
```

By evaluating the potential energy for each position we can verify that trajectory with the analytical formula. To compare both solutions, we plot both trajectories over each other using matplotlib, which embeds the generated plot inside the IPython notebook.

```

In [11]: L.image(zoom=1.0)
Out[11]:

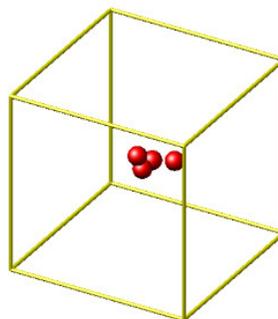
```



```

In [14]: L.image(zoom=1.0)
Out[14]:

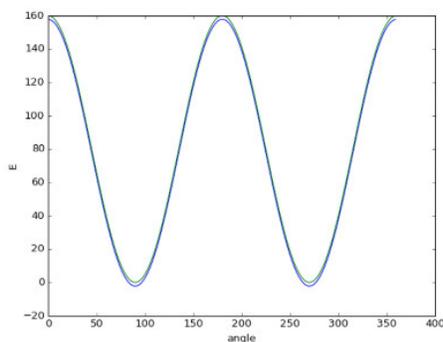
```



```

In [27]: plt.plot(range(360), pe, range(360), E_analytical)
plt.xlabel('angle')
plt.ylabel('E')

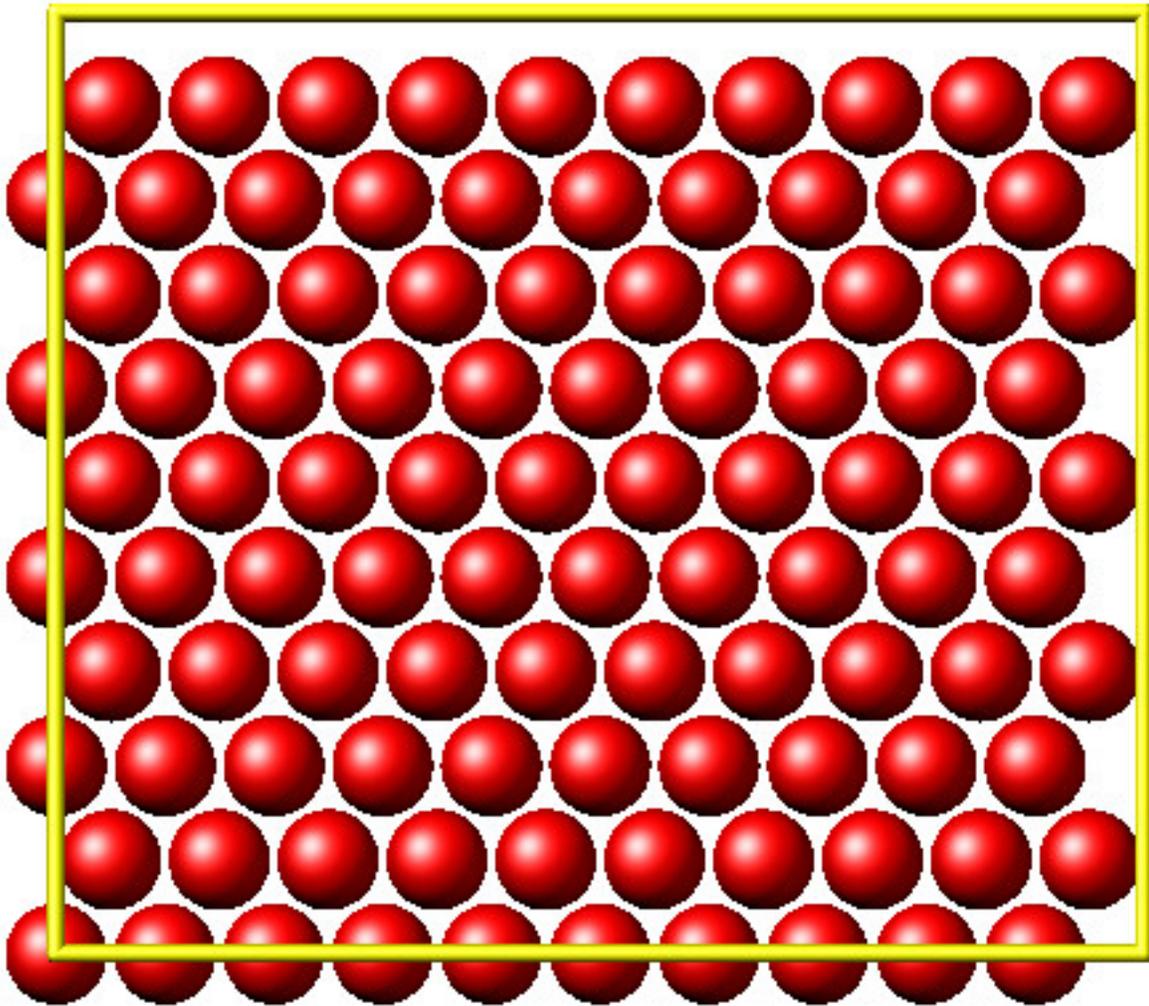
```



## Running a Monte Carlo relaxation

This second example shows how to use PyLammps to create a 2D Monte Carlo Relaxation simulation, computing and plotting energy terms and even embedding video output.

Initially, a 2D system is created in a state with minimal energy.

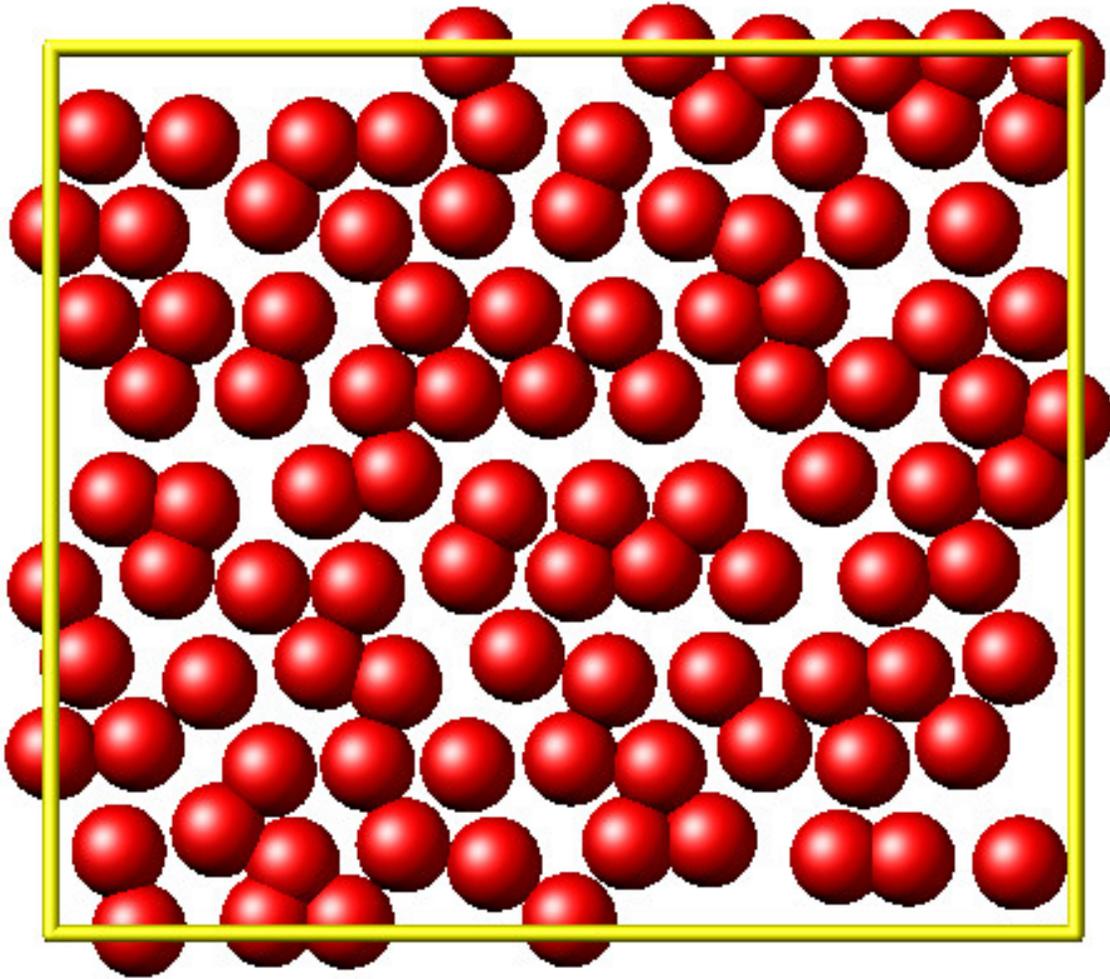


It is then disordered by moving each atom by a random delta.

```
random.seed(27848)
deltaperturb = 0.2

for i in range(L.system.natoms):
    x, y = L.atoms[i].position
    dx = deltaperturb * random.uniform(-1, 1)
    dy = deltaperturb * random.uniform(-1, 1)
    L.atoms[i].position = (x+dx, y+dy)

L.run(0)
```



Finally, the Monte Carlo algorithm is implemented in Python. It continuously moves random atoms by a random delta and only accepts certain moves.

```
estart = L.eval("pe")
elast = estart

naccept = 0
energies = [estart]

niterations = 3000
deltamove = 0.1
kT = 0.05

natoms = L.system.natoms

for i in range(niterations):
    iatom = random.randrange(0, natoms)
    current_atom = L.atoms[iatom]
```

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```
x0, y0 = current_atom.position

dx = deltamove \* random.uniform(-1, 1)
dy = deltamove \* random.uniform(-1, 1)

current_atom.position = (x0+dx, y0+dy)

L.run(1, "pre no post no")

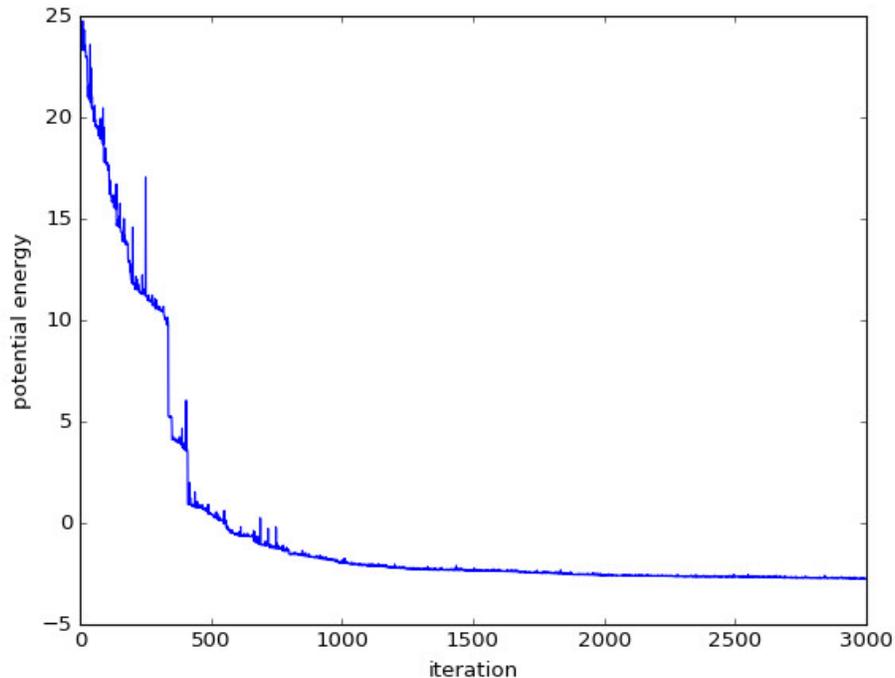
e = L.eval("pe")
energies.append(e)

if e <= elast:
    naccept += 1
    elast = e
elif random.random() <= math.exp(natoms\*(elast-e)/kT):
    naccept += 1
    elast = e
else:
    current_atom.position = (x0, y0)
```

The energies of each iteration are collected in a Python list and finally plotted using matplotlib.

```
In [20]: plt.xlabel('iteration')
plt.ylabel('potential energy')
plt.plot(energies)
```

Figure 1



```
Out[20]: [<matplotlib.lines.Line2D at 0x7f26a40123c8>]
```

The IPython notebook also shows how to use dump commands and embed video files inside of the IPython notebook.

### Using PyLammps and mpi4py (Experimental)

PyLammps can be run in parallel using mpi4py. This python package can be installed using

```
pip install mpi4py
```

The following is a short example which reads in an existing LAMMPS input file and executes it in parallel. You can find `in.melt` in the `examples/melt` folder.

```
from mpi4py import MPI
from lammps import PyLammps

L = PyLammps()
L.file("in.melt")

if MPI.COMM_WORLD.rank == 0:
    print("Potential energy: ", L.eval("pe"))
```

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```
MPI.Finalize()
```

To run this script (melt.py) in parallel using 4 MPI processes we invoke the following mpirun command:

```
mpirun -np 4 python melt.py
```

**Warning:** Any command must be executed by all MPI processes. However, evaluations and querying the system state is only available on rank 0.

## Feedback and Contributing

If you find this Python interface useful, please feel free to provide feedback and ideas on how to improve it to Richard Berger ([richard.berger@outlook.com](mailto:richard.berger@outlook.com)). We also want to encourage people to write tutorial style IPython notebooks showcasing LAMMPS usage and maybe their latest research results.

## 8.6.4 Using LAMMPS on Windows 10 with WSL

written by Richard Berger

---

It's always been tricky for us to have LAMMPS users and developers work on Windows. We primarily develop LAMMPS to run on Linux clusters. To teach LAMMPS in workshop settings, we had to redirect Windows users to Linux Virtual Machines such as VirtualBox or Unix-like compilation with Cygwin.

With the latest updates in Windows 10 (Version 2004, Build 19041 or higher), Microsoft has added a new way to work on Linux-based code. The Windows Subsystem for Linux (WSL). With WSL Version 2, you now get a Linux Virtual Machine that transparently integrates into Windows. All you need is to ensure you have the latest Windows updates installed and enable this new feature. Linux VMs are then easily installed using the Microsoft Store.

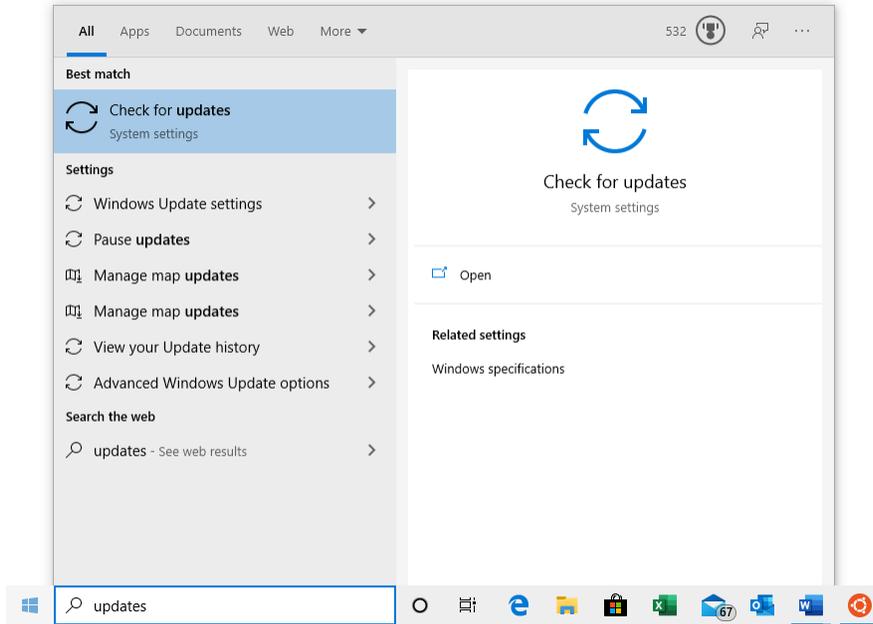
In this tutorial, I'll show you how to set up and compile LAMMPS for both serial and MPI usage in WSL2.

### Installation

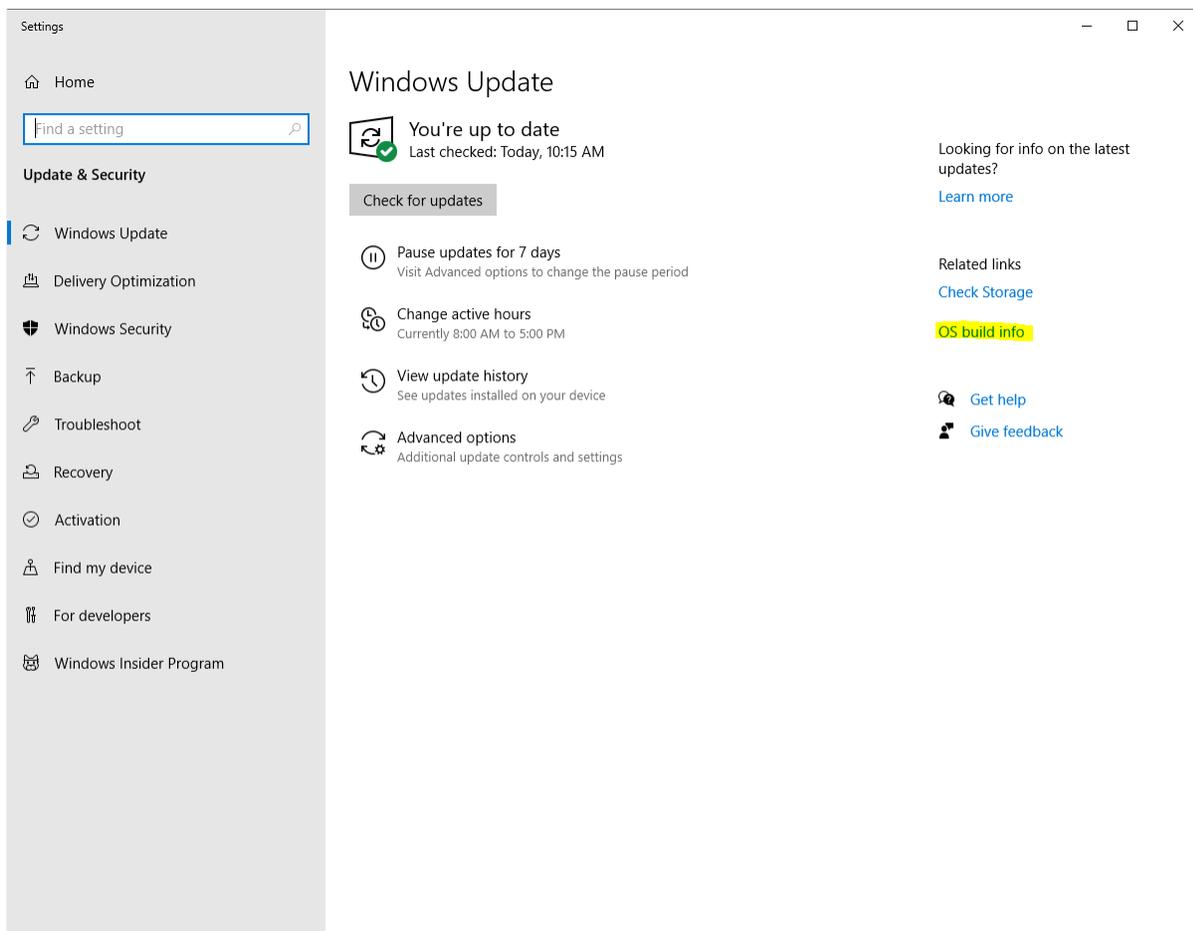
#### Upgrade to the latest Windows 10

Type "Updates" in Windows Start and select "Check for Updates".



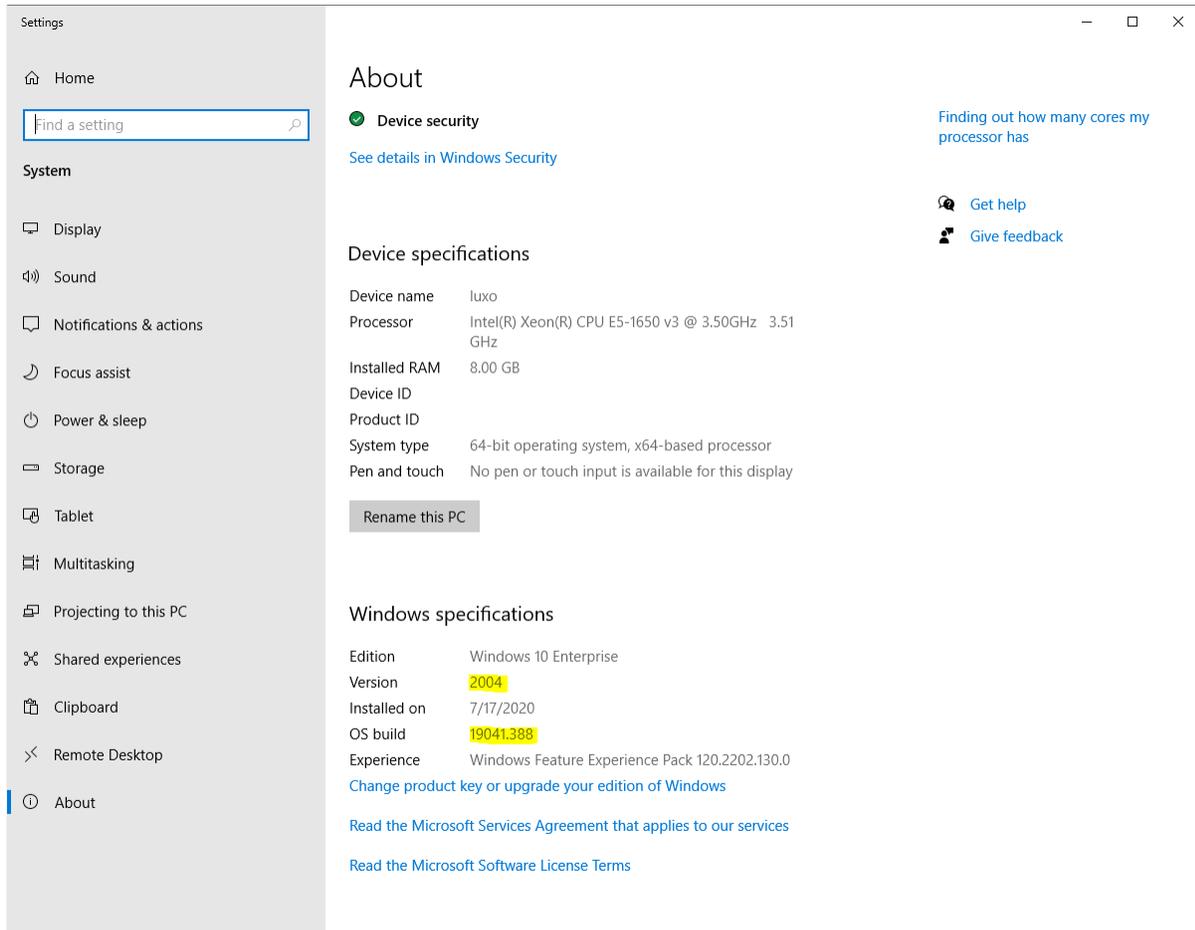


Install all pending updates and reboot your system as many times as necessary. Continue until your Windows installation is updated.



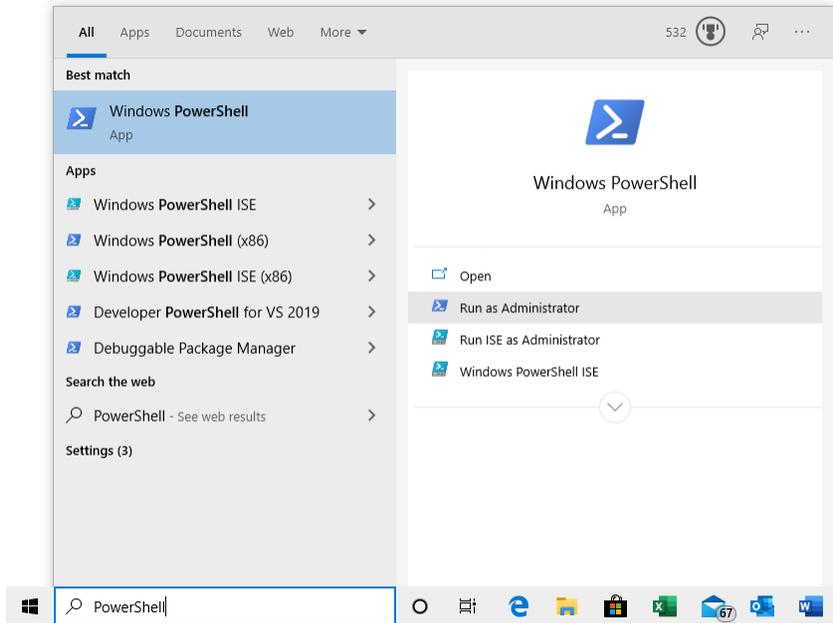
Verify your system has at least **version 2004 and build 19041 or later**. You can find this information by clicking on

“OS build info”.



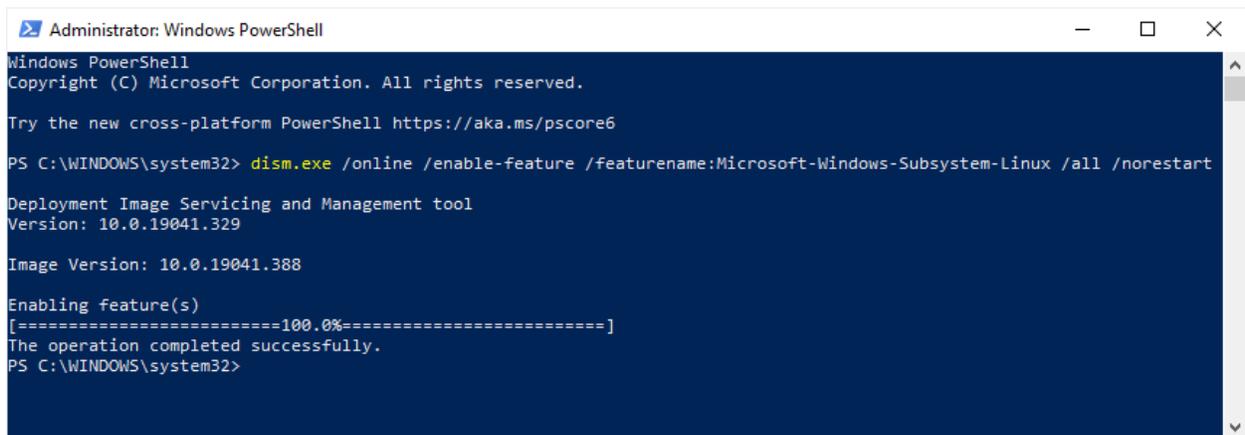
## Enable WSL

Next, we must install two additional Windows features to enable WSL support. Open a PowerShell window as an administrator. Type “PowerShell” in Windows Start and select “Run as Administrator”.



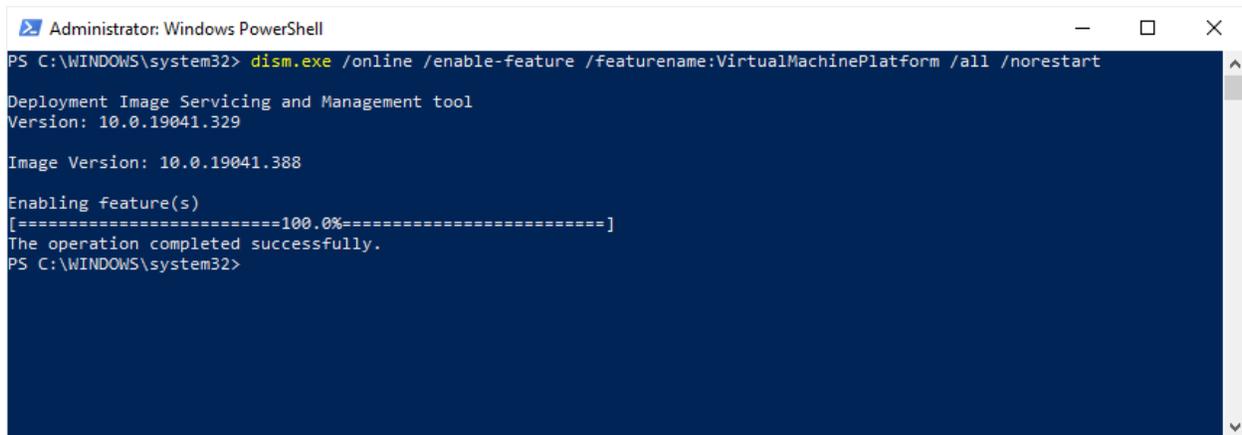
Windows will ask you for administrator access. After you accept a new command line window will appear. Type in the following command to install WSL:

```
dism.exe /online /enable-feature /featurename:Microsoft-Windows-Subsystem-Linux /all /
↵norestart
```



Next, enable the VirtualMachinePlatform feature using the following command:

```
dism.exe /online /enable-feature /featurename:VirtualMachinePlatform /all /norestart
```



```
Administrator: Windows PowerShell
PS C:\WINDOWS\system32> dism.exe /online /enable-feature /featurename:VirtualMachinePlatform /all /norestart

Deployment Image Servicing and Management tool
Version: 10.0.19041.329

Image Version: 10.0.19041.388

Enabling feature(s)
[=====100.0%=====]
The operation completed successfully.
PS C:\WINDOWS\system32>
```

Finally, reboot your system.

### Update WSL kernel component

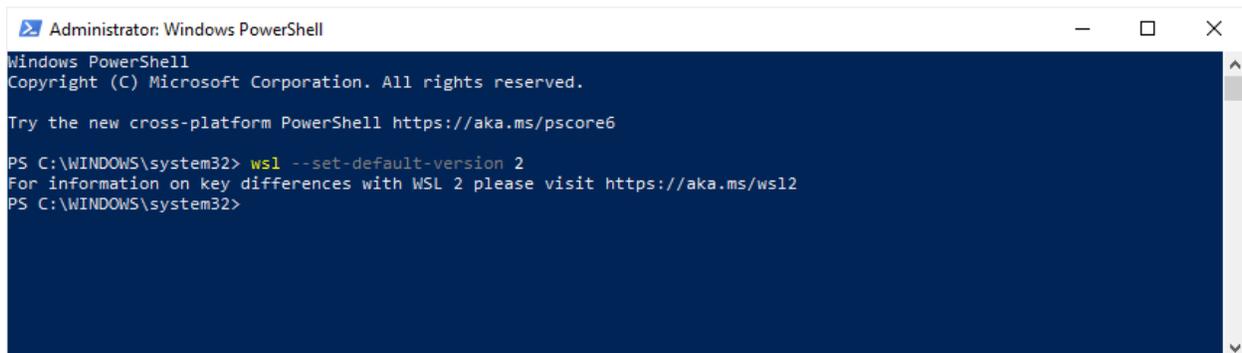
Download and install the WSL Kernel Component Update. Afterwards, reboot your system.

### Set WSL2 as default

Again, open PowerShell as administrator and run the following command:

```
wsl --set-default-version 2
```

This command ensures that all future Linux installations will use WSL version 2.



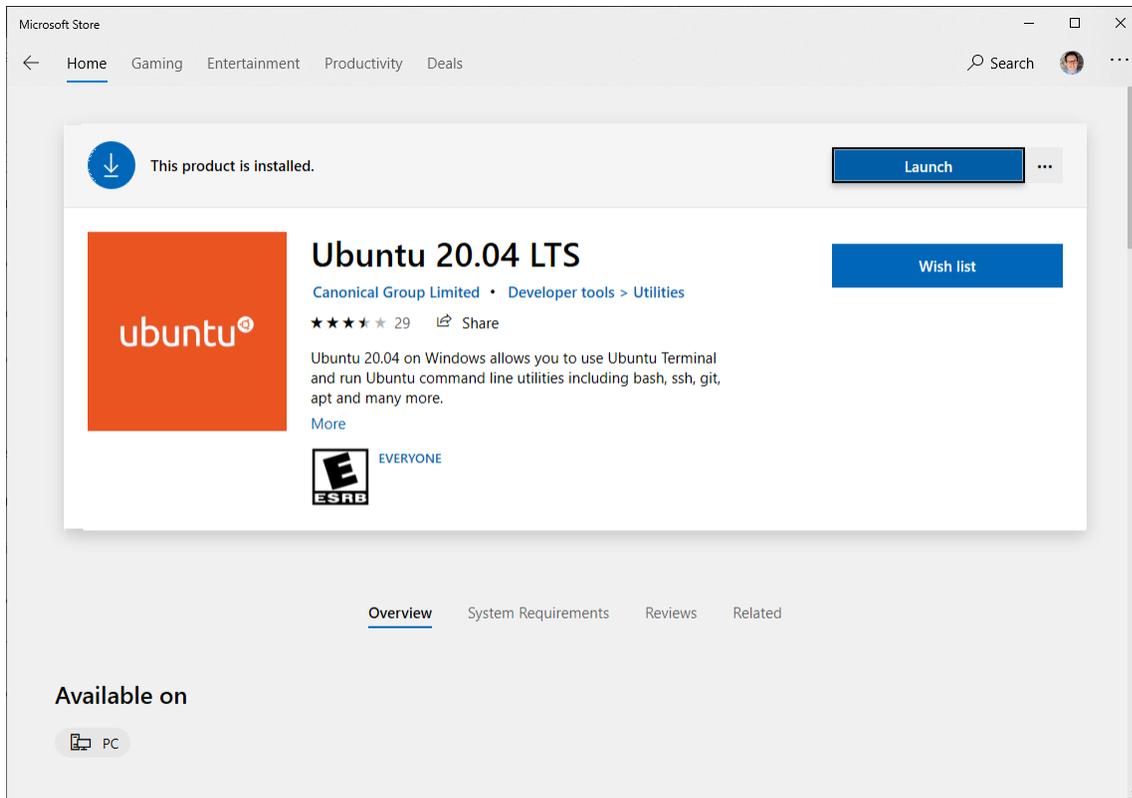
```
Administrator: Windows PowerShell
Windows PowerShell
Copyright (C) Microsoft Corporation. All rights reserved.

Try the new cross-platform PowerShell https://aka.ms/pscore6

PS C:\WINDOWS\system32> wsl --set-default-version 2
For information on key differences with WSL 2 please visit https://aka.ms/wsl2
PS C:\WINDOWS\system32>
```

### Install a Linux Distribution

Next, we need to install a Linux distribution via the Microsoft Store. Install [Ubuntu 20.04 LTS](#). Once installed, you can launch it like any other application from the Start Menu.



## Initial Setup

The first time you launch the Ubuntu Linux console, it will prompt you for a UNIX username and password. You will need this password to perform sudo commands later. Once completed, your Linux shell is ready for use. All your actions and commands will run as the Linux user you specified.

```
richard@luxo: ~
Retype new password:
passwd: password updated successfully
Installation successful!
To run a command as administrator (user "root"), use "sudo <command>".
See "man sudo_root" for details.

Welcome to Ubuntu 20.04 LTS (GNU/Linux 4.4.0-19041-Microsoft x86_64)

 * Documentation:  https://help.ubuntu.com
 * Management:    https://landscape.canonical.com
 * Support:       https://ubuntu.com/advantage

System information as of Fri Jul 17 13:51:44 EDT 2020

System load:  0.52   Processes:      7
Usage of /home: unknown   Users logged in:  0
Memory usage: 53%    IPv4 address for eth0: 10.0.2.15
Swap usage:   0%

0 updates can be installed immediately.
0 of these updates are security updates.

The list of available updates is more than a week old.
To check for new updates run: sudo apt update

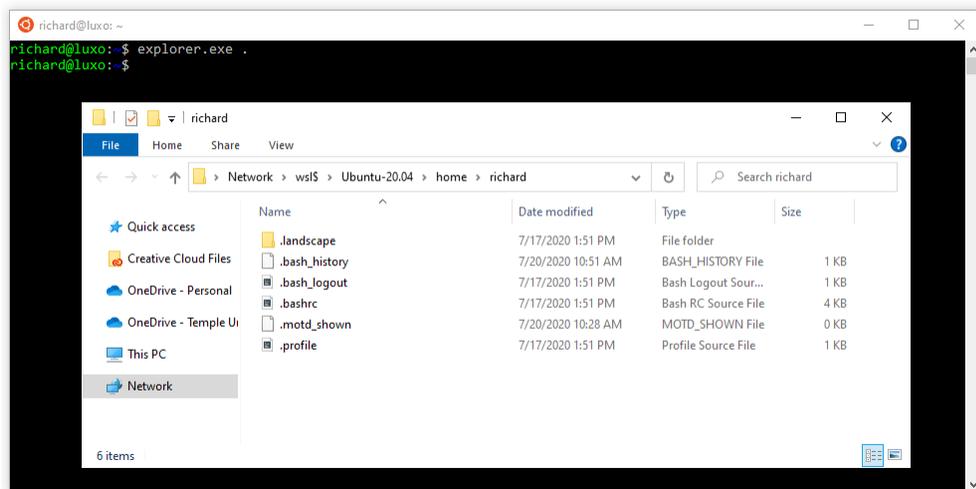
This message is shown once once a day. To disable it please create the
/home/richard/.hushlogin file.
richard@luxo: ~$
```

## Windows Explorer / WSL integration

Your Linux installation will have its own Linux filesystem, which contains the Ubuntu files. Your Linux user will have a regular Linux home directory in `/home/<USERNAME>`. This directory is different from your Windows User directory. Windows and Linux filesystems are connected through WSL.

All hard drives in Windows are accessible in the `/mnt` directory in Linux. E.g., WSL maps the C hard drive to the `/mnt/c` directory. That means you can access your Windows User directory in `/mnt/c/Users/<WINDOWS_USERNAME>`.

The Windows Explorer can also access the Linux filesystem. To illustrate this integration, open an Ubuntu console and navigate to a directory of your choice. To view this location in Windows Explorer, use the `explorer.exe .` command (do not forget the final dot!).



## Compiling LAMMPS

You now have a fully functioning Ubuntu installation and can follow most guides to install LAMMPS on a Linux system. Here are some of the essential steps to follow:

### Install prerequisite packages

Before we can begin, we need to download the necessary compiler toolchain and libraries to compile LAMMPS. In our Ubuntu-based Linux installation, we will use the `apt` package manager to install additional packages.

First, upgrade all existing packages using `apt update` and `apt upgrade`.

```
sudo apt update
sudo apt upgrade -y
```

Next, install the following packages with `apt install`:

```
sudo apt install -y cmake build-essential ccache gfortran openmpi-bin libopenmpi-dev \
libfftw3-dev libjpeg-dev libpng-dev python3-dev python3-pip \
python3-virtualenv libblas-dev liblapack-dev libhdf5-serial-dev \
hdf5-tools
```

## Download LAMMPS

Obtain a copy of the LAMMPS source code and go into it using the `cd` command.

### Option 1: Download a LAMMPS tarball using `wget`

```
wget https://github.com/lammps/lammps/archive/stable_3Mar2020.tar.gz
tar xvzf stable_3Mar2020.tar.gz
cd lammps
```

### Option 2: Download a LAMMPS development version from GitHub

```
git clone --depth=1 https://github.com/lammps/lammps.git
cd lammps
```

## Configure and Compile LAMMPS with CMake

A beginner-friendly way to compile LAMMPS is to use CMake. Create a build directory to compile LAMMPS and move into it. This directory will store the build configuration and any binaries generated during compilation.

```
mkdir build
cd build
```

There are countless ways to compile LAMMPS. It is beyond the scope of this tutorial. If you want to find out more about what can be enabled, please consult the extensive [documentation](#).

To compile a minimal version of LAMMPS, we're going to use a preset. Presets are a way to specify a collection of CMake options using a file.

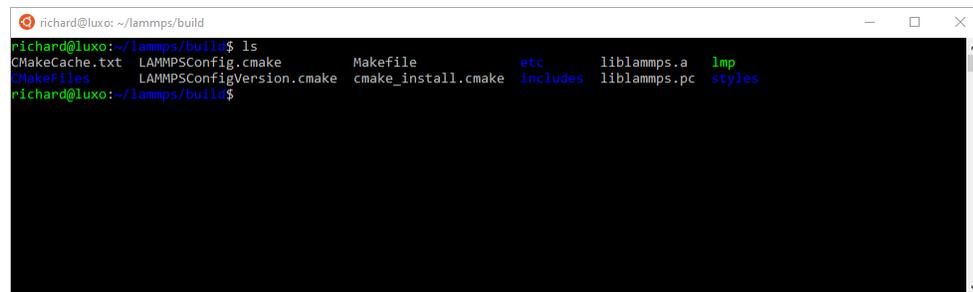
```
cmake ../cmake/presets/basic.cmake ../cmake
```

This command configures the build and generates the necessary Makefiles. To compile the binary, run the `make` command.

```
make -j 4
```

The `-j` option specifies how many parallel processes will perform the compilation. This option can significantly speed up compilation times. Use a number that corresponds to the number of processors in your system.

After the compilation completes successfully, you will have an executable called `lmp` in the `build` directory.



```
richard@luxo: ~/lammps/build
richard@luxo:~/lammps/build$ ls
CMakeCache.txt  LAMMPSConfig.cmake  Makefile          etc               liblammps.a  lmp
CMakeFiles     LAMMPSConfigVersion.cmake  cmake_install.cmake  includes         liblammps.pc  styles
richard@luxo:~/lammps/build$
```

Please take note of the absolute path of your build directory. You will need to know the location to execute the LAMMPS binary later.

One way of getting the absolute path of the current directory is through the \$PWD variable:

```
# prints out the current value of the PWD variable
echo $PWD
```

Let us save this value in a temporary variable LAMMPS\_BUILD\_DIR for future use:

```
LAMMPS_BUILD_DIR=$PWD
```

The full path of the LAMMPS binary then is \$LAMMPS\_BUILD\_DIR/lmp.

---

### Running an example script

Now that we have a LAMMPS binary, we will run a script from the examples folder.

Switch into the examples/melt folder:

```
cd ../examples/melt
```

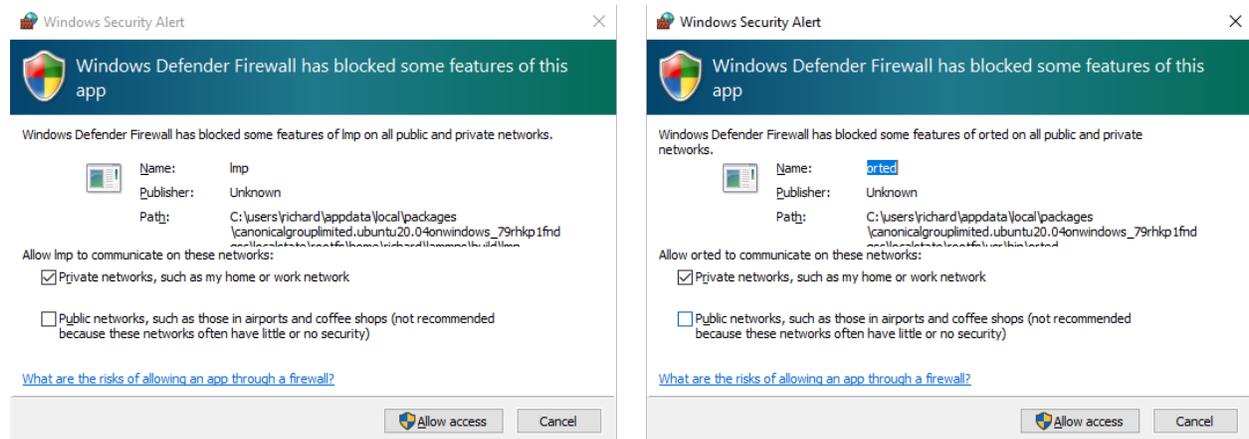
To run this example in serial, use the following command line:

```
$LAMMPS_BUILD_DIR/lmp -in in.melt
```

To run the same script in parallel using MPI with 4 processes, do the following:

```
mpirun -np 4 $LAMMPS_BUILD_DIR/lmp -in in.melt
```

If you run LAMMPS for the first time, the Windows Firewall might prompt you to confirm access. LAMMPS is accessing the network stack to enable parallel computation. Allow the access.



In either serial or MPI case, LAMMPS executes and will output something similar to this:

```
LAMMPS (30 Jun 2020)
...
...
...
```

(continues on next page)



(continued from previous page)

```
Total # of neighbors = 151513
Ave neighs/atom = 37.878250
Neighbor list builds = 12
Dangerous builds not checked
Total wall time: 0:00:00
```

**Congratulations! You've successfully compiled and executed LAMMPS on WSL!**

### Final steps

It is cumbersome to always specify the path of your LAMMPS binary. You can avoid this by adding the absolute path of your build directory to your PATH environment variable.

```
export PATH=$LAMMPS_BUILD_DIR:$PATH
```

You can then run LAMMPS input scripts like this:

```
lmp -in in.melt
```

or

```
mpirun -np 4 lmp -in in.melt
```

**Note:** The value of this PATH variable will disappear once you close your console window. To persist this setting edit the \$HOME/.bashrc file using your favorite text editor and add this line:

```
export PATH=/full/path/to/your/lammps/build:$PATH
```

**Example:** If the LAMMPS executable *lmp* has the following absolute path:

```
/home/<USERNAME>/lammps/build/lmp
```

the PATH variable should be:

```
export PATH=/home/<USERNAME>/lammps/build:$PATH
```

Once set up, all your Ubuntu consoles will always have access to your *lmp* binary without having to specify its location.

### Conclusion

I hope this gives you good overview on how to start compiling and running LAMMPS on Windows. WSL makes preparing and running scripts on Windows a much better experience.

If you are completely new to Linux, I highly recommend investing some time in studying Linux online tutorials. E.g., tutorials about Bash Shell and Basic Unix commands (e.g., [Linux Journey](#)). Acquiring these skills will make you much more productive in this environment.

**See also:**

- [Windows Subsystem for Linux Documentation](#)



## EXAMPLE SCRIPTS

The LAMMPS distribution includes an examples sub-directory with many sample problems. Many are 2d models that run quickly and are straightforward to visualize, requiring at most a couple of minutes to run on a desktop machine. Each problem has an input script (in.\*) and produces a log file (log.\*) when it runs. Some use a data file (data.\*) of initial coordinates as additional input. A few sample log file run on different machines and different numbers of processors are included in the directories to compare your answers to. E.g. a log file like log.date.crack.foo.P means the “crack” example was run on P processors of machine “foo” on that date (i.e. with that version of LAMMPS).

Many of the input files have commented-out lines for creating dump files and image files.

If you uncomment the *dump* command in the input script, a text dump file will be produced, which can be animated by various [visualization programs](#).

If you uncomment the *dump image* command in the input script, and assuming you have built LAMMPS with a JPG library, JPG snapshot images will be produced when the simulation runs. They can be quickly post-processed into a movie using commands described on the [dump image](#) doc page.

Animations of many of the examples can be viewed on the Movies section of the [LAMMPS website](#).

There are two kinds of sub-directories in the examples folder. Lower case named directories contain one or a few simple, quick-to-run problems. Upper case named directories contain up to several complex scripts that illustrate a particular kind of simulation method or model. Some of these run for longer times, e.g. to measure a particular quantity.

Lists of both kinds of directories are given below.

---

### 9.1 Lowercase directories

accelerate	run with various acceleration options (OpenMP, GPU, Phi)
airebo	polyethylene with AIREBO potential
atm	Axilrod-Teller-Muto potential example
balance	dynamic load balancing, 2d system
body	body particles, 2d system
bpm	BPM simulations of pouring elastic grains and plate impact
cmap	CMAF 5-body contributions to CHARMM force field
colloid	big colloid particles in a small particle solvent, 2d system
comb	models using the COMB potential
controller	use of fix controller as a thermostat
coreshell	core/shell model using CORESHELL package
crack	crack propagation in a 2d solid
deposit	deposit atoms and molecules on a surface

continues on next page

Table 1 – continued from previous page

dipole	point dipolar particles, 2d system
dreiding	methanol via Dreiding FF
eim	NaCl using the EIM potential
ellipse	ellipsoidal particles in spherical solvent, 2d system
flow	Couette and Poiseuille flow in a 2d channel
friction	frictional contact of spherical asperities between 2d surfaces
mc	Monte Carlo features via fix gcmc, widom and other commands
granregion	use of fix wall/region/gran as boundary on granular particles
hugoniostat	Hugoniostat shock dynamics
hyper	global and local hyperdynamics of diffusion on Pt surface
indent	spherical indenter into a 2d solid
kim	use of potentials from the <a href="#">OpenKIM Repository</a>
latte	examples for using fix latte for DFTB via the LATTE library
mdi	use of the MDI package and MolSSI MDI code coupling library
meam	MEAM test for SiC and shear (same as shear examples)
melt	rapid melt of 3d LJ system
micelle	self-assembly of small lipid-like molecules into 2d bilayers
min	energy minimization of 2d LJ melt
mscg	parameterize a multi-scale coarse-graining (MSCG) model
msst	MSST shock dynamics
multi	multi neighboring for systems with large interaction disparities
nb3b	use of non-bonded 3-body harmonic pair style
neb	nudged elastic band (NEB) calculation for barrier finding
nemd	non-equilibrium MD of 2d sheared system
obstacle	flow around two voids in a 2d channel
peptide	dynamics of a small solvated peptide chain (5-mer)
peri	Peridynamic model of cylinder impacted by indenter
pour	pouring of granular particles into a 3d box, then chute flow
prd	parallel replica dynamics of vacancy diffusion in bulk Si
python	using embedded Python in a LAMMPS input script
qeq	use of the QEQ package for charge equilibration
rdf-adf	computing radial and angle distribution functions for water
reax	RDX and TATB models using the ReaxFF
rerun	use of rerun and read_dump commands
rigid	rigid bodies modeled as independent or coupled
shear	sideways shear applied to 2d solid, with and without a void
snap	NVE dynamics for BCC tantalum crystal using SNAP potential
srd	stochastic rotation dynamics (SRD) particles as solvent
streitz	use of Streitz/Mintmire potential with charge equilibration
tad	temperature-accelerated dynamics of vacancy diffusion in bulk Si
threebody	regression test input for a variety of manybody potentials
tracker	track interactions in LJ melt
vashishta	use of the Vashishta potential
voronoi	Voronoi tessellation via compute voronoi/atom command

Here is how you can run and visualize one of the sample problems:

```
cd indent
cp ../../src/lmp_linux .           # copy LAMMPS executable to this dir
lmp_linux -in in.indent           # run the problem
```

Running the simulation produces the files *dump.indent* and *log.lammps*. You can visualize the dump file of snapshots

with a variety of third-party tools highlighted on the [Visualization](#) page of the LAMMPS website.

If you uncomment the *dump image* line(s) in the input script a series of JPG images will be produced by the run (assuming you built LAMMPS with JPG support; see the [Build\\_settings](#) page for details). These can be viewed individually or turned into a movie or animated by tools like ImageMagick or QuickTime or various Windows-based tools. See the [dump image](#) page for more details. E.g. this Imagemagick command would create a GIF file suitable for viewing in a browser.

```
% convert -loop 1 *.jpg foo.gif
```

## 9.2 Uppercase directories

ASPHERE	various aspherical particle models, using ellipsoids, rigid bodies, line/triangle particles, etc
COUPLE	examples of how to use LAMMPS as a library
DIFFUSE	compute diffusion coefficients via several methods
ELASTIC	compute elastic constants at zero temperature
ELASTIC_T	compute elastic constants at finite temperature
HEAT	compute thermal conductivity for LJ and water via fix ehex
KAPPA	compute thermal conductivity via several methods
MC-LOOP	using LAMMPS in a Monte Carlo mode to relax the energy of a system in a input script loop
PACKAGES	examples for specific packages and contributed commands
SPIN	examples for features of the SPIN package
UNITS	examples that run the same simulation in lj, real, metal units
VISCOSITY	compute viscosity via several methods

Nearly all of these directories have README files which give more details on how to understand and use their contents.

The PACKAGES directory has a large number of sub-directories which correspond by name to specific packages. They contain scripts that illustrate how to use the command(s) provided in those packages. Many of the sub-directories have their own README files which give further instructions. See the [Packages\\_details](#) doc page for more info on specific packages.



## AUXILIARY TOOLS

LAMMPS is designed to be a computational kernel for performing molecular dynamics computations. Additional pre- and post-processing steps are often necessary to setup and analyze a simulation. A list of such tools can be found on the [LAMMPS webpage](#) at these links:

- [Pre/Post processing](#)
- [External LAMMPS packages & tools](#)
- [Pizza.py toolkit](#)

The last link for [Pizza.py](#) is a Python-based tool developed at Sandia which provides tools for doing setup, analysis, plotting, and visualization for LAMMPS simulations.

Additional tools included in the LAMMPS distribution are described on this page.

Note that many users write their own setup or analysis tools or use other existing codes and convert their output to a LAMMPS input format or vice versa. The tools listed here are included in the LAMMPS distribution as examples of auxiliary tools. Some of them are not actively supported by the LAMMPS developers, as they were contributed by LAMMPS users. If you have problems using them, we can direct you to the authors.

The source code for each of these codes is in the tools sub-directory of the LAMMPS distribution. There is a Makefile (which you may need to edit for your platform) which will build several of the tools which reside in that directory. Most of them are larger packages in their own sub-directories with their own Makefiles and/or README files.

---

### 10.1 Pre-processing tools

<i>amber2lmp</i>	<i>ch2lmp</i>	<i>chain</i>	<i>createatoms</i>	<i>drude</i>	<i>eam database</i>
<i>eam generate</i>	<i>eff</i>	<i>ipp</i>	<i>micelle2d</i>	<i>moltemplate</i>	<i>msi2lmp</i>
<i>polybond</i>	<i>stl_bin2txt</i>				

### 10.2 Post-processing tools

<i>amber2lmp</i>	<i>binary2txt</i>	<i>ch2lmp</i>	<i>colvars</i>	<i>eff</i>	<i>fep</i>
<i>lmp2arc</i>	<i>lmp2cfg</i>	<i>matlab</i>	<i>phonon</i>	<i>pymol_asphere</i>	<i>python</i>
<i>replica</i>	<i>smd</i>	<i>spin</i>	<i>xmgrace</i>		

## 10.3 Miscellaneous tools

---

<i>LAMMPS coding stan- dards</i>	<i>emacs</i>	<i>i-pi</i>	<i>kate</i>	<i>LAMMPS shell</i>	<i>LAMMPS magic patterns for file(1)</i>
<i>Offline build tool</i>	<i>singular- ity/apptainer</i>	<i>SWIG inter- face</i>	<i>vim</i>		

---

## 10.4 Tool descriptions

### 10.4.1 amber2Imp tool

The `amber2Imp` sub-directory contains two Python scripts for converting files back-and-forth between the AMBER MD code and LAMMPS. See the README file in `amber2Imp` for more information.

These tools were written by Keir Novik while he was at Queen Mary University of London. Keir is no longer there and cannot support these tools which are out-of-date with respect to the current LAMMPS version (and maybe with respect to AMBER as well). Since we don't use these tools at Sandia, you will need to experiment with them and make necessary modifications yourself.

---

### 10.4.2 binary2txt tool

The file `binary2txt.cpp` converts one or more binary LAMMPS dump file into ASCII text files. The syntax for running the tool is

```
binary2txt file1 file2 ...
```

which creates `file1.txt`, `file2.txt`, etc. This tool must be compiled on a platform that can read the binary file created by a LAMMPS run, since binary files are not compatible across all platforms.

---

### 10.4.3 ch2Imp tool

The `ch2Imp` sub-directory contains tools for converting files back-and-forth between the CHARMM MD code and LAMMPS.

They are intended to make it easy to use CHARMM as a builder and as a post-processor for LAMMPS. Using `charmm2lammps.pl`, you can convert a PDB file with associated CHARMM info, including CHARMM force field data, into its LAMMPS equivalent. Support for the CMAP correction of CHARMM22 and later is available as an option. This tool can also add solvent water molecules and Na<sup>+</sup> or Cl<sup>-</sup> ions to the system. Using `lammps2pdb.pl` you can convert LAMMPS atom dumps into PDB files.

See the README file in the `ch2Imp` sub-directory for more information.

These tools were created by Pieter in't Veld (`pjintve at sandia.gov`) and Paul Crozier (`pscrozi at sandia.gov`) at Sandia.

CMAP support added and tested by Xiaohu Hu (`hux2 at ornl.gov`) and Robert A. Latour (`latourr at clemson.edu`), David Hyde-Volpe, and Tigran Abramyan, (Clemson University) and Chris Lorenz (`chris.lorenz at kcl.ac.uk`), King's College London.

---



### 10.4.4 chain tool

The file `chain.f90` creates a LAMMPS data file containing bead-spring polymer chains and/or monomer solvent atoms. It uses a text file containing chain definition parameters as an input. The created chains and solvent atoms can strongly overlap, so LAMMPS needs to run the system initially with a “soft” pair potential to un-overlap it. The syntax for running the tool is

```
chain < def.chain > data.file
```

See the `def.chain` or `def.chain.ab` files in the `tools` directory for examples of definition files. This tool was used to create the system for the *chain benchmark*.

### 10.4.5 LAMMPS coding standard

The `coding_standard` folder contains multiple python scripts to check for and apply some LAMMPS coding conventions. The following scripts are available:

```
permissions.py # detects if sources have executable permissions and scripts have not
whitespace.py # detects TAB characters and trailing whitespace
homepage.py   # detects outdated LAMMPS homepage URLs (pointing to sandia.gov instead
↳of lammps.org)
errordocs.py  # detects deprecated error docs in header files
```

The tools need to be given the main folder of the LAMMPS distribution or individual file names as argument and will by default check them and report any non-compliance. With the optional `-f` argument the corresponding script will try to change the non-compliant file(s) to match the conventions.

For convenience this scripts can also be invoked by the make file in the `src` folder with, *make check-whitespace* or *make fix-whitespace* to either detect or edit the files. Correspondingly for the other python scripts. *make check* will run all checks.

### 10.4.6 colvars tools

The `colvars` directory contains a collection of tools for post-processing data produced by the `colvars` collective variable library. To compile the tools, edit the makefile for your system and run “make”.

Please report problems and issues the `colvars` library and its tools at: <https://github.com/colvars/colvars/issues>

`abf_integrate`:

MC-based integration of multidimensional free energy gradient Version 20110511

```
./abf_integrate < filename > [-n < nsteps >] [-t < temp >] [-m [0|1] (metadynamics)] [-h
↳< hill_height >] [-f < variable_hill_factor >]
```

The LAMMPS interface to the `colvars` collective variable library, as well as these tools, were created by Axel Kohlmeyer ([akohlmey@gmail.com](mailto:akohlmey@gmail.com)) while at ICTP, Italy.

### 10.4.7 createatoms tool

The tools/createatoms directory contains a Fortran program called createAtoms.f which can generate a variety of interesting crystal structures and geometries and output the resulting list of atom coordinates in LAMMPS or other formats.

See the included Manual.pdf for details.

The tool is authored by Xiaowang Zhou (Sandia), xzhou at sandia.gov.

---

### 10.4.8 drude tool

The tools/drude directory contains a Python script called polarizer.py which can add Drude oscillators to a LAMMPS data file in the required format.

See the header of the polarizer.py file for details.

The tool is authored by Agilio Padua and Alain Dequidt: agilio.padua at ens-lyon.fr, alain.dequidt at uca.fr

---

### 10.4.9 eam database tool

The tools/eam\_database directory contains a Fortran and a Python program that will generate EAM alloy setfl potential files for any combination of the 17 elements: Cu, Ag, Au, Ni, Pd, Pt, Al, Pb, Fe, Mo, Ta, W, Mg, Co, Ti, Zr, Cr. The files can then be used with the *pair\_style eam/alloy* command.

The Fortran version of the tool was authored by Xiaowang Zhou (Sandia), xzhou at sandia.gov, with updates from Lucas Hale (NIST) lucas.hale at nist.gov and is based on his paper:

X. W. Zhou, R. A. Johnson, and H. N. G. Wadley, Phys. Rev. B, 69, 144113 (2004).

The parameters for Cr were taken from:

Lin Z B, Johnson R A and Zhigilei L V, Phys. Rev. B 77 214108 (2008).

The Python version of the tool was authored by Germain Clavier (TU Eindhoven) g.m.g.c.clavier at tue.nl or germain.clavier at gmail.com

---

**Note:** The parameters in the database are only optimized for individual elements. The mixed parameters for interactions between different elements generated by this tool are derived from simple mixing rules and are thus inferior to parameterizations that are specifically optimized for specific mixtures and combinations of elements.

---

### 10.4.10 eam generate tool

The tools/eam\_generate directory contains several one-file C programs that convert an analytic formula into a tabulated *embedded atom method (EAM)* setfl potential file. The potentials they produce are in the potentials directory, and can be used with the *pair\_style eam/alloy* command.

The source files and potentials were provided by Gerolf Ziegenhain (gerolf at ziegenhain.com).

---

### 10.4.11 eff tool

The tools/eff directory contains various scripts for generating structures and post-processing output for simulations using the electron force field (eFF).

These tools were provided by Andres Jaramillo-Botero at CalTech (ajaramil at wag.caltech.edu).

---

### 10.4.12 emacs tool

The tools/emacs directory contains an Emacs Lisp add-on file for GNU Emacs that enables a lammeps-mode for editing input scripts when using GNU Emacs, with various highlighting options set up.

These tools were provided by Aidan Thompson at Sandia (athomps at sandia.gov).

---

### 10.4.13 fep tool

The tools/fep directory contains Python scripts useful for post-processing results from performing free-energy perturbation simulations using the FEP package.

The scripts were contributed by Agilio Padua (ENS de Lyon), agilio.padua at ens-lyon.fr.

See README file in the tools/fep directory.

---

### 10.4.14 i-pi tool

The tools/i-pi directory contains a version of the i-PI package, with all the LAMMPS-unrelated files removed. It is provided so that it can be used with the *fix ipi* command to perform path-integral molecular dynamics (PIMD).

The i-PI package was created and is maintained by Michele Ceriotti, michele.ceriotti at gmail.com, to interface to a variety of molecular dynamics codes.

See the tools/i-pi/manual.pdf file for an overview of i-PI, and the *fix ipi* page for further details on running PIMD calculations with LAMMPS.

---

### 10.4.15 ipp tool

The tools/ipp directory contains a Perl script ipp which can be used to facilitate the creation of a complicated file (say, a lammeps input script or tools/createatoms input file) using a template file.

ipp was created and is maintained by Reese Jones (Sandia), rjones at sandia.gov.

See two examples in the tools/ipp directory. One of them is for the tools/createatoms tool's input file.

---

### 10.4.16 kate tool

The file in the tools/kate directory is an add-on to the Kate editor in the KDE suite that allow syntax highlighting of LAMMPS input scripts. See the README.txt file for details.

The file was provided by Alessandro Luigi Sellerio (alessandro.sellerio at ieni.cnr.it).

---

### 10.4.17 LAMMPS shell

New in version 9Oct2020.

#### Overview

The LAMMPS Shell, `lammops-shell` is a program that functions very similar to the regular LAMMPS executable but has several modifications and additions that make it more powerful for interactive sessions, i.e. where you type LAMMPS commands from the prompt instead of reading them from a file.

- It uses the readline and history libraries to provide command line editing and context aware TAB-expansion (details on that below).
- When processing an input file with the `'-in'` or `'-i'` flag from the command line, it does not exit at the end of that input file but stops at a prompt, so that additional commands can be issued
- Errors will not abort the shell but return to the prompt.
- It has additional commands aimed at interactive use (details below).
- Interrupting a calculation with CTRL-C will not terminate the session but rather enforce a timeout to cleanly stop an ongoing run (more info on timeouts is in the *timer command* documentation).

These enhancements make the LAMMPS shell an attractive choice for interactive LAMMPS sessions in graphical desktop environments (e.g. Gnome, KDE, Cinnamon, XFCE, Windows).

#### TAB-expansion

When writing commands interactively at the shell prompt, you can hit the TAB key at any time to try and complete the text. This completion is context aware and will expand any first word only to commands available in that executable.

- For style commands it will expand to available styles of the corresponding category (e.g. pair styles after a *pair\_style* command).
- For *compute*, *fix*, or *dump* it will also expand only to already defined groups for the group-ID keyword.
- For commands like *compute\_modify*, *fix\_modify*, or *dump\_modify* it will expand to known compute/fix/dump IDs only.
- When typing references to computes, fixes, or variables with a `"c_"`, `"f_"`, or `"v_"` prefix, respectively, then the expansion will be to known compute/fix IDs and variable names. Variable name expansion is also available for the `${name}` variable syntax.
- In all other cases TAB expansion will complete to names of files and directories.

## Command line editing and history

When typing commands, command line editing similar to what BASH provides is available. Thus it is possible to move around the currently line and perform various cut and insert and edit operations. Previous commands can be retrieved by scrolling up (and down) or searching (e.g. with CTRL-r).

Also history expansion through using the exclamation mark ‘!’ can be performed. Examples: ‘!’ will be replaced with the previous command, ‘!-2’ will repeat the command before that, ‘!30’ will be replaced with event number 30 in the command history list, and ‘!run’ with the last command line that started with “run”. Adding a “:p” to such a history expansion will result that the expansion is printed and added to the history list, but NOT executed. On exit the LAMMPS shell will write the history list to a file “.lammps\_history” in the current working directory. If such a file exists when the LAMMPS shell is launched it will be read to populate the history list.

This is realized via the readline library and can thus be customized with an `.inputrc` file in the home directory. For application specific customization, the LAMMPS shell uses the name “lammps-shell”. For more information about using and customizing an application using readline, please see the available documentation at: <http://www.gnu.org/s/readline/#Documentation>

## Additional commands

The following commands are added to the LAMMPS shell on top of the regular LAMMPS commands:

```

help (or ?)    print a brief help message
history        display the current command history list
clear_history  wipe out the current command history list
save_history <range> <file>
                write commands from the history to file.
                The range is given as <from>-<to>, where <from> and <to>
                may be empty. Example: save_history 100- in.recent
source <file>  read commands from file (same as "include")
pwd            print current working directory
cd <directory> change current working directory (same as pwd if no directory)
mem           print current and maximum memory usage
|<command>    execute <command> as a shell command and return to the command prompt
exit          exit the LAMMPS shell cleanly (unlike the "quit" command)

```

Please note that some known shell operations are implemented in the LAMMPS *shell command* in a platform neutral fashion, while using the ‘|’ character will always pass the following text to the operating system’s shell command.

## Compilation

Compilation of the LAMMPS shell can be enabled by setting the CMake variable `BUILD_LAMMPS_SHELL` to “on” or using the makefile in the `tools/lammps-shell` folder to compile after building LAMMPS using the conventional make procedure. The makefile will likely need customization depending on the features and settings used for compiling LAMMPS.

## Limitations

The LAMMPS shell was not designed for use with MPI parallelization via `mpirun` or `mpiexec` or `srun`.

## Readline customization

The behavior of the readline functionality can be customized in the `${HOME}/.inputrc` file. This can be used to alter the default settings or change the key-bindings. The LAMMPS Shell sets the application name `lammcs-shell`, so settings can be either applied globally or only for the LAMMPS shell by bracketing them between `$if lammcs-shell` and `$endif` like in the following example:

```
$if lammcs-shell
# disable "beep" or "screen flash"
set bell-style none
# bind the "Insert" key to toggle overwrite mode
"\e[2~": overwrite-mode
$endif
```

More details about this are in the [readline documentation](#).

## LAMMPS Shell tips and tricks

Below are some suggestions for how to use and customize the LAMMPS shell.

### Enable tilde expansion

Adding `set expand-tilde on` to `${HOME}/.inputrc` is recommended as this will change the filename expansion behavior to replace any text starting with “~” by the full path to the corresponding user’s home directory. While the expansion of filenames **will** happen on all arguments where the context is not known (e.g. `~/compile/lamm<TAB>` will expand to `~/compile/lammcs/`), it will not replace the tilde by default. But since LAMMPS does not do tilde expansion itself (unlike a shell), this will result in errors. Instead the tilde-expression should be expanded into a valid path, where the plain “~/” stands for the current user’s home directory and “~/someuser/” stands for “/home/someuser” or whatever the full path to that user’s home directory is.

### File extension association

Since the LAMMPS shell (unlike the regular LAMMPS executable) does not exit when an input file is passed on the command line with the “-in” or “-i” flag (the behavior is like for `python -i <filename>`), it makes the LAMMPS shell suitable for associating it with input files based on their filename extension (e.g. “.lmp”). Since `lammcs-shell` is a console application, you have to run it inside a terminal program with a command line like this:

```
xterm -title "LAMMPS Shell" -e /path/to/lammcs-shell -i in.file.lmp
```

## Use history to create an input file

When experimenting with commands to interactively to figure out a suitable choice of settings or simply the correct syntax, you may want to record part of your commands to a file for later use. This can be done with the `save_history` commands, which allows to selectively write a section of the command history to a file (Example: `save_history 25-30 in.run`). This file can be further edited (Example: `|vim in.run`) and then the file read back in and tried out (Example: `source in.run`). If the input also creates a system box, you first need to use the `clear_command` command.

### 10.4.18 Imp2arc tool

The `imp2arc` sub-directory contains a tool for converting LAMMPS output files to the format for Accelrys' Insight MD code (formerly MSI/Biosym and its Discover MD code). See the README file for more information.

This tool was written by John Carpenter (Cray), Michael Peachey (Cray), and Steve Lustig (Dupont). John is now at the Mayo Clinic (`jec` at `mayo.edu`), but still fields questions about the tool.

This tool was updated for the current LAMMPS C++ version by Jeff Greathouse at Sandia (`jagreat` at `sandia.gov`).

### 10.4.19 Imp2cfg tool

The `imp2cfg` sub-directory contains a tool for converting LAMMPS output files into a series of `*.cfg` files which can be read into the `AtomEye` visualizer. See the README file for more information.

This tool was written by Ara Kooser at Sandia (`askoose` at `sandia.gov`).

### 10.4.20 Magic patterns for the “file” command

New in version 10Mar2021.

The `file` magic contains patterns that are used by the `file` program available on most Unix-like operating systems which enables it to detect various LAMMPS files and print some useful information about them. To enable these patterns, append or copy the contents of the file `.magic` in your home directory or (as administrator) to `/etc/magic` (for a system-wide installation). Afterwards the `file` command should be able to detect most LAMMPS restarts, dump, data and log files. Examples:

```
$ file *.*
dihedral-quadratic.restart:  LAMMPS binary restart file (rev 2), Version 10 Mar 2021,
↳Little Endian
mol-pair-wf_cut.restart:    LAMMPS binary restart file (rev 2), Version 24 Dec 2020,
↳Little Endian
atom.bin:                   LAMMPS atom style binary dump (rev 2), Little Endian,
↳First time step: 445570
custom.bin:                 LAMMPS custom style binary dump (rev 2), Little Endian,
↳First time step: 100
bn1.lammpstrj:              LAMMPS text mode dump, First time step: 5000
data.fourmol:               LAMMPS data file written by LAMMPS
pnc.data:                   LAMMPS data file written by msi2lmp
```

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data.spce:	LAMMPS data file written by TopoTools
B.data:	LAMMPS data file written by OVITO
log.lammps:	LAMMPS log file written by version 10 Feb 2021

---

### 10.4.21 matlab tool

The matlab sub-directory contains several **MATLAB** scripts for post-processing LAMMPS output. The scripts include readers for log and dump files, a reader for EAM potential files, and a converter that reads LAMMPS dump files and produces CFG files that can be visualized with the **AtomEye** visualizer.

See the README.pdf file for more information.

These scripts were written by Arun Subramaniyan at Purdue Univ (asubrama at purdue.edu).

---

### 10.4.22 micelle2d tool

The file micelle2d.f creates a LAMMPS data file containing short lipid chains in a monomer solution. It uses a text file containing lipid definition parameters as an input. The created molecules and solvent atoms can strongly overlap, so LAMMPS needs to run the system initially with a “soft” pair potential to un-overlap it. The syntax for running the tool is

```
micelle2d < def.micelle2d > data.file
```

See the def.micelle2d file in the tools directory for an example of a definition file. This tool was used to create the system for the *micelle example*.

---

### 10.4.23 moltemplate tool

The moltemplate sub-directory contains instructions for installing moltemplate, a Python-based tool for building molecular systems based on a text-file description, and creating LAMMPS data files that encode their molecular topology as lists of bonds, angles, dihedrals, etc. See the README.txt file for more information.

This tool was written by Andrew Jewett (jewett.aj at gmail.com), who supports it. It has its own WWW page at <https://moltemplate.org>. The latest sources can be found on its [GitHub page](#)

---

### 10.4.24 msi2lmp tool

The msi2lmp sub-directory contains a tool for creating LAMMPS template input and data files from BIOVIA’s **Materials Studio** files (formerly **Accelrys’ Insight MD** code, formerly **MSI/Biosym** and its **Discover MD** code).

This tool was written by John Carpenter (Cray), Michael Peachey (Cray), and Steve Lustig (Dupont). Several people contributed changes to remove bugs and adapt its output to changes in LAMMPS.

This tool has several known limitations and is no longer under active development, so there are no changes except for the occasional bug fix.

---



See the README file in the tools/msi2lmp folder for more information.

## 10.4.25 Scripts for building LAMMPS when offline

In some situations it might be necessary to build LAMMPS on a system without direct internet access. The scripts in tools/offline folder allow you to pre-load external dependencies for both the documentation build and for building LAMMPS with CMake.

It does so by

1. downloading necessary pip packages,
2. cloning git repositories
3. downloading tarballs

to a designated cache folder.

As of April 2021, all of these downloads make up around 600MB. By default, the offline scripts will download everything into the \$HOME/.cache/lammps folder, but this can be changed by setting the LAMMPS\_CACHING\_DIR environment variable.

Once the caches have been initialized, they can be used for building the LAMMPS documentation or compiling LAMMPS using CMake on an offline system.

The use\_caches.sh script must be sourced into the current shell to initialize the offline build environment. Note that it must use the same LAMMPS\_CACHING\_DIR. This script does the following:

1. Set up environment variables that modify the behavior of both, pip and git
2. Start a simple local HTTP server using Python to host files for CMake

Afterwards, it will print out instruction on how to modify the CMake command line to make sure it uses the local HTTP server.

To undo the environment changes and shutdown the local HTTP server, run the deactivate\_caches command.

### Examples

For all of the examples below, you first need to create the cache, which requires an internet connection.

```
./tools/offline/init_caches.sh
```

Afterwards, you can disconnect or copy the contents of the LAMMPS\_CACHING\_DIR folder to an offline system.

### Documentation Build

The documentation build will create a new virtual environment that typically first installs dependencies from pip. With the offline environment loaded, these installations will instead grab the necessary packages from your local cache.

```
# if LAMMPS_CACHING_DIR is different from default, make sure to set it first
# export LAMMPS_CACHING_DIR=path/to/folder
source tools/offline/use_caches.sh
cd doc/
make html
```

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```
deactivate_caches
```

## CMake Build

When compiling certain packages with external dependencies, the CMake build system will download necessary files or sources from the web. For more flexibility the CMake configuration allows users to specify the URL of each of these dependencies. What the `init_caches.sh` script does is create a CMake “preset” file, which sets the URLs for all of the known dependencies and redirects the download to the local cache.

```
# if LAMMPS_CACHING_DIR is different from default, make sure to set it first
# export LAMMPS_CACHING_DIR=path/to/folder
source tools/offline/use_caches.sh

mkdir build
cd build
cmake -D LAMMPS_DOWNLOADS_URL=${HTTP_CACHE_URL} -C "${LAMMPS_HTTP_CACHE_CONFIG}" -C ../
↪cmake/presets/most.cmake ../cmake
make -j 8

deactivate_caches
```

---

### 10.4.26 phonon tool

The phonon sub-directory contains a post-processing tool useful for analyzing the output of the *fix phonon* command in the PHONON package.

See the README file for instruction on building the tool and what library it needs. And see the examples/PACKAGES/phonon directory for example problems that can be post-processed with this tool.

This tool was written by Ling-Ti Kong at Shanghai Jiao Tong University.

---

### 10.4.27 polybond tool

The polybond sub-directory contains a Python-based tool useful for performing “programmable polymer bonding”. The Python file `lmpsdata.py` provides a “Lmpsdata” class with various methods which can be invoked by a user-written Python script to create data files with complex bonding topologies.

See the Manual.pdf for details and example scripts.

This tool was written by Zachary Kraus at Georgia Tech.

---

### 10.4.28 pymol\_asphere tool

The `pymol_asphere` sub-directory contains a tool for converting a LAMMPS dump file that contains orientation info for ellipsoidal particles into an input file for the [PyMol visualization package](#) or its [open source variant](#).

Specifically, the tool triangulates the ellipsoids so they can be viewed as true ellipsoidal particles within PyMol. See the `README` and `examples` directory within `pymol_asphere` for more information.

This tool was written by Mike Brown at Sandia.

---

### 10.4.29 python tool

The `python` sub-directory contains several Python scripts that perform common LAMMPS post-processing tasks, such as:

- extract thermodynamic info from a log file as columns of numbers
- plot two columns of thermodynamic info from a log file using GnuPlot
- sort the snapshots in a dump file by atom ID
- convert multiple *NEB* dump files into one dump file for viz
- convert dump files into XYZ, CFG, or PDB format for viz by other packages

These are simple scripts built on [Pizza.py](#) modules. See the `README` for more info on `Pizza.py` and how to use these scripts.

---

### 10.4.30 replica tool

The `tools/replica` directory contains the `reorder_remd_traj` python script which can be used to reorder the replica trajectories (resulting from the use of the `temper` command) according to temperature. This will produce discontinuous trajectories with all frames at the same temperature in each trajectory. Additional options can be used to calculate the canonical configurational log-weight for each frame at each temperature using the `pymbar` package. See the `README.md` file for further details. Try out the peptide example provided.

This tool was written by (and is maintained by) Tanmoy Sanyal, while at the Shell lab at UC Santa Barbara. (tanmoy dot 7989 at gmail.com)

---

### 10.4.31 smd tool

The `smd` sub-directory contains a C++ file `dump2vtk_tris.cpp` and `Makefile` which can be compiled and used to convert triangle output files created by the Smooth-Mach Dynamics (MACHDYN) package into a VTK-compatible unstructured grid file. It could then be read in and visualized by VTK.

See the header of `dump2vtk.cpp` for more details.

This tool was written by the MACHDYN package author, Georg Ganzenmuller at the Fraunhofer-Institute for High-Speed Dynamics, Ernst Mach Institute in Germany ([georg.ganzenmueller@emi.fhg.de](mailto:georg.ganzenmueller@emi.fhg.de)).

---

### 10.4.32 spin tool

The spin sub-directory contains a C file `interpolate.c` which can be compiled and used to perform a cubic polynomial interpolation of the MEP following a GNEB calculation.

See the README file in `tools/spin/interpolate_gneb` for more details.

This tool was written by the SPIN package author, Julien Tranchida at Sandia National Labs (`jtranch` at `sandia.gov`, and by Aleksei Ivanov, at University of Iceland (`ali5` at `hi.is`).

---

### 10.4.33 singularity/apptainer tool

The singularity sub-directory contains container definitions files that can be used to build container images for building and testing LAMMPS on specific OS variants using the [Apptainer](#) or [Singularity](#) container software. Contributions for additional variants are welcome. For more details please see the README.md file in that folder.

---

### 10.4.34 stl\_bin2txt tool

The file `stl_bin2txt.cpp` converts binary STL files - like they are frequently offered for download on the web - into ASCII format STL files that LAMMPS can read with the `create_atoms mesh` or the `fix smd/wall_surface` commands. The syntax for running the tool is

```
stl_bin2txt infile.stl outfile.stl
```

which creates `outfile.stl` from `infile.stl`. This tool must be compiled on a platform compatible with the byte-ordering that was used to create the binary file. This usually is a so-called little endian hardware (like x86).

---

### 10.4.35 SWIG interface

The [SWIG tool](#) offers a mostly automated way to incorporate compiled code modules into scripting languages. It processes the function prototypes in C and generates wrappers for a wide variety of scripting languages from it. Thus it can also be applied to the [C language library interface](#) of LAMMPS so that build a wrapper that allows to call LAMMPS from programming languages like: C#/Mono, Lua, Java, JavaScript, Perl, Python, R, Ruby, Tcl, and more.

#### What is included

We provide here an “interface file”, `lammps.i`, that has the content of the `library.h` file adapted so SWIG can process it. That will create wrappers for all the functions that are present in the LAMMPS C library interface. Please note that not all kinds of C functions can be automatically translated, so you would have to add custom functions to be able to utilize those where the automatic translation does not work. A few functions for converting pointers and accessing arrays are predefined. We provide the file here on an “as is” basis to help people getting started, but not as a fully tested and supported feature of the LAMMPS distribution. Any contributions to complete this are, of course, welcome. Please also note, that for the case of creating a Python wrapper, a fully supported [Ctypes based lammps module](#) already exists. That module is designed to be object oriented while SWIG will generate a 1:1 translation of the functions in the interface file.

## Building the wrapper

When using CMake, the build steps for building a wrapper module are integrated for the languages: Java, Lua, Perl5, Python, Ruby, and Tcl. These require that the LAMMPS library is build as a shared library and all necessary development headers and libraries are present.

```
-D WITH_SWIG=on      # to enable building any SWIG wrapper
-D BUILD_SWIG_JAVA=on # to enable building the Java wrapper
-D BUILD_SWIG_LUA=on  # to enable building the Lua wrapper
-D BUILD_SWIG_PERL5=on # to enable building the Perl 5.x wrapper
-D BUILD_SWIG_PYTHON=on # to enable building the Python wrapper
-D BUILD_SWIG_RUBY=on  # to enable building the Ruby wrapper
-D BUILD_SWIG_TCL=on  # to enable building the Tcl wrapper
```

Manual building allows a little more flexibility. E.g. one can choose the name of the module and build and use a dynamically loaded object for Tcl with:

```
$ swig -tcl -module tcllammps lammps.i
$ gcc -fPIC -shared $(pkgconf --cflags tcl) -o tcllammps.so \
    lammps_wrap.c -L ../src/ -llammps
$ tclsh
```

Or one can build an extended Tcl shell command with the wrapped functions included with:

```
$ swig -tcl -module tcllmps lammps_shell.i
$ gcc -o tcllmpsh lammps_wrap.c -Xlinker -export-dynamic \
    -DHAVE_CONFIG_H $(pkgconf --cflags tcl) \
    $(pkgconf --libs tcl) -L ../src -llammps
```

In both cases it is assumed that the LAMMPS library was compiled as a shared library in the src folder. Otherwise the last part of the commands needs to be adjusted.

## Utility functions

Definitions for several utility functions required to manage and access data passed or returned as pointers are included in the lammps.i file. So most of the functionality of the library interface should be accessible. What works and what does not depends a bit on the individual language for which the wrappers are built and how well SWIG supports those. The [SWIG documentation](#) has very detailed instructions and recommendations.

## Usage examples

The tools/swig folder has multiple shell scripts, run\_<name>\_example.sh that will create a small example script and demonstrate how to load the wrapper and run LAMMPS through it in the corresponding programming language.

For illustration purposes below is a part of the Tcl example script.

```
% load ./tcllammps.so
% set lmp [lammps_open_no_mpi 0 NULL NULL]
% lammps_command $lmp "units real"
% lammps_command $lmp "lattice fcc 2.5"
% lammps_command $lmp "region box block -5 5 -5 5 -5 5"
% lammps_command $lmp "create_box 1 box"
% lammps_command $lmp "create_atoms 1 box"
```

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```
%  
% set dt [doublep_value [voidp_to_doublep [lammmps_extract_global $lmp dt]]]  
% puts "LAMMPS version $ver"  
% puts [format "Number of created atoms: %g" [lammmps_get_natoms $lmp]]  
% puts "Current size of timestep: $dt"  
% puts "LAMMPS version: [lammmps_version $lmp]"  
% lammmps_close $lmp
```

---

### 10.4.36 vim tool

The files in the tools/vim directory are add-ons to the VIM editor that allow easier editing of LAMMPS input scripts. See the README.txt file for details.

These files were provided by Gerolf Ziegenhain (gerolf at ziegenhain.com)

---

### 10.4.37 xmgrace tool

The files in the tools/xmgrace directory can be used to plot the thermodynamic data in LAMMPS log files via the xmgrace plotting package. There are several tools in the directory that can be used in post-processing mode. The lammplot.cpp file can be compiled and used to create plots from the current state of a running LAMMPS simulation.

See the README file for details.

These files were provided by Vikas Varshney (vv0210 at gmail.com)

## ERRORS

These doc pages describe many of the error and warning message you can encounter when using LAMMPS. The common problems include conceptual issues. The messages and warnings doc pages give complete lists of all the messages the code may generate, with additional details for many of them.

### 11.1 Common problems

If two LAMMPS runs do not produce the exact same answer on different machines or different numbers of processors, this is typically not a bug. In theory you should get identical answers on any number of processors and on any machine. In practice, numerical round-off can cause slight differences and eventual divergence of molecular dynamics phase space trajectories within a few 100s or few 1000s of timesteps. However, the statistical properties of the two runs (e.g. average energy or temperature) should still be the same.

If the *velocity* command is used to set initial atom velocities, a particular atom can be assigned a different velocity when the problem is run on a different number of processors or on different machines. If this happens, the phase space trajectories of the two simulations will rapidly diverge. See the discussion of the *loop* option in the *velocity* command for details and options that avoid this issue.

Similarly, the *create\_atoms* command generates a lattice of atoms. For the same physical system, the ordering and numbering of atoms by atom ID may be different depending on the number of processors.

Some commands use random number generators which may be setup to produce different random number streams on each processor and hence will produce different effects when run on different numbers of processors. A commonly-used example is the *fix langevin* command for thermostating.

A LAMMPS simulation typically has two stages, setup and run. Most LAMMPS errors are detected at setup time; others like a bond stretching too far may not occur until the middle of a run.

LAMMPS tries to flag errors and print informative error messages so you can fix the problem. For most errors it will also print the last input script command that it was processing. Of course, LAMMPS cannot figure out your physics or numerical mistakes, like choosing too big a timestep, specifying erroneous force field coefficients, or putting 2 atoms on top of each other! If you run into errors that LAMMPS does not catch that you think it should flag, please send an email to the [developers](#).

If you get an error message about an invalid command in your input script, you can determine what command is causing the problem by looking in the `log.lammps` file or using the *echo command* to see it on the screen. If you get an error like “Invalid ... style”, with ... being *fix*, *compute*, *pair*, etc, it means that you mistyped the style name or that the command is part of an optional package which was not compiled into your executable. The list of available styles in your executable can be listed by using *the -h command-line switch*. The installation and compilation of optional packages is explained on the [Build packages](#) doc page.

For a given command, LAMMPS expects certain arguments in a specified order. If you mess this up, LAMMPS will often flag the error, but it may also simply read a bogus argument and assign a value that is valid, but not what you wanted. E.g. trying to read the string “abc” as an integer value of 0. Careful reading of the associated doc page for the

command should allow you to fix these problems. In most cases, where LAMMPS expects to read a number, either integer or floating point, it performs a stringent test on whether the provided input actually is an integer or floating-point number, respectively, and reject the input with an error message (for instance, when an integer is required, but a floating-point number 1.0 is provided):

```
ERROR: Expected integer parameter instead of '1.0' in input script or data file
```

Some commands allow for using variable references in place of numeric constants so that the value can be evaluated and may change over the course of a run. This is typically done with the syntax `v_name` for a parameter, where name is the name of the variable. On the other hand, immediate variable expansion with the syntax `${name}` is performed while reading the input and before parsing commands,

---

**Note:** Using a variable reference (i.e. `v_name`) is only allowed if the documentation of the corresponding command explicitly says it is. Otherwise, you will receive an error message of this kind:

---

```
ERROR: Expected floating point parameter instead of 'v_name' in input script or data file
```

Generally, LAMMPS will print a message to the screen and logfile and exit gracefully when it encounters a fatal error. Sometimes it will print a WARNING to the screen and logfile and continue on; you can decide if the WARNING is important or not. A WARNING message that is generated in the middle of a run is only printed to the screen, not to the logfile, to avoid cluttering up thermodynamic output. If LAMMPS crashes or hangs without spitting out an error message first then it could be a bug (see [this section](#)) or one of the following cases:

LAMMPS runs in the available memory a processor allows to be allocated. Most reasonable MD runs are compute limited, not memory limited, so this should not be a bottleneck on most platforms. Almost all large memory allocations in the code are done via C-style malloc's which will generate an error message if you run out of memory. Smaller chunks of memory are allocated via C++ "new" statements. If you are unlucky you could run out of memory just when one of these small requests is made, in which case the code will crash or hang (in parallel), since LAMMPS does not trap on those errors.

Illegal arithmetic can cause LAMMPS to run slow or crash. This is typically due to invalid physics and numerics that your simulation is computing. If you see wild thermodynamic values or NaN values in your LAMMPS output, something is wrong with your simulation. If you suspect this is happening, it is a good idea to print out thermodynamic info frequently (e.g. every timestep) via the `thermo` so you can monitor what is happening. Visualizing the atom movement is also a good idea to insure your model is behaving as you expect.

In parallel, one way LAMMPS can hang is due to how different MPI implementations handle buffering of messages. If the code hangs without an error message, it may be that you need to specify an MPI setting or two (usually via an environment variable) to enable buffering or boost the sizes of messages that can be buffered.

## 11.2 Error and warning details

Many errors or warnings are self-explanatory and thus straightforward to resolve. However, there are also cases, where there is no single cause and explanation, where LAMMPS can only detect symptoms of an error but not the exact cause, or where the explanation needs to be more detailed than what can be fit into a message printed by the program. The following are discussions of such cases.



### 11.2.1 Unknown identifier in data file

This error happens when LAMMPS encounters a line of text in an unexpected format while reading a data file. This is most commonly caused by inconsistent header and section data. The header section informs LAMMPS how many entries or lines are expected in the various sections (like Atoms, Masses, Pair Coeffs, *etc.*) of the data file. If there is a mismatch, LAMMPS will either keep reading beyond the end of a section or stop reading before the section has ended.

Such a mismatch can happen unexpectedly when the first line of the data is *not* a comment as required by the format. That would result in LAMMPS expecting, for instance, 0 atoms because the “atoms” header line is treated as a comment.

## 11.3 Reporting bugs

If you are confident that you have found a bug in LAMMPS, please follow the steps outlined below:

- Check the [New features and bug fixes](#) section of the [LAMMPS WWW site](#) or the [GitHub Releases](#) page to see if the bug has already been addressed in a patch release.
- Check that your issue can be reproduced with the latest development version of LAMMPS.
- Check the manual carefully to verify that the unexpected behavior you are observing is indeed in conflict with the documentation
- Check the [GitHub Issue](#) page if your issue has already been reported and if it is still open.
- Check the [GitHub Pull Requests](#) page to see if there is already a fix for your bug pending.
- Check the [LAMMPS forum at MatSci](#) to see if the issue has been discussed before.

If none of these steps yields any useful information, please file a new bug report on the [GitHub Issue](#) page. The website will offer you to select a suitable template with explanations and then you should replace those explanations with the information that you can provide to reproduce your issue.

The most useful thing you can do to help us verify and fix a bug is to isolate the problem. Run it on the smallest number of atoms and fewest number of processors with the simplest input script that reproduces the bug. Try to identify what command or combination of commands is causing the problem and upload the complete input deck as a tar or zip archive. Please avoid using binary restart files unless the issue requires it. In the latter case you should also include an input deck to quickly generate this restart from a data file or a simple additional input. This input deck can be used with tools like a debugger or `valgrind` to further *debug the crash*.

You may also post a message in the [development category of the LAMMPS forum at MatSci](#) describing the problem with the same kind of information. The forum can provide a faster response, especially if the bug reported is actually expected behavior or other LAMMPS users have come across it before.

## 11.4 Debugging crashes

If LAMMPS crashes with a “segmentation fault” or a “bus error” or similar message, then you can use the following two methods to further narrow down the origin of the issue. This will help the LAMMPS developers (or yourself) to understand the reason for the crash and apply a fix (either to the input script or the source code). This requires that your LAMMPS executable includes the required *debug information*. Otherwise it is not possible to look up the names of functions or variables.

The following patch will introduce a bug into the code for pair style `lj/cut` when using the `examples/melt/in.melt` input. We use it to show how to identify the origin of a segmentation fault.

```

--- a/src/pair_lj_cut.cpp
+++ b/src/pair_lj_cut.cpp
@@ -81,6 +81,7 @@ void PairLJCut::compute(int eflag, int vflag)
    int nlocal = atom->nlocal;
    double *special_lj = force->special_lj;
    int newton_pair = force->newton_pair;
+   double comx = 0.0;

    inum = list->inum;
    ilist = list->ilist;
@@ -134,8 +135,10 @@ void PairLJCut::compute(int eflag, int vflag)
                                evdwl,0.0,fpair,delx,dely,delz);
    }
}
- }

+   comx += atom->rmass[i]*x[i][0]; /* BUG */
+ }
+ printf("comx = %g\n",comx);
+ if (vflag_fdotr) virial_fdotr_compute();
}

```

After recompiling LAMMPS and running the input you should get something like this:

```

$ ./lmp -in in.melt
LAMMPS (19 Mar 2020)
  using 1 OpenMP thread(s) per MPI task
Lattice spacing in x,y,z = 1.6796 1.6796 1.6796
Created orthogonal box = (0 0 0) to (16.796 16.796 16.796)
  1 by 1 by 1 MPI processor grid
Created 4000 atoms
  create_atoms CPU = 0.000432253 secs
Neighbor list info ...
  update every 20 steps, delay 0 steps, check no
  max neighbors/atom: 2000, page size: 100000
  master list distance cutoff = 2.8
  ghost atom cutoff = 2.8
  binsize = 1.4, bins = 12 12 12
  1 neighbor lists, perpetual/occasional/extra = 1 0 0
  (1) pair lj/cut, perpetual
    attributes: half, newton on
    pair build: half/bin/atomonly/newton
    stencil: half/bin/3d/newton
    bin: standard
Setting up Verlet run ...
  Unit style      : lj
  Current step   : 0
  Time step      : 0.005
Segmentation fault (core dumped)

```

### 11.4.1 Using the GDB debugger to get a stack trace

There are two options to use the GDB debugger for identifying the origin of the segmentation fault or similar crash. The GDB debugger has many more features and options, as can be seen for example its [online documentation](#).

#### Run LAMMPS from within the debugger

Running LAMMPS under the control of the debugger as shown below only works for a single MPI rank (for debugging a program running in parallel you usually need a parallel debugger program). A simple way to launch GDB is to prefix the LAMMPS command line with `gdb --args` and then type the command “run” at the GDB prompt. This will launch the debugger, load the LAMMPS executable and its debug info, and then run it. When it reaches the code causing the segmentation fault, it will stop with a message why it stopped, print the current line of code, and drop back to the GDB prompt.

```
[...]
Setting up Verlet run ...
  Unit style      : lj
  Current step    : 0
  Time step       : 0.005

Program received signal SIGSEGV, Segmentation fault.
0x00000000006653ab in LAMMPS_NS::PairLJCut::compute (this=0x829740, eflag=1, vflag=
↳<optimized out>) at /home/akohlmey/compile/lammps/src/pair_lj_cut.cpp:139
139      comx += atom->rmass[i]*x[i][0]; /* BUG */
(gdb)
```

Now typing the command “where” will show the stack of functions starting from the current function back to “main()”.

```
(gdb) where
#0  0x00000000006653ab in LAMMPS_NS::PairLJCut::compute (this=0x829740, eflag=1, vflag=
↳<optimized out>) at /home/akohlmey/compile/lammps/src/pair_lj_cut.cpp:139
#1  0x00000000004cf0a2 in LAMMPS_NS::Verlet::setup (this=0x7e6c90, flag=1) at /home/
↳akohlmey/compile/lammps/src/verlet.cpp:131
#2  0x000000000049db42 in LAMMPS_NS::Run::command (this=this@entry=0x7fffffffcca0,
↳narg=narg@entry=1, arg=arg@entry=0x7e8750)
    at /home/akohlmey/compile/lammps/src/run.cpp:177
#3  0x000000000041258a in LAMMPS_NS::Input::command_creator<LAMMPS_NS::Run> (lmp=
↳<optimized out>, narg=1, arg=0x7e8750)
    at /home/akohlmey/compile/lammps/src/input.cpp:878
#4  0x0000000000410ad3 in LAMMPS_NS::Input::execute_command (this=0x7d1410) at /home/
↳akohlmey/compile/lammps/src/input.cpp:864
#5  0x00000000004111fb in LAMMPS_NS::Input::file (this=0x7d1410) at /home/akohlmey/
↳compile/lammps/src/input.cpp:229
#6  0x000000000040933a in main (argc=<optimized out>, argv=<optimized out>) at /home/
↳akohlmey/compile/lammps/src/main.cpp:65
(gdb)
```

You can also print the value of variables and see if there is anything unexpected. Segmentation faults, for example, commonly happen when a pointer variable is not assigned and still initialized to NULL.

```
(gdb) print x
$1 = (double **) 0x7ffff7ca1010
(gdb) print i
```

(continues on next page)

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```

$2 = 0
(gdb) print x[0]
$3 = (double *) 0x7ffff6d80010
(gdb) print x[0][0]
$4 = 0
(gdb) print x[1][0]
$5 = 0.83979809569125363
(gdb) print atom->rmass
$6 = (double *) 0x0
(gdb)

```

### Inspect a core dump file with the debugger

When an executable crashes with a “core dumped” message, it creates a file “core” or “core.<PID#>” which contains the information about the current state. This file may be located in the folder where you ran LAMMPS or in some hidden folder managed by the systemd daemon. In the latter case, you need to “extract” the core file with the `coredumpctl` utility to the current folder. Example: `coredumpctl -o core dump lmp`. Now you can launch the debugger to load the executable, its debug info and the core dump and drop you to a prompt like before.

```

$ gdb lmp core
Reading symbols from lmp...
[New LWP 1928535]
[Thread debugging using libthread_db enabled]
Using host libthread_db library "/lib64/libthread_db.so.1".
Core was generated by `./lmp -in in.melt'.
Program terminated with signal SIGSEGV, Segmentation fault.
#0  0x00000000006653ab in LAMMPS_NS::PairLJCut::compute (this=0x1b10740, eflag=1, vflag=
↳<optimized out>)
    at /home/akohlmeier/compile/lammps/src/pair_lj_cut.cpp:139
139      comx += atom->rmass[i]*x[i][0]; /* BUG */
(gdb)

```

From here on, you use the same commands as shown before to get a stack trace and print current values of (pointer) variables.

### 11.4.2 Using valgrind to get a stack trace

The `valgrind` suite of tools allows to closely inspect the behavior of a compiled program by essentially emulating a CPU and instrumenting the program while running. This slows down execution quite significantly, but can also report issues that are not resulting in a crash. The default valgrind tool is a memory checker and you can use it by prefixing the normal command line with `valgrind`. Unlike GDB, this will also work for parallel execution, but it is recommended to redirect the valgrind output to a file (e.g. with `--log-file=crash-%p.txt`, the `%p` will be substituted with the process ID) so that the messages of the multiple valgrind instances to the console are not mixed.

```

$ valgrind ./lmp -in in.melt
==1933642== Memcheck, a memory error detector
==1933642== Copyright (C) 2002-2017, and GNU GPL'd, by Julian Seward et al.
==1933642== Using Valgrind-3.15.0 and LibVEX; rerun with -h for copyright info
==1933642== Command: ./lmp -in in.melt
==1933642==

```

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```

LAMMPS (19 Mar 2020)
OMP_NUM_THREADS environment is not set. Defaulting to 1 thread. (src/comm.cpp:94)
  using 1 OpenMP thread(s) per MPI task
Lattice spacing in x,y,z = 1.6796 1.6796 1.6796
Created orthogonal box = (0 0 0) to (16.796 16.796 16.796)
  1 by 1 by 1 MPI processor grid
Created 4000 atoms
  create_atoms CPU = 0.032964 secs
Neighbor list info ...
  update every 20 steps, delay 0 steps, check no
  max neighbors/atom: 2000, page size: 100000
  master list distance cutoff = 2.8
  ghost atom cutoff = 2.8
  binsize = 1.4, bins = 12 12 12
  1 neighbor lists, perpetual/occasional/extra = 1 0 0
  (1) pair lj/cut, perpetual
    attributes: half, newton on
    pair build: half/bin/atomonly/newton
    stencil: half/bin/3d/newton
    bin: standard
Setting up Verlet run ...
  Unit style      : lj
  Current step    : 0
  Time step       : 0.005
==1933642== Invalid read of size 8
==1933642==    at 0x6653AB: LAMMPS_NS::PairLJCut::compute(int, int) (pair_lj_cut.cpp:139)
==1933642==    by 0x4CF0A1: LAMMPS_NS::Verlet::setup(int) (verlet.cpp:131)
==1933642==    by 0x49DB41: LAMMPS_NS::Run::command(int, char**) (run.cpp:177)
==1933642==    by 0x412589: void LAMMPS_NS::Input::command_creator<LAMMPS_NS::Run>
↳(LAMMPS_NS::LAMMPS*, int, char**) (input.cpp:881)
==1933642==    by 0x410AD2: LAMMPS_NS::Input::execute_command() (input.cpp:864)
==1933642==    by 0x4111FA: LAMMPS_NS::Input::file() (input.cpp:229)
==1933642==    by 0x409339: main (main.cpp:65)
==1933642== Address 0x0 is not stack'd, malloc'd or (recently) free'd
==1933642==

```

As you can see, the stack trace information is similar to that obtained from GDB. In addition you get a more specific hint about what cause the segmentation fault, i.e. that it is a NULL pointer dereference. To find out which pointer exactly was NULL, you need to use the debugger, though.

## 11.5 Error messages

This is an alphabetic list of the ERROR messages LAMMPS prints out and the reason why. If the explanation here is not sufficient, the documentation for the offending command may help. Error messages also list the source file and line number where the error was generated. For example, a message like this:

```
ERROR: Illegal velocity command (velocity.cpp:78)
```

means that line #78 in the file src/velocity.cpp generated the error. Looking in the source code may help you figure out what went wrong.

Doc page with [WARNING messages](#)

***1-3 bond count is inconsistent***

An inconsistency was detected when computing the number of 1-3 neighbors for each atom. This likely means something is wrong with the bond topologies you have defined.

***1-4 bond count is inconsistent***

An inconsistency was detected when computing the number of 1-4 neighbors for each atom. This likely means something is wrong with the bond topologies you have defined.

***Accelerator sharing is not currently supported on system***

Multiple MPI processes cannot share the accelerator on your system. For NVIDIA GPUs, see the `nvidia-smi` command to change this setting.

***All angle coeffs are not set***

All angle coefficients must be set in the data file or by the `angle_coeff` command before running a simulation.

***All atom IDs = 0 but atom\_modify id = yes***

Self-explanatory.

***All atoms of a swapped type must have same charge.***

Self-explanatory.

***All atoms of a swapped type must have the same charge.***

Self-explanatory.

***All bond coeffs are not set***

All bond coefficients must be set in the data file or by the `bond_coeff` command before running a simulation.

***All dihedral coeffs are not set***

All dihedral coefficients must be set in the data file or by the `dihedral_coeff` command before running a simulation.

***All improper coeffs are not set***

All improper coefficients must be set in the data file or by the `improper_coeff` command before running a simulation.

***All masses are not set***

For atom styles that define masses for each atom type, all masses must be set in the data file or by the `mass` command before running a simulation. They must also be set before using the `velocity` command.

***All mol IDs should be set for fix gcmc group atoms***

The molecule flag is on, yet not all molecule ids in the fix group have been set to non-zero positive values by the user. This is an error since all atoms in the fix gcmc group are eligible for deletion, rotation, and translation and therefore must have valid molecule ids.

***All pair coeffs are not set***

All pair coefficients must be set in the data file or by the `pair_coeff` command before running a simulation.

***All read\_dump x,y,z fields must be specified for scaled, triclinic coords***

For triclinic boxes and scaled coordinates you must specify all 3 of the x,y,z fields, else LAMMPS cannot reconstruct the unscaled coordinates.

***All universe/uloop variables must have same # of values***

Self-explanatory.

***All variables in next command must be same style***

Self-explanatory.

***Angle atom missing in delete\_bonds***

The `delete_bonds` command cannot find one or more atoms in a particular angle on a particular processor. The pairwise cutoff is too short or the atoms are too far apart to make a valid angle.

***Angle atom missing in set command***

The set command cannot find one or more atoms in a particular angle on a particular processor. The pairwise cutoff is too short or the atoms are too far apart to make a valid angle.

***Angle atoms %d %d %d missing on proc %d at step %ld***

One or more of 3 atoms needed to compute a particular angle are missing on this processor. Typically this is because the pairwise cutoff is set too short or the angle has blown apart and an atom is too far away.

***Angle atoms missing on proc %d at step %ld***

One or more of 3 atoms needed to compute a particular angle are missing on this processor. Typically this is because the pairwise cutoff is set too short or the angle has blown apart and an atom is too far away.

***Angle coeff for hybrid has invalid style***

Angle style hybrid uses another angle style as one of its coefficients. The angle style used in the angle\_coeff command or read from a restart file is not recognized.

***Angle coeffs are not set***

No angle coefficients have been assigned in the data file or via the angle\_coeff command.

***Angle extent > half of periodic box length***

This error was detected by the neigh\_modify check yes setting. It is an error because the angle atoms are so far apart it is ambiguous how it should be defined.

***Angle potential must be defined for SHAKE***

When shaking angles, an angle\_style potential must be used.

***Angle style hybrid cannot have hybrid as an argument***

Self-explanatory.

***Angle style hybrid cannot have none as an argument***

Self-explanatory.

***Angle style hybrid cannot use same angle style twice***

Self-explanatory.

***Angle table must range from 0 to 180 degrees***

Self-explanatory.

***Angle table parameters did not set N***

List of angle table parameters must include N setting.

***Angle\_coeff command before angle\_style is defined***

Coefficients cannot be set in the data file or via the angle\_coeff command until an angle\_style has been assigned.

***Angle\_coeff command before simulation box is defined***

The angle\_coeff command cannot be used before a read\_data, read\_restart, or create\_box command.

***Angle\_coeff command when no angles allowed***

The chosen atom style does not allow for angles to be defined.

***Angle\_style command when no angles allowed***

The chosen atom style does not allow for angles to be defined.

***Angles assigned incorrectly***

Angles read in from the data file were not assigned correctly to atoms. This means there is something invalid about the topology definitions.

***Angles defined but no angle types***

The data file header lists angles but no angle types.

***Append boundary must be shrink/minimum***

The boundary style of the face where atoms are added must be of type m (shrink/minimum).

***Arccos of invalid value in variable formula***

Argument of arccos() must be between -1 and 1.

***Arcsin of invalid value in variable formula***

Argument of arcsin() must be between -1 and 1.

***Assigning body parameters to non-body atom***

Self-explanatory.

***Assigning ellipsoid parameters to non-ellipsoid atom***

Self-explanatory.

***Assigning line parameters to non-line atom***

Self-explanatory.

***Assigning quat to non-body atom***

Self-explanatory.

***Assigning tri parameters to non-tri atom***

Self-explanatory.

***At least one atom of each swapped type must be present to define charges.***

Self-explanatory.

***Atom IDs must be consecutive for velocity create loop all***

Self-explanatory.

***Atom IDs must be used for molecular systems***

Atom IDs are used to identify and find partner atoms in bonds.

***Atom count changed in fix neb***

This is not allowed in a NEB calculation.

***Atom count is inconsistent, cannot write data file***

The sum of atoms across processors does not equal the global number of atoms. Probably some atoms have been lost.

***Atom count is inconsistent, cannot write restart file***

Sum of atoms across processors does not equal initial total count. This is probably because you have lost some atoms.

***Atom in too many rigid bodies - boost MAXBODY***

Fix poems has a parameter MAXBODY (in fix\_poems.cpp) which determines the maximum number of rigid bodies a single atom can belong to (i.e. a multibody joint). The bodies you have defined exceed this limit.

***Atom sort did not operate correctly***

This is an internal LAMMPS error. Please report it to the developers.

***Atom style hybrid cannot have hybrid as an argument***

Self-explanatory.

***Atom style hybrid cannot use same atom style twice***

Self-explanatory.

***Atom style template molecule must have atom types***

The defined molecule(s) does not specify atom types.

***Atom style was redefined after using fix property/atom***

This is not allowed.

***Atom type must be zero in fix gcmc mol command***

Self-explanatory.



***Atom vector in equal-style variable formula***

Atom vectors generate one value per atom which is not allowed in an equal-style variable.

***Atom-style variable in equal-style variable formula***

Atom-style variables generate one value per atom which is not allowed in an equal-style variable.

***Atom\_modify id command after simulation box is defined***

The atom\_modify id command cannot be used after a read\_data, read\_restart, or create\_box command.

***Atom\_modify map command after simulation box is defined***

The atom\_modify map command cannot be used after a read\_data, read\_restart, or create\_box command.

***Atom\_modify sort and first options cannot be used together***

Self-explanatory.

***Atom\_style command after simulation box is defined***

The atom\_style command cannot be used after a read\_data, read\_restart, or create\_box command.

***Atom\_style line can only be used in 2d simulations***

Self-explanatory.

***Atom\_style tri can only be used in 3d simulations***

Self-explanatory.

***Atomfile variable could not read values***

Check the file assigned to the variable.

***Atomfile variable in equal-style variable formula***

Self-explanatory.

***Atomfile-style variable in equal-style variable formula***

Self-explanatory.

***Attempt to pop empty stack in fix box/relax***

Internal LAMMPS error. Please report it to the developers.

***Attempt to push beyond stack limit in fix box/relax***

Internal LAMMPS error. Please report it to the developers.

***Attempting to rescale a 0.0 temperature***

Cannot rescale a temperature that is already 0.0.

***Attempting to insert more particles than available lattice points***

Self-explanatory.

***Bad FENE bond***

Two atoms in a FENE bond have become so far apart that the bond cannot be computed.

***Bad TIP4P angle type for PPPM/TIP4P***

Specified angle type is not valid.

***Bad TIP4P angle type for PPPMDisp/TIP4P***

Specified angle type is not valid.

***Bad TIP4P bond type for PPPM/TIP4P***

Specified bond type is not valid.

***Bad TIP4P bond type for PPPMDisp/TIP4P***

Specified bond type is not valid.

***Bad fix ID in fix append/atoms command***

The value of the fix\_id for keyword spatial must start with "f\_".

***Bad grid of processors***

The 3d grid of processors defined by the processors command does not match the number of processors LAMMPS is being run on.

***Bad kspace\_modify kmax/ewald parameter***

Kspace\_modify values for the kmax/ewald keyword must be integers > 0

***Bad kspace\_modify slab parameter***

Kspace\_modify value for the slab/volume keyword must be >= 2.0.

***Bad matrix inversion in mldivide3***

This error should not occur unless the matrix is badly formed.

***Bad principal moments***

Fix rigid did not compute the principal moments of inertia of a rigid group of atoms correctly.

***Bad quadratic solve for particle/line collision***

This is an internal error. It should normally not occur.

***Bad quadratic solve for particle/tri collision***

This is an internal error. It should normally not occur.

***Bad real space Coulombic cutoff in fix tune/kspace***

Fix tune/kspace tried to find the optimal real space Coulombic cutoff using the Newton-Rhaphson method, but found a non-positive or NaN cutoff

***Balance command before simulation box is defined***

The balance command cannot be used before a read\_data, read\_restart, or create\_box command.

***Balance produced bad splits***

This should not occur. It means two or more cutting plane locations are on top of each other or out of order. Report the problem to the developers.

***Balance rcb cannot be used with comm\_style brick***

Comm\_style tiled must be used instead.

***Balance shift string is invalid***

The string can only contain the characters “x”, “y”, or “z”.

***Bias compute does not calculate a velocity bias***

The specified compute must compute a bias for temperature.

***Bias compute does not calculate temperature***

The specified compute must compute temperature.

***Bias compute group does not match compute group***

The specified compute must operate on the same group as the parent compute.

***Big particle in fix srd cannot be point particle***

Big particles must be extended spheroids or ellipsoids.

***Bigint setting in lmptype.h is invalid***

Size of bigint is less than size of tagint.

***Bigint setting in lmptype.h is not compatible***

Format of bigint stored in restart file is not consistent with LAMMPS version you are running. See the settings in src/lmptype.h

***Bitmapped lookup tables require int/float be same size***

Cannot use pair tables on this machine, because of word sizes. Use the pair\_modify command with table 0 instead.

***Bitmapped table in file does not match requested table***

Setting for bitmapped table in pair\_coeff command must match table in file exactly.

***Bitmapped table is incorrect length in table file***

Number of table entries is not a correct power of 2.

***Bond and angle potentials must be defined for TIP4P***

Cannot use TIP4P pair potential unless bond and angle potentials are defined.

***Bond atom missing in box size check***

The second atom needed to compute a particular bond is missing on this processor. Typically this is because the pairwise cutoff is set too short or the bond has blown apart and an atom is too far away.

***Bond atom missing in delete\_bonds***

The delete\_bonds command cannot find one or more atoms in a particular bond on a particular processor. The pairwise cutoff is too short or the atoms are too far apart to make a valid bond.

***Bond atom missing in image check***

The second atom in a particular bond is missing on this processor. Typically this is because the pairwise cutoff is set too short or the bond has blown apart and an atom is too far away.

***Bond atom missing in set command***

The set command cannot find one or more atoms in a particular bond on a particular processor. The pairwise cutoff is too short or the atoms are too far apart to make a valid bond.

***Bond atoms %d %d missing on proc %d at step %ld***

The second atom needed to compute a particular bond is missing on this processor. Typically this is because the pairwise cutoff is set too short or the bond has blown apart and an atom is too far away.

***Bond atoms missing on proc %d at step %ld***

The second atom needed to compute a particular bond is missing on this processor. Typically this is because the pairwise cutoff is set too short or the bond has blown apart and an atom is too far away.

***Bond coeff for hybrid has invalid style***

Bond style hybrid uses another bond style as one of its coefficients. The bond style used in the bond\_coeff command or read from a restart file is not recognized.

***Bond coeffs are not set***

No bond coefficients have been assigned in the data file or via the bond\_coeff command.

***Bond extent > half of periodic box length***

This error was detected by the neigh\_modify check yes setting. It is an error because the bond atoms are so far apart it is ambiguous how it should be defined.

***Bond potential must be defined for SHAKE***

Cannot use fix shake unless bond potential is defined.

***Bond style hybrid cannot have hybrid as an argument***

Self-explanatory.

***Bond style hybrid cannot have none as an argument***

Self-explanatory.

***Bond style hybrid cannot use same bond style twice***

Self-explanatory.

***Bond style quartic cannot be used with 3,4-body interactions***

No angle, dihedral, or improper styles can be defined when using bond style quartic.

***Bond style quartic cannot be used with atom style template***

This bond style can change the bond topology which is not allowed with this atom style.

***Bond style quartic requires special\_bonds = 1,1,1***

This is a restriction of the current bond quartic implementation.

***Bond table parameters did not set N***

List of bond table parameters must include N setting.

***Bond table values are not increasing***

The values in the tabulated file must be monotonically increasing.

***BondAngle coeff for hybrid angle has invalid format***

No “ba” field should appear in data file entry.

***BondBond coeff for hybrid angle has invalid format***

No “bb” field should appear in data file entry.

***Bond\_coeff command before bond\_style is defined***

Coefficients cannot be set in the data file or via the bond\_coeff command until an bond\_style has been assigned.

***Bond\_coeff command before simulation box is defined***

The bond\_coeff command cannot be used before a read\_data, read\_restart, or create\_box command.

***Bond\_coeff command when no bonds allowed***

The chosen atom style does not allow for bonds to be defined.

***Bond\_style command when no bonds allowed***

The chosen atom style does not allow for bonds to be defined.

***Bonds assigned incorrectly***

Bonds read in from the data file were not assigned correctly to atoms. This means there is something invalid about the topology definitions.

***Bonds defined but no bond types***

The data file header lists bonds but no bond types.

***Both restart files must use % or neither***

Self-explanatory.

***Both restart files must use MPI-IO or neither***

Self-explanatory.

***Both sides of boundary must be periodic***

Cannot specify a boundary as periodic only on the lo or hi side. Must be periodic on both sides.

***Boundary command after simulation box is defined***

The boundary command cannot be used after a read\_data, read\_restart, or create\_box command.

***Box bounds are invalid***

The box boundaries specified in the read\_data file are invalid. The lo value must be less than the hi value for all 3 dimensions.

***Box command after simulation box is defined***

The box command cannot be used after a read\_data, read\_restart, or create\_box command.

***CPU neighbor lists must be used for ellipsoid/sphere mix.***

When using Gay-Berne or RE-squared pair styles with both ellipsoidal and spherical particles, the neighbor list must be built on the CPU

***Can not specify Pxy/Pxz/Pyz in fix box/relax with non-triclinic box***

Only triclinic boxes can be used with off-diagonal pressure components. See the region prism command for details.

***Can not specify Pxy/Pxz/Pyz in fix nvt/npt/nph with non-triclinic box***

Only triclinic boxes can be used with off-diagonal pressure components. See the region prism command for details.

***Can only use -plog with multiple partitions***

Self-explanatory. See page discussion of command-line switches.

***Can only use -pscreen with multiple partitions***

Self-explanatory. See page discussion of command-line switches.

***Can only use Kokkos supported regions with Kokkos package***

Self-explanatory.

***Can only use NEB with 1-processor replicas***

This is current restriction for NEB as implemented in LAMMPS.

***Can only use TAD with 1-processor replicas for NEB***

This is current restriction for NEB as implemented in LAMMPS.

***Cannot (yet) do analytic differentiation with ppm/gpu***

This is a current restriction of this command.

***Cannot (yet) request ghost atoms with Kokkos half neighbor list***

This feature is not yet supported.

***Cannot (yet) use 'electron' units with dipoles***

This feature is not yet supported.

***Cannot (yet) use Ewald with triclinic box and slab correction***

This feature is not yet supported.

***Cannot (yet) use K-space slab correction with compute group/group for triclinic systems***

This option is not yet supported.

***Cannot (yet) use MSM with 2d simulation***

This feature is not yet supported.

***Cannot (yet) use PPPM with triclinic box and TIP4P***

This feature is not yet supported.

***Cannot (yet) use PPPM with triclinic box and kspace\_modify diff ad***

This feature is not yet supported.

***Cannot (yet) use PPPM with triclinic box and slab correction***

This feature is not yet supported.

***Cannot (yet) use kspace slab correction with long-range dipoles and non-neutral systems or per-atom energy***

This feature is not yet supported.

***Cannot (yet) use kspace\_modify diff ad with compute group/group***

This option is not yet supported.

***Cannot (yet) use kspace\_style ppm/stagger with triclinic systems***

This feature is not yet supported.

***Cannot (yet) use molecular templates with Kokkos***

Self-explanatory.

***Cannot (yet) use respa with Kokkos***

Self-explanatory.

***Cannot (yet) use rigid bodies with fix deform and Kokkos***

Self-explanatory.

***Cannot (yet) use rigid bodies with fix nh and Kokkos***

Self-explanatory.

***Cannot (yet) use single precision with MSM (remove -DFFT\_SINGLE from Makefile and re-compile)***

Single precision cannot be used with MSM.

***Cannot add atoms to fix move variable***

Atoms can not be added afterwards to this fix option.

***Cannot append atoms to a triclinic box***

The simulation box must be defined with edges aligned with the Cartesian axes.

***Cannot balance in z dimension for 2d simulation***

Self-explanatory.

***Cannot change box ortho/triclinic with certain fixes defined***

This is because those fixes store the shape of the box. You need to use unfix to discard the fix, change the box, then redefine a new fix.

***Cannot change box ortho/triclinic with dumps defined***

This is because some dumps store the shape of the box. You need to use undump to discard the dump, change the box, then redefine a new dump.

***Cannot change box tilt factors for orthogonal box***

Cannot use tilt factors unless the simulation box is non-orthogonal.

***Cannot change box to orthogonal when tilt is non-zero***

Self-explanatory.

***Cannot change box z boundary to non-periodic for a 2d simulation***

Self-explanatory.

***Cannot change dump\_modify every for dump dcd***

The frequency of writing dump dcd snapshots cannot be changed.

***Cannot change dump\_modify every for dump xtc***

The frequency of writing dump xtc snapshots cannot be changed.

***Cannot change timestep once fix srd is setup***

This is because various SRD properties depend on the timestep size.

***Cannot change timestep with fix pour***

This is because fix pour pre-computes the time delay for particles to fall out of the insertion volume due to gravity.

***Cannot change to comm\_style brick from tiled layout***

Self-explanatory.

***Cannot change\_box after reading restart file with per-atom info***

This is because the restart file info cannot be migrated with the atoms. You can get around this by performing a 0-timestep run which will assign the restart file info to actual atoms.

***Cannot change\_box in xz or yz for 2d simulation***

Self-explanatory.

***Cannot change\_box in z dimension for 2d simulation***

Self-explanatory.

***Cannot clear group all***

This operation is not allowed.

***Cannot close restart file - MPI error: %s***

This error was generated by MPI when reading/writing an MPI-IO restart file.

***Cannot compute initial g\_ewald\_disp***

LAMMPS failed to compute an initial guess for the PPPM\_disp g\_ewald\_6 factor that partitions the computation between real space and k-space for Dispersion interactions.

***Cannot create an atom map unless atoms have IDs***

The simulation requires a mapping from global atom IDs to local atoms, but the atoms that have been defined have no IDs.

***Cannot create atoms with undefined lattice***

Must use the lattice command before using the create\_atoms command.

***Cannot create/grow a vector/array of pointers for %s***

LAMMPS code is making an illegal call to the templated memory allocators, to create a vector or array of pointers.

***Cannot create \_atoms after reading restart file with per-atom info***

The per-atom info was stored to be used when by a fix that you may re-define. If you add atoms before re-defining the fix, then there will not be a correct amount of per-atom info.

***Cannot create \_box after simulation box is defined***

A simulation box can only be defined once.

***Cannot currently use pair reax with pair hybrid***

This is not yet supported.

***Cannot currently use ppm/gpu with fix balance.***

Self-explanatory.

***Cannot delete group all***

Self-explanatory.

***Cannot delete group currently used by a compute***

Self-explanatory.

***Cannot delete group currently used by a dump***

Self-explanatory.

***Cannot delete group currently used by a fix***

Self-explanatory.

***Cannot delete group currently used by atom\_modify first***

Self-explanatory.

***Cannot delete \_atoms bond yes for non-molecular systems***

Self-explanatory.

***Cannot displace \_atoms after reading restart file with per-atom info***

This is because the restart file info cannot be migrated with the atoms. You can get around this by performing a 0-timestep run which will assign the restart file info to actual atoms.

***Cannot do GCMC on atoms in atom\_modify first group***

This is a restriction due to the way atoms are organized in a list to enable the atom\_modify first command.

***Cannot do atom/swap on atoms in atom\_modify first group***

This is a restriction due to the way atoms are organized in a list to enable the atom\_modify first command.

***Cannot dump sort on atom IDs with no atom IDs defined***

Self-explanatory.

***Cannot dump sort when multiple dump files are written***

In this mode, each processor dumps its atoms to a file, so no sorting is allowed.

***Cannot embed Python when also extending Python with LAMMPS***

When running LAMMPS via Python through the LAMMPS library interface you cannot also use the input script python command.

***Cannot evaporate atoms in atom\_modify first group***

This is a restriction due to the way atoms are organized in a list to enable the atom\_modify first command.

***Cannot find create\_bonds group ID***

Self-explanatory.

***Cannot find delete\_bonds group ID***

Group ID used in the delete\_bonds command does not exist.

***Cannot find specified group ID for core particles***

Self-explanatory.

***Cannot find specified group ID for shell particles***

Self-explanatory.

***Cannot have both pair\_modify shift and tail set to yes***

These 2 options are contradictory.

***Cannot intersect groups using a dynamic group***

This operation is not allowed.

***Cannot mix molecular and molecule template atom styles***

Self-explanatory.

***Cannot open -reorder file***

Self-explanatory.

***Cannot open ADP potential file %s***

The specified ADP potential file cannot be opened. Check that the path and name are correct.

***Cannot open AIREBO potential file %s***

The specified AIREBO potential file cannot be opened. Check that the path and name are correct.

***Cannot open BOP potential file %s***

The specified BOP potential file cannot be opened. Check that the path and name are correct.

***Cannot open COMB potential file %s***

The specified COMB potential file cannot be opened. Check that the path and name are correct.

***Cannot open COMB3 lib.comb3 file***

The COMB3 library file cannot be opened. Check that the path and name are correct.

***Cannot open COMB3 potential file %s***

The specified COMB3 potential file cannot be opened. Check that the path and name are correct.

***Cannot open EAM potential file %s***

The specified EAM potential file cannot be opened. Check that the path and name are correct.

***Cannot open EIM potential file %s***

The specified EIM potential file cannot be opened. Check that the path and name are correct.

***Cannot open LCBOP potential file %s***

The specified LCBOP potential file cannot be opened. Check that the path and name are correct.

***Cannot open MEAM potential file %s***

The specified MEAM potential file cannot be opened. Check that the path and name are correct.

***Cannot open SNAP coefficient file %s***

The specified SNAP coefficient file cannot be opened. Check that the path and name are correct.



***Cannot open SNAP parameter file %s***

The specified SNAP parameter file cannot be opened. Check that the path and name are correct.

***Cannot open Stillinger-Weber potential file %s***

The specified SW potential file cannot be opened. Check that the path and name are correct.

***Cannot open Tersoff potential file %s***

The specified potential file cannot be opened. Check that the path and name are correct.

***Cannot open Vashishta potential file %s***

The specified Vashishta potential file cannot be opened. Check that the path and name are correct.

***Cannot open balance output file***

Self-explanatory.

***Cannot open coul/streitz potential file %s***

The specified coul/streitz potential file cannot be opened. Check that the path and name are correct.

***Cannot open custom file***

Self-explanatory.

***Cannot open data file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open dir to search for restart file***

Using a "\*" in the name of the restart file will open the current directory to search for matching file names.

***Cannot open dump file***

Self-explanatory.

***Cannot open dump file %s***

The output file for the dump command cannot be opened. Check that the path and name are correct.

***Cannot open file %s***

The specified file cannot be opened. Check that the path and name are correct. If the file is a compressed file, also check that the gzip executable can be found and run.

***Cannot open file variable file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix ave/chunk file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix ave/correlate file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix ave/histo file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix ave/time file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix balance output file***

Self-explanatory.

***Cannot open fix poems file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix print file %s***

The output file generated by the fix print command cannot be opened

***Cannot open fix qeq parameter file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix qeq/comb file %s***

The output file for the fix qeq/combs command cannot be opened. Check that the path and name are correct.

***Cannot open fix reax/bonds file %s***

The output file for the fix reax/bonds command cannot be opened. Check that the path and name are correct.

***Cannot open fix rigid infile %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix rigid restart file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix rigid/small infile %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open fix tmd file %s***

The output file for the fix tmd command cannot be opened. Check that the path and name are correct.

***Cannot open fix ttm file %s***

The output file for the fix ttm command cannot be opened. Check that the path and name are correct.

***Cannot open gzipped file***

LAMMPS was compiled without support for reading and writing gzipped files through a pipeline to the gzip program with -DLAMMPS\_GZIP.

***Cannot open input script %s***

Self-explanatory.

***Cannot open log.cite file***

This file is created when you use some LAMMPS features, to indicate what paper you should cite on behalf of those who implemented the feature. Check that you have write privileges into the directory you are running in.

***Cannot open log.lammps for writing***

The default LAMMPS log file cannot be opened. Check that the directory you are running in allows for files to be created.

***Cannot open logfile***

The LAMMPS log file named in a command-line argument cannot be opened. Check that the path and name are correct.

***Cannot open logfile %s***

The LAMMPS log file specified in the input script cannot be opened. Check that the path and name are correct.

***Cannot open molecule file %s***

The specified file cannot be opened. Check that the path and name are correct.

***Cannot open nb3b/harmonic potential file %s***

The specified potential file cannot be opened. Check that the path and name are correct.

***Cannot open pair\_write file***

The specified output file for pair energies and forces cannot be opened. Check that the path and name are correct.

***Cannot open polymorphic potential file %s***

The specified polymorphic potential file cannot be opened. Check that the path and name are correct.

***Cannot open print file %s***

Self-explanatory.

***Cannot open processors output file***

Self-explanatory.

***Cannot open restart file %s***

Self-explanatory.

***Cannot open restart file for reading - MPI error: %s***

This error was generated by MPI when reading/writing an MPI-IO restart file.

***Cannot open restart file for writing - MPI error: %s***

This error was generated by MPI when reading/writing an MPI-IO restart file.

***Cannot open screen file***

The screen file specified as a command-line argument cannot be opened. Check that the directory you are running in allows for files to be created.

***Cannot open temporary file for world counter.***

Self-explanatory.

***Cannot open universe log file***

For a multi-partition run, the master log file cannot be opened. Check that the directory you are running in allows for files to be created.

***Cannot open universe screen file***

For a multi-partition run, the master screen file cannot be opened. Check that the directory you are running in allows for files to be created.

***Cannot read from restart file - MPI error: %s***

This error was generated by MPI when reading/writing an MPI-IO restart file.

***Cannot read\_data without add keyword after simulation box is defined***

Self-explanatory.

***Cannot read\_restart after simulation box is defined***

The read\_restart command cannot be used after a read\_data, read\_restart, or create\_box command.

***Cannot redefine variable as a different style***

An equal-style variable can be re-defined but only if it was originally an equal-style variable.

***Cannot replicate 2d simulation in z dimension***

The replicate command cannot replicate a 2d simulation in the z dimension.

***Cannot replicate with fixes that store atom quantities***

Either fixes are defined that create and store atom-based vectors or a restart file was read which included atom-based vectors for fixes. The replicate command cannot duplicate that information for new atoms. You should use the replicate command before fixes are applied to the system.

***Cannot reset timestep with a dynamic region defined***

Dynamic regions (see the region command) have a time dependence. Thus you cannot change the timestep when one or more of these are defined.

***Cannot reset timestep with a time-dependent fix defined***

You cannot reset the timestep when a fix that keeps track of elapsed time is in place.

***Cannot run 2d simulation with non-periodic Z dimension***

Use the boundary command to make the z dimension periodic in order to run a 2d simulation.

***Cannot set bond topology types for atom style template***

The bond, angle, etc types cannot be changed for this atom style since they are static settings in the molecule template files.

***Cannot set both respa pair and inner/middle/outer***

In the rRESPA integrator, you must compute pairwise potentials either all together (pair), or in pieces (inner/middle/outer). You can't do both.

***Cannot set cutoff/multi before simulation box is defined***

Self-explanatory.

***Cannot set dpd/theta for this atom style***

Self-explanatory.

***Cannot set dump\_modify flush for dump xtc***

Self-explanatory.

***Cannot set mass for this atom style***

This atom style does not support mass settings for each atom type. Instead they are defined on a per-atom basis in the data file.

***Cannot set meso/cv for this atom style***

Self-explanatory.

***Cannot set meso/e for this atom style***

Self-explanatory.

***Cannot set meso/rho for this atom style***

Self-explanatory.

***Cannot set non-zero image flag for non-periodic dimension***

Self-explanatory.

***Cannot set non-zero z velocity for 2d simulation***

Self-explanatory.

***Cannot set quaternion for atom that has none***

Self-explanatory.

***Cannot set quaternion with xy components for 2d system***

Self-explanatory.

***Cannot set respa hybrid and any of pair/inner/middle/outer***

In the rRESPA integrator, you must compute pairwise potentials either all together (pair), with different cutoff regions (inner/middle/outer), or per hybrid sub-style (hybrid). You cannot mix those.

***Cannot set respa middle without inner/outer***

In the rRESPA integrator, you must define both a inner and outer setting in order to use a middle setting.

***Cannot set restart file size - MPI error: %s***

This error was generated by MPI when reading/writing an MPI-IO restart file.

***Cannot set smd/contact/radius for this atom style***

Self-explanatory.

***Cannot set smd/mass/density for this atom style***

Self-explanatory.

***Cannot set temperature for fix rigid/nph***

The temp keyword cannot be specified.

***Cannot set theta for atom that is not a line***

Self-explanatory.

***Cannot set this attribute for this atom style***

The attribute being set does not exist for the defined atom style.

***Cannot set variable z velocity for 2d simulation***

Self-explanatory.

***Cannot skew triclinic box in z for 2d simulation***

Self-explanatory.

***Cannot subtract groups using a dynamic group***

This operation is not allowed.

**Cannot union groups using a dynamic group**

This operation is not allowed.

**Cannot use -kokkos on without KOKKOS installed**

Self-explanatory.

**Cannot use -reorder after -partition**

Self-explanatory. See page discussion of command-line switches.

**Cannot use Ewald with 2d simulation**

The kspace style ewald cannot be used in 2d simulations. You can use 2d Ewald in a 3d simulation; see the kspace\_modify command.

**Cannot use Ewald/disp solver on system with no charge, dipole, or LJ particles**

No atoms in system have a non-zero charge or dipole, or are LJ particles. Change charges/dipoles or change options of the kspace solver/pair style.

**Cannot use EwaldDisp with 2d simulation**

This is a current restriction of this command.

**Cannot use Kokkos pair style with rRESPA inner/middle**

Self-explanatory.

**Cannot use NEB unless atom map exists**

Use the atom\_modify command to create an atom map.

**Cannot use NEB with a single replica**

Self-explanatory.

**Cannot use NEB with atom\_modify sort enabled**

This is current restriction for NEB implemented in LAMMPS.

**Cannot use PPPM with 2d simulation**

The kspace style pppm cannot be used in 2d simulations. You can use 2d PPPM in a 3d simulation; see the kspace\_modify command.

**Cannot use PPPMDisp with 2d simulation**

The kspace style pppm/disp cannot be used in 2d simulations. You can use 2d pppm/disp in a 3d simulation; see the kspace\_modify command.

**Cannot use PRD with a changing box**

The current box dimensions are not copied between replicas

**Cannot use PRD with a time-dependent fix defined**

PRD alters the timestep in ways that will mess up these fixes.

**Cannot use PRD with a time-dependent region defined**

PRD alters the timestep in ways that will mess up these regions.

**Cannot use PRD with atom\_modify sort enabled**

This is a current restriction of PRD. You must turn off sorting, which is enabled by default, via the atom\_modify command.

**Cannot use PRD with multi-processor replicas unless atom map exists**

Use the atom\_modify command to create an atom map.

**Cannot use TAD unless atom map exists for NEB**

See atom\_modify map command to set this.

**Cannot use TAD with a single replica for NEB**

NEB requires multiple replicas.

***Cannot use TAD with atom\_modify sort enabled for NEB***

This is a current restriction of NEB.

***Cannot use a damped dynamics min style with fix box/relax***

This is a current restriction in LAMMPS. Use another minimizer style.

***Cannot use a damped dynamics min style with per-atom DOF***

This is a current restriction in LAMMPS. Use another minimizer style.

***Cannot use append/atoms in periodic dimension***

The boundary style of the face where atoms are added can not be of type p (periodic).

***Cannot use atomfile-style variable unless atom map exists***

Self-explanatory. See the atom\_modify command to create a map.

***Cannot use both com and bias with compute temp/chunk***

Self-explanatory.

***Cannot use chosen neighbor list style with buck/coul/cut/kk***

Self-explanatory.

***Cannot use chosen neighbor list style with buck/coul/long/kk***

Self-explanatory.

***Cannot use chosen neighbor list style with buck/kk***

That style is not supported by Kokkos.

***Cannot use chosen neighbor list style with coul/cut/kk***

That style is not supported by Kokkos.

***Cannot use chosen neighbor list style with coul/debye/kk***

Self-explanatory.

***Cannot use chosen neighbor list style with coul/dsf/kk***

That style is not supported by Kokkos.

***Cannot use chosen neighbor list style with coul/wolf/kk***

That style is not supported by Kokkos.

***Cannot use chosen neighbor list style with lj/charmm/coul/charmm/implicit/kk***

Self-explanatory.

***Cannot use chosen neighbor list style with lj/charmm/coul/charmm/kk***

Self-explanatory.

***Cannot use chosen neighbor list style with lj/charmm/coul/long/kk***

Self-explanatory.

***Cannot use chosen neighbor list style with lj/class2/coul/cut/kk***

Self-explanatory.

***Cannot use chosen neighbor list style with lj/class2/coul/long/kk***

Self-explanatory.

***Cannot use chosen neighbor list style with lj/class2/kk***

Self-explanatory.

***Cannot use chosen neighbor list style with lj/cut/coul/cut/kk***

That style is not supported by Kokkos.

***Cannot use chosen neighbor list style with lj/cut/coul/debye/kk***

Self-explanatory.

**Cannot use chosen neighbor list style with lj/cut/coul/long/kk**

That style is not supported by Kokkos.

**Cannot use chosen neighbor list style with lj/cut/kk**

That style is not supported by Kokkos.

**Cannot use chosen neighbor list style with lj/expand/kk**

Self-explanatory.

**Cannot use chosen neighbor list style with lj/gromacs/coul/gromacs/kk**

Self-explanatory.

**Cannot use chosen neighbor list style with lj/gromacs/kk**

Self-explanatory.

**Cannot use chosen neighbor list style with lj/spica/kk**

That style is not supported by Kokkos.

**Cannot use chosen neighbor list style with pair eam/kk**

That style is not supported by Kokkos.

**Cannot use chosen neighbor list style with pair eam/kk/alloy**

Self-explanatory.

**Cannot use chosen neighbor list style with pair eam/kk/fs**

Self-explanatory.

**Cannot use chosen neighbor list style with pair sw/kk**

Self-explanatory.

**Cannot use chosen neighbor list style with tersoff/kk**

Self-explanatory.

**Cannot use chosen neighbor list style with tersoff/zbl/kk**

Self-explanatory.

**Cannot use compute chunk/atom bin z for 2d model**

Self-explanatory.

**Cannot use compute cluster/atom unless atoms have IDs**

Atom IDs are used to identify clusters.

**Cannot use create\_atoms rotate unless single style**

Self-explanatory.

**Cannot use create\_bonds unless atoms have IDs**

This command requires a mapping from global atom IDs to local atoms, but the atoms that have been defined have no IDs.

**Cannot use create\_bonds with non-molecular system**

Self-explanatory.

**Cannot use cwiggle in variable formula between runs**

This is a function of elapsed time.

**Cannot use delete\_atoms bond yes with atom\_style template**

This is because the bonds for that atom style are hardwired in the molecule template.

**Cannot use delete\_atoms unless atoms have IDs**

Your atoms do not have IDs, so the delete\_atoms command cannot be used.

**Cannot use delete\_bonds with non-molecular system**

Your choice of atom style does not have bonds.

***Cannot use dump\_modify fileper without % in dump file name***

Self-explanatory.

***Cannot use dump\_modify nfile without % in dump file name***

Self-explanatory.

***Cannot use dynamic group with fix adapt atom***

This is not yet supported.

***Cannot use fix TMD unless atom map exists***

Using this fix requires the ability to lookup an atom index, which is provided by an atom map. An atom map does not exist (by default) for non-molecular problems. Using the atom\_modify map command will force an atom map to be created.

***Cannot use fix bond/break with non-molecular systems***

Only systems with bonds that can be changed can be used. Atom\_style template does not qualify.

***Cannot use fix bond/create with non-molecular systems***

Only systems with bonds that can be changed can be used. Atom\_style template does not qualify.

***Cannot use fix bond/swap with non-molecular systems***

Only systems with bonds that can be changed can be used. Atom\_style template does not qualify.

***Cannot use fix box/relax on a 2nd non-periodic dimension***

When specifying an off-diagonal pressure component, the second of the two dimensions must be periodic. E.g. if the xy component is specified, then the y dimension must be periodic.

***Cannot use fix box/relax on a non-periodic dimension***

When specifying a diagonal pressure component, the dimension must be periodic.

***Cannot use fix box/relax with both relaxation and scaling on a tilt factor***

When specifying scaling on a tilt factor component, that component can not also be controlled by the barostat. E.g. if scalexy yes is specified and also keyword tri or xy, this is wrong.

***Cannot use fix box/relax with tilt factor scaling on a 2nd non-periodic dimension***

When specifying scaling on a tilt factor component, the second of the two dimensions must be periodic. E.g. if the xy component is specified, then the y dimension must be periodic.

***Cannot use fix deform on a shrink-wrapped boundary***

The x, y, z options cannot be applied to shrink-wrapped dimensions.

***Cannot use fix deform tilt on a shrink-wrapped 2nd dim***

This is because the shrink-wrapping will change the value of the strain implied by the tilt factor.

***Cannot use fix deform trape on a box with zero tilt***

The trape style alters the current strain.

***Cannot use fix deposit rigid and not molecule***

Self-explanatory.

***Cannot use fix deposit rigid and shake***

These two attributes are conflicting.

***Cannot use fix deposit shake and not molecule***

Self-explanatory.

***Cannot use fix enforce2d with 3d simulation***

Self-explanatory.

***Cannot use fix gcmc in a 2d simulation***

Fix gcmc is set up to run in 3d only. No 2d simulations with fix gcmc are allowed.



**Cannot use fix gcmc shake and not molecule**

Self-explanatory.

**Cannot use fix msst without per-type mass defined**

Self-explanatory.

**Cannot use fix npt and fix deform on same component of stress tensor**

This would be changing the same box dimension twice.

**Cannot use fix nvt/npt/nph on a 2nd non-periodic dimension**

When specifying an off-diagonal pressure component, the second of the two dimensions must be periodic. E.g. if the xy component is specified, then the y dimension must be periodic.

**Cannot use fix nvt/npt/nph on a non-periodic dimension**

When specifying a diagonal pressure component, the dimension must be periodic.

**Cannot use fix nvt/npt/nph with both xy dynamics and xy scaling**

Self-explanatory.

**Cannot use fix nvt/npt/nph with both xz dynamics and xz scaling**

Self-explanatory.

**Cannot use fix nvt/npt/nph with both yz dynamics and yz scaling**

Self-explanatory.

**Cannot use fix nvt/npt/nph with xy scaling when y is non-periodic dimension**

The second dimension in the barostated tilt factor must be periodic.

**Cannot use fix nvt/npt/nph with xz scaling when z is non-periodic dimension**

The second dimension in the barostated tilt factor must be periodic.

**Cannot use fix nvt/npt/nph with yz scaling when z is non-periodic dimension**

The second dimension in the barostated tilt factor must be periodic.

**Cannot use fix pour rigid and not molecule**

Self-explanatory.

**Cannot use fix pour rigid and shake**

These two attributes are conflicting.

**Cannot use fix pour shake and not molecule**

Self-explanatory.

**Cannot use fix pour with triclinic box**

This option is not yet supported.

**Cannot use fix press/berendsen and fix deform on same component of stress tensor**

These commands both change the box size/shape, so you cannot use both together.

**Cannot use fix press/berendsen on a non-periodic dimension**

Self-explanatory.

**Cannot use fix press/berendsen with triclinic box**

Self-explanatory.

**Cannot use fix reax/bonds without pair\_style reax**

Self-explanatory.

**Cannot use fix rigid npt/nph and fix deform on same component of stress tensor**

This would be changing the same box dimension twice.

**Cannot use fix rigid npt/nph on a non-periodic dimension**

When specifying a diagonal pressure component, the dimension must be periodic.

***Cannot use fix rigid/small npt/nph on a non-periodic dimension***

When specifying a diagonal pressure component, the dimension must be periodic.

***Cannot use fix shake with non-molecular system***

Your choice of atom style does not have bonds.

***Cannot use fix ttm with 2d simulation***

This is a current restriction of this fix due to the grid it creates.

***Cannot use fix ttm with triclinic box***

This is a current restriction of this fix due to the grid it creates.

***Cannot use fix tune/kSPACE without a kSPACE style***

Self-explanatory.

***Cannot use fix tune/kSPACE without a pair style***

This fix (tune/kSPACE) can only be used when a pair style has been specified.

***Cannot use fix wall in periodic dimension***

Self-explanatory.

***Cannot use fix wall zlo/zhi for a 2d simulation***

Self-explanatory.

***Cannot use fix wall/reflect in periodic dimension***

Self-explanatory.

***Cannot use fix wall/reflect zlo/zhi for a 2d simulation***

Self-explanatory.

***Cannot use fix wall/srd in periodic dimension***

Self-explanatory.

***Cannot use fix wall/srd more than once***

Nor is there a need to since multiple walls can be specified in one command.

***Cannot use fix wall/srd without fix srd***

Self-explanatory.

***Cannot use fix wall/srd zlo/zhi for a 2d simulation***

Self-explanatory.

***Cannot use fix\_deposit unless atoms have IDs***

Self-explanatory.

***Cannot use fix\_pour unless atoms have IDs***

Self-explanatory.

***Cannot use include command within an if command***

Self-explanatory.

***Cannot use lines with fix srd unless overlap is set***

This is because line segments are connected to each other.

***Cannot use multiple fix wall commands with pair brownian***

Self-explanatory.

***Cannot use multiple fix wall commands with pair lubricate***

Self-explanatory.

***Cannot use multiple fix wall commands with pair lubricate/poly***

Self-explanatory.

***Cannot use multiple fix wall commands with pair lubricateU***

Self-explanatory.

***Cannot use neigh\_modify exclude with GPU neighbor builds***

This is a current limitation of the GPU implementation in LAMMPS.

***Cannot use neighbor bins - box size << cutoff***

Too many neighbor bins will be created. This typically happens when the simulation box is very small in some dimension, compared to the neighbor cutoff. Use the “nsq” style instead of “bin” style.

***Cannot use newton pair with beck/gpu pair style***

Self-explanatory.

***Cannot use newton pair with born/coul/long/gpu pair style***

Self-explanatory.

***Cannot use newton pair with born/coul/wolf/gpu pair style***

Self-explanatory.

***Cannot use newton pair with born/gpu pair style***

Self-explanatory.

***Cannot use newton pair with buck/coul/cut/gpu pair style***

Self-explanatory.

***Cannot use newton pair with buck/coul/long/gpu pair style***

Self-explanatory.

***Cannot use newton pair with buck/gpu pair style***

Self-explanatory.

***Cannot use newton pair with colloid/gpu pair style***

Self-explanatory.

***Cannot use newton pair with coul/cut/gpu pair style***

Self-explanatory.

***Cannot use newton pair with coul/debye/gpu pair style***

Self-explanatory.

***Cannot use newton pair with coul/dsf/gpu pair style***

Self-explanatory.

***Cannot use newton pair with coul/long/gpu pair style***

Self-explanatory.

***Cannot use newton pair with dipole/cut/gpu pair style***

Self-explanatory.

***Cannot use newton pair with dipole/sf/gpu pair style***

Self-explanatory.

***Cannot use newton pair with dpd/gpu pair style***

Self-explanatory.

***Cannot use newton pair with dpd/tstat/gpu pair style***

Self-explanatory.

***Cannot use newton pair with eam/alloy/gpu pair style***

Self-explanatory.

***Cannot use newton pair with eam/fs/gpu pair style***

Self-explanatory.

*Cannot use newton pair with eam/gpu pair style*

Self-explanatory.

*Cannot use newton pair with gauss/gpu pair style*

Self-explanatory.

*Cannot use newton pair with gayberne/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/charmm/coul/long/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/class2/coul/long/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/class2/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/cubic/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/cut/coul/cut/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/cut/coul/debye/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/cut/coul/dsf/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/cut/coul/long/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/cut/coul/msm/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/cut/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/expand/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/gromacs/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/spica/coul/long/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj/spica/gpu pair style*

Self-explanatory.

*Cannot use newton pair with lj96/cut/gpu pair style*

Self-explanatory.

*Cannot use newton pair with mie/cut/gpu pair style*

Self-explanatory.

*Cannot use newton pair with morse/gpu pair style*

Self-explanatory.

*Cannot use newton pair with resquared/gpu pair style*

Self-explanatory.

**Cannot use newton pair with soft/gpu pair style**

Self-explanatory.

**Cannot use newton pair with table/gpu pair style**

Self-explanatory.

**Cannot use newton pair with yukawa/colloid/gpu pair style**

Self-explanatory.

**Cannot use newton pair with yukawa/gpu pair style**

Self-explanatory.

**Cannot use newton pair with zbl/gpu pair style**

Self-explanatory.

**Cannot use non-zero forces in an energy minimization**

Fix setforce cannot be used in this manner. Use fix addforce instead.

**Cannot use non-periodic boundares with fix ttm**

This fix requires a fully periodic simulation box.

**Cannot use non-periodic boundaries with Ewald**

For kspace style ewald, all 3 dimensions must have periodic boundaries unless you use the kspace\_modify command to define a 2d slab with a non-periodic z dimension.

**Cannot use non-periodic boundaries with EwaldDisp**

For kspace style ewald/disp, all 3 dimensions must have periodic boundaries unless you use the kspace\_modify command to define a 2d slab with a non-periodic z dimension.

**Cannot use non-periodic boundaries with PPPM**

For kspace style pppm, all 3 dimensions must have periodic boundaries unless you use the kspace\_modify command to define a 2d slab with a non-periodic z dimension.

**Cannot use non-periodic boundaries with PPPMDisp**

For kspace style pppm/disp, all 3 dimensions must have periodic boundaries unless you use the kspace\_modify command to define a 2d slab with a non-periodic z dimension.

**Cannot use order greater than 8 with pppm/gpu.**

Self-explanatory.

**Cannot use package gpu neigh yes with triclinic box**

This is a current restriction in LAMMPS.

**Cannot use pair tail corrections with 2d simulations**

The correction factors are only currently defined for 3d systems.

**Cannot use processors part command without using partitions**

See the command-line -partition switch.

**Cannot use ramp in variable formula between runs**

This is because the ramp() function is time dependent.

**Cannot use read\_data add before simulation box is defined**

Self-explanatory.

**Cannot use read\_data extra with add flag**

Self-explanatory.

**Cannot use read\_data offset without add flag**

Self-explanatory.

**Cannot use read\_data shift without add flag**

Self-explanatory.

***Cannot use region INF or EDGE when box does not exist***

Regions that extend to the box boundaries can only be used after the create\_box command has been used.

***Cannot use set atom with no atom IDs defined***

Atom IDs are not defined, so they cannot be used to identify an atom.

***Cannot use set mol with no molecule IDs defined***

Self-explanatory.

***Cannot use swiggle in variable formula between runs***

This is a function of elapsed time.

***Cannot use tris with fix srd unless overlap is set***

This is because triangles are connected to each other.

***Cannot use variable energy with constant efield in fix efield***

LAMMPS computes the energy itself when the E-field is constant.

***Cannot use variable energy with constant force in fix addforce***

This is because for constant force, LAMMPS can compute the change in energy directly.

***Cannot use variable every setting for dump dcd***

The format of DCD dump files requires snapshots be output at a constant frequency.

***Cannot use variable every setting for dump xtc***

The format of this file requires snapshots at regular intervals.

***Cannot use vdisplace in variable formula between runs***

This is a function of elapsed time.

***Cannot use velocity bias command without temp keyword***

Self-explanatory.

***Cannot use velocity create loop all unless atoms have IDs***

Atoms in the simulation do not have IDs, so this style of velocity creation cannot be performed.

***Cannot use wall in periodic dimension***

Self-explanatory.

***Cannot use write\_restart fileper without % in restart file name***

Self-explanatory.

***Cannot use write\_restart nfile without % in restart file name***

Self-explanatory.

***Cannot wiggle and shear fix wall/gran***

Cannot specify both options at the same time.

***Cannot write to restart file - MPI error: %s***

This error was generated by MPI when reading/writing an MPI-IO restart file.

***Cannot yet use KSpace solver with grid with comm style tiled***

This is current restriction in LAMMPS.

***Cannot yet use comm\_style tiled with multi-mode comm***

Self-explanatory.

***Cannot yet use comm\_style tiled with triclinic box***

Self-explanatory.

***Cannot yet use compute tally with Kokkos***

This feature is not yet supported.

***Cannot yet use fix bond/break with this improper style***

This is a current restriction in LAMMPS.

***Cannot yet use fix bond/create with this improper style***

This is a current restriction in LAMMPS.

***Cannot yet use minimize with Kokkos***

This feature is not yet supported.

***Cannot yet use pair hybrid with Kokkos***

This feature is not yet supported.

***Cannot zero Langevin force of 0 atoms***

The group has zero atoms, so you cannot request its force be zeroed.

***Cannot zero gld force for zero atoms***

There are no atoms currently in the group.

***Cannot zero momentum of no atoms***

Self-explanatory.

***Change\_box command before simulation box is defined***

Self-explanatory.

***Change\_box volume used incorrectly***

The “dim volume” option must be used immediately following one or two settings for “dim1 ...” (and optionally “dim2 ...”) and must be for a different dimension, i.e.  $\text{dim} \neq \text{dim1}$  and  $\text{dim} \neq \text{dim2}$ .

***Chunk/atom compute does not exist for compute angmom/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for compute com/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for compute gyration/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for compute inertia/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for compute msd/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for compute omega/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for compute property/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for compute temp/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for compute torque/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for compute vcm/chunk***

Self-explanatory.

***Chunk/atom compute does not exist for fix ave/chunk***

Self-explanatory.

***Comm tiled invalid index in box drop brick***

Internal error check in comm\_style tiled which should not occur. Contact the developers.

***Comm tiled mis-match in box drop brick***

Internal error check in comm\_style tiled which should not occur. Contact the developers.

***Comm\_modify group != atom\_modify first group***

Self-explanatory.

***Communication cutoff for comm\_style tiled cannot exceed periodic box length***

Self-explanatory.

***Communication cutoff too small for SNAP micro load balancing***

This can happen if you change the neighbor skin after your pair\_style command or if your box dimensions grow during a run. You can set the cutoff explicitly via the comm\_modify cutoff command.

***Compute %s does not allow use of dynamic group***

Dynamic groups have not yet been enabled for this compute.

***Compute for fix pafi does not calculate a local array***

Self-explanatory.

***Compute for fix pafi must have 9 fields per atom***

Self-explanatory.

***Compute ID for compute chunk /atom does not exist***

Self-explanatory.

***Compute ID for compute chunk/atom does not exist***

Self-explanatory.

***Compute gyration ID does not exist for compute gyration/shape***

Self-explanatory. Provide a valid compute ID.

***Compute gyration/shape compute ID does not point to a gyration compute***

Self-explanatory. Provide an ID of a compute gyration command.

***Compute ID for compute reduce does not exist***

Self-explanatory.

***Compute ID for compute slice does not exist***

Self-explanatory.

***Compute ID for fix ave/atom does not exist***

Self-explanatory.

***Compute ID for fix ave/chunk does not exist***

Self-explanatory.

***Compute ID for fix ave/correlate does not exist***

Self-explanatory.

***Compute ID for fix ave/histo does not exist***

Self-explanatory.

***Compute ID for fix ave/time does not exist***

Self-explanatory.

***Compute ID for fix numdiff does not exist***

Self-explanatory.

***Compute ID for fix numdiff/virial does not exist***

Self-explanatory.

***Compute ID for fix store/state does not exist***

Self-explanatory.



***Compute ID for fix vector does not exist***

Self-explanatory.

***Compute ID must be alphanumeric or underscore characters***

Self-explanatory.

***Compute angle/local used when angles are not allowed***

The atom style does not support angles.

***Compute angmom/chunk does not use chunk/atom compute***

The style of the specified compute is not chunk/atom.

***Compute body/local requires atom style body***

Self-explanatory.

***Compute bond/local used when bonds are not allowed***

The atom style does not support bonds.

***Compute centro/atom requires a pair style be defined***

This is because the computation of the centro-symmetry values uses a pairwise neighbor list.

***Compute chunk/atom bin/cylinder radius is too large for periodic box***

Radius cannot be bigger than 1/2 of a non-axis periodic dimension.

***Compute chunk/atom bin/sphere radius is too large for periodic box***

Radius cannot be bigger than 1/2 of any periodic dimension.

***Compute chunk/atom compute array is accessed out-of-range***

The index for the array is out of bounds.

***Compute chunk/atom compute does not calculate a per-atom array***

Self-explanatory.

***Compute chunk/atom compute does not calculate a per-atom vector***

Self-explanatory.

***Compute chunk/atom compute does not calculate per-atom values***

Self-explanatory.

***Compute chunk/atom cylinder axis must be z for 2d***

Self-explanatory.

***Compute chunk/atom fix array is accessed out-of-range***

the index for the array is out of bounds.

***Compute chunk/atom fix does not calculate a per-atom array***

Self-explanatory.

***Compute chunk/atom fix does not calculate a per-atom vector***

Self-explanatory.

***Compute chunk/atom fix does not calculate per-atom values***

Self-explanatory.

***Compute chunk/atom for triclinic boxes requires units reduced***

Self-explanatory.

***Compute chunk/atom ids once but nchunk is not once***

You cannot assign chunks IDs to atom permanently if the number of chunks may change.

***Compute chunk/atom molecule for non-molecular system***

Self-explanatory.

***Compute chunk/atom sphere z origin must be 0.0 for 2d***

Self-explanatory.

***Compute chunk/atom stores no IDs for compute property/chunk***

It will only store IDs if its compress option is enabled.

***Compute chunk/atom stores no coord1 for compute property/chunk***

Only certain binning options for compute chunk/atom store coordinates.

***Compute chunk/atom stores no coord2 for compute property/chunk***

Only certain binning options for compute chunk/atom store coordinates.

***Compute chunk/atom stores no coord3 for compute property/chunk***

Only certain binning options for compute chunk/atom store coordinates.

***Compute chunk/atom variable is not atom-style variable***

Self-explanatory.

***Compute chunk/atom without bins cannot use discard mixed***

That discard option only applies to the binning styles.

***Compute cluster/atom cutoff is longer than pairwise cutoff***

Cannot identify clusters beyond cutoff.

***Compute cluster/atom requires a pair style be defined***

This is so that the pair style defines a cutoff distance which is used to find clusters.

***Compute cna/atom cutoff is longer than pairwise cutoff***

Self-explanatory.

***Compute cna/atom requires a pair style be defined***

Self-explanatory.

***Compute com/chunk does not use chunk/atom compute***

The style of the specified compute is not chunk/atom.

***Compute contact/atom requires a pair style be defined***

Self-explanatory.

***Compute contact/atom requires atom style sphere***

Self-explanatory.

***Compute coord/atom cutoff is longer than pairwise cutoff***

Cannot compute coordination at distances longer than the pair cutoff, since those atoms are not in the neighbor list.

***Compute coord/atom requires a pair style be defined***

Self-explanatory.

***Compute damage/atom requires peridynamic potential***

Damage is a Peridynamic-specific metric. It requires you to be running a Peridynamics simulation.

***Compute dihedral/local used when dihedrals are not allowed***

The atom style does not support dihedrals.

***Compute dilatation/atom cannot be used with this pair style***

Self-explanatory.

***Compute dilatation/atom requires Peridynamic pair style***

Self-explanatory.

***Compute does not allow an extra compute or fix to be reset***

This is an internal LAMMPS error. Please report it to the developers.

***Compute erotate/asphere requires atom style ellipsoid or line or tri***

Self-explanatory.

***Compute erotate/asphere requires extended particles***

This compute cannot be used with point particles.

***Compute erotate/rigid with non-rigid fix-ID***

Self-explanatory.

***Compute erotate/sphere requires atom style sphere***

Self-explanatory.

***Compute erotate/sphere/atom requires atom style sphere***

Self-explanatory.

***Compute event/displace has invalid fix event assigned***

This is an internal LAMMPS error. Please report it to the developers.

***Compute group/group group ID does not exist***

Self-explanatory.

***Compute gyration/chunk does not use chunk/atom compute***

The style of the specified compute is not chunk/atom.

***Compute heat/flux compute ID does not compute ke/atom***

Self-explanatory.

***Compute heat/flux compute ID does not compute pe/atom***

Self-explanatory.

***Compute heat/flux compute ID does not compute stress/atom***

Self-explanatory.

***Compute hexorder/atom cutoff is longer than pairwise cutoff***

Cannot compute order parameter beyond cutoff.

***Compute hexorder/atom requires a pair style be defined***

Self-explanatory.

***Compute improper/local used when impropers are not allowed***

The atom style does not support impropers.

***Compute inertia/chunk does not use chunk/atom compute***

The style of the specified compute is not chunk/atom.

***Compute ke/rigid with non-rigid fix-ID***

Self-explanatory.

***Compute msd/chunk does not use chunk/atom compute***

The style of the specified compute is not chunk/atom.

***Compute msd/chunk nchunk is not static***

This is required because the MSD cannot be computed consistently if the number of chunks is changing. Compute chunk/atom allows setting nchunk to be static.

***Compute nve/asphere requires atom style ellipsoid***

Self-explanatory.

***Compute nvt/nph/npt asphere requires atom style ellipsoid***

Self-explanatory.

***Compute nvt/nph/npt body requires atom style body***

Self-explanatory.

***Compute omega/chunk does not use chunk/atom compute***

The style of the specified compute is not chunk/atom.

***Compute orientorder/atom cutoff is longer than pairwise cutoff***

Cannot compute order parameter beyond cutoff.

***Compute orientorder/atom requires a pair style be defined***

Self-explanatory.

***Compute pair must use group all***

Pair styles accumulate energy on all atoms.

***Compute pe must use group all***

Energies computed by potentials (pair, bond, etc) are computed on all atoms.

***Compute plasticity/atom cannot be used with this pair style***

Self-explanatory.

***Compute plasticity/atom requires Peridynamic pair style***

Self-explanatory.

***Compute pressure must use group all***

Virial contributions computed by potentials (pair, bond, etc) are computed on all atoms.

***Compute pressure requires temperature ID to include kinetic energy***

The keflag cannot be used unless a temperature compute is provided.

***Compute pressure temperature ID does not compute temperature***

The compute ID assigned to a pressure computation must compute temperature.

***Compute property/atom floating point vector does not exist***

The command is accessing a vector added by the fix property/atom command, that does not exist.

***Compute property/atom for atom property that is not allocated***

Self-explanatory.

***Compute property/atom integer vector does not exist***

The command is accessing a vector added by the fix property/atom command, that does not exist.

***Compute property/chunk does not use chunk/atom compute***

The style of the specified compute is not chunk/atom.

***Compute property/local cannot use these inputs together***

Only inputs that generate the same number of datums can be used together. E.g. bond and angle quantities cannot be mixed.

***Compute property/local does not (yet) work with atom\_style template***

Self-explanatory.

***Compute property/local for property that is not allocated***

Self-explanatory.

***Compute rdf requires a pair style be defined***

Self-explanatory.

***Compute reduce compute array is accessed out-of-range***

An index for the array is out of bounds.

***Compute reduce compute calculates global values***

A compute that calculates peratom or local values is required.

***Compute reduce compute does not calculate a local array***

Self-explanatory.

***Compute reduce compute does not calculate a local vector***

Self-explanatory.

***Compute reduce compute does not calculate a per-atom array***

Self-explanatory.

***Compute reduce compute does not calculate a per-atom vector***

Self-explanatory.

***Compute reduce fix array is accessed out-of-range***

An index for the array is out of bounds.

***Compute reduce fix calculates global values***

A fix that calculates peratom or local values is required.

***Compute reduce fix does not calculate a local array***

Self-explanatory.

***Compute reduce fix does not calculate a local vector***

Self-explanatory.

***Compute reduce fix does not calculate a per-atom array***

Self-explanatory.

***Compute reduce fix does not calculate a per-atom vector***

Self-explanatory.

***Compute reduce replace requires min or max mode***

Self-explanatory.

***Compute reduce variable is not atom-style variable***

Self-explanatory.

***Compute slice compute array is accessed out-of-range***

An index for the array is out of bounds.

***Compute slice compute does not calculate a global array***

Self-explanatory.

***Compute slice compute does not calculate a global vector***

Self-explanatory.

***Compute slice compute does not calculate global vector or array***

Self-explanatory.

***Compute slice compute vector is accessed out-of-range***

The index for the vector is out of bounds.

***Compute slice fix array is accessed out-of-range***

An index for the array is out of bounds.

***Compute slice fix does not calculate a global array***

Self-explanatory.

***Compute slice fix does not calculate a global vector***

Self-explanatory.

***Compute slice fix does not calculate global vector or array***

Self-explanatory.

***Compute slice fix vector is accessed out-of-range***

The index for the vector is out of bounds.

- Compute sna/atom cutoff is longer than pairwise cutoff*  
Self-explanatory.
- Compute sna/atom requires a pair style be defined*  
Self-explanatory.
- Compute snad/atom cutoff is longer than pairwise cutoff*  
Self-explanatory.
- Compute snad/atom requires a pair style be defined*  
Self-explanatory.
- Compute snav/atom cutoff is longer than pairwise cutoff*  
Self-explanatory.
- Compute snav/atom requires a pair style be defined*  
Self-explanatory.
- Compute stress/atom temperature ID does not compute temperature*  
The specified compute must compute temperature.
- Compute temp/asphere requires atom style ellipsoid*  
Self-explanatory.
- Compute temp/asphere requires extended particles*  
This compute cannot be used with point particles.
- Compute temp/body requires atom style body*  
Self-explanatory.
- Compute temp/body requires bodies*  
This compute can only be applied to body particles.
- Compute temp/chunk does not use chunk/atom compute*  
The style of the specified compute is not chunk/atom.
- Compute temp/cs requires ghost atoms store velocity*  
Use the comm\_modify vel yes command to enable this.
- Compute temp/cs used when bonds are not allowed*  
This compute only works on pairs of bonded particles.
- Compute temp/partial cannot use vz for 2d systemx*  
Self-explanatory.
- Compute temp/profile cannot bin z for 2d systems*  
Self-explanatory.
- Compute temp/profile cannot use vz for 2d systemx*  
Self-explanatory.
- Compute temp/sphere requires atom style sphere*  
Self-explanatory.
- Compute ti kspace style does not exist*  
Self-explanatory.
- Compute ti pair style does not exist*  
Self-explanatory.
- Compute ti tail when pair style does not compute tail corrections*  
Self-explanatory.

***Compute torque/chunk does not use chunk/atom compute***

The style of the specified compute is not chunk/atom.

***Compute used in dump between runs is not current***

The compute was not invoked on the current timestep, therefore it cannot be used in a dump between runs.

***Compute used in variable between runs is not current***

Computes cannot be invoked by a variable in between runs. Thus they must have been evaluated on the last timestep of the previous run in order for their value(s) to be accessed. See the page for the variable command for more info.

***Compute used in variable thermo keyword between runs is not current***

Some thermo keywords rely on a compute to calculate their value(s). Computes cannot be invoked by a variable in between runs. Thus they must have been evaluated on the last timestep of the previous run in order for their value(s) to be accessed. See the page for the variable command for more info.

***Compute vcm/chunk does not use chunk/atom compute***

The style of the specified compute is not chunk/atom.

***Computed temperature for fix temp/berendsen cannot be 0.0***

Self-explanatory.

***Computed temperature for fix temp/rescale cannot be 0.0***

Cannot rescale the temperature to a new value if the current temperature is 0.0.

***Core/shell partner atom not found***

Could not find one of the atoms in the bond pair.

***Core/shell partners were not all found***

Could not find or more atoms in the bond pairs.

***Could not adjust g\_ewald\_6***

The Newton-Raphson solver failed to converge to a good value for g\_ewald. This error should not occur for typical problems. Please send an email to the developers.

***Could not compute g\_ewald***

The Newton-Raphson solver failed to converge to a good value for g\_ewald. This error should not occur for typical problems. Please send an email to the developers.

***Could not compute grid size***

The code is unable to compute a grid size consistent with the desired accuracy. This error should not occur for typical problems. Please send an email to the developers.

***Could not compute grid size for Coulomb interaction***

The code is unable to compute a grid size consistent with the desired accuracy. This error should not occur for typical problems. Please send an email to the developers.

***Could not compute grid size for Dispersion***

The code is unable to compute a grid size consistent with the desired accuracy. This error should not occur for typical problems. Please send an email to the developers.

***Could not create 3d FFT plan***

The FFT setup for the PPPM solver failed, typically due to lack of memory. This is an unusual error. Check the size of the FFT grid you are requesting.

***Could not create 3d grid of processors***

The specified constraints did not allow a Px by Py by Pz grid to be created where  $P_x * P_y * P_z = P$  = total number of processors.

***Could not create 3d remap plan***

The FFT setup in pppm failed.

***Could not create Python function arguments***

This is an internal Python error, possibly because the number of inputs to the function is too large.

***Could not create numa grid of processors***

The specified constraints did not allow this style of grid to be created. Usually this is because the total processor count is not a multiple of the cores/node or the user specified processor count is > 1 in one of the dimensions.

***Could not create twolevel 3d grid of processors***

The specified constraints did not allow this style of grid to be created.

***Could not evaluate Python function input variable***

Self-explanatory.

***Could not find Python function***

The provided Python code was run successfully, but it not define a callable function with the required name.

***Could not find atom\_modify first group ID***

Self-explanatory.

***Could not find change\_box group ID***

Group ID used in the change\_box command does not exist.

***Could not find compute ID for PRD***

Self-explanatory.

***Could not find compute ID for TAD***

Self-explanatory.

***Could not find compute ID for temperature bias***

Self-explanatory.

***Could not find compute ID to delete***

Self-explanatory.

***Could not find compute displace/atom fix ID***

Self-explanatory.

***Could not find compute event/displace fix ID***

Self-explanatory.

***Could not find compute group ID***

Self-explanatory.

***Could not find compute heat/flux compute ID***

Self-explanatory.

***Could not find compute msd fix ID***

Self-explanatory.

***Could not find compute msd/chunk fix ID***

The compute creates an internal fix, which has been deleted.

***Could not find compute pressure temperature ID***

The compute ID for calculating temperature does not exist.

***Could not find compute stress/atom temperature ID***

Self-explanatory.

***Could not find compute vacf fix ID***

Self-explanatory.

***Could not find compute/voronoi surface group ID***

Self-explanatory.



**Could not find compute\_modify ID**

Self-explanatory.

**Could not find custom per-atom property ID**

Self-explanatory.

**Could not find delete\_atoms group ID**

Group ID used in the delete\_atoms command does not exist.

**Could not find delete\_atoms region ID**

Region ID used in the delete\_atoms command does not exist.

**Could not find displace\_atoms group ID**

Group ID used in the displace\_atoms command does not exist.

**Could not find dump custom compute ID**

Self-explanatory.

**Could not find dump custom fix ID**

Self-explanatory.

**Could not find dump custom variable name**

Self-explanatory.

**Could not find dump group ID**

A group ID used in the dump command does not exist.

**Could not find dump local compute ID**

Self-explanatory.

**Could not find dump local fix ID**

Self-explanatory.

**Could not find dump modify compute ID**

Self-explanatory.

**Could not find dump modify custom atom floating point property ID**

Self-explanatory.

**Could not find dump modify custom atom integer property ID**

Self-explanatory.

**Could not find dump modify fix ID**

Self-explanatory.

**Could not find dump modify variable name**

Self-explanatory.

**Could not find fix ID to delete**

Self-explanatory.

**Could not find fix adapt storage fix ID**

This should not happen unless you explicitly deleted a secondary fix that fix adapt created internally.

**Could not find fix halt variable name**

Self-explanatory.

**Could not find fix gcmc exclusion group ID**

Self-explanatory.

**Could not find fix gcmc rotation group ID**

Self-explanatory.

***Could not find fix group ID***

A group ID used in the fix command does not exist.

***Could not find fix msst compute ID***

Self-explanatory.

***Could not find fix poems group ID***

A group ID used in the fix poems command does not exist.

***Could not find fix recenter group ID***

A group ID used in the fix recenter command does not exist.

***Could not find fix rigid group ID***

A group ID used in the fix rigid command does not exist.

***Could not find fix srd group ID***

Self-explanatory.

***Could not find fix\_modify ID***

A fix ID used in the fix\_modify command does not exist.

***Could not find fix\_modify pressure ID***

The compute ID for computing pressure does not exist.

***Could not find fix\_modify temperature ID***

The compute ID for computing temperature does not exist.

***Could not find group clear group ID***

Self-explanatory.

***Could not find group delete group ID***

Self-explanatory.

***Could not find pair fix ID***

A fix is created internally by the pair style to store shear history information. You cannot delete it.

***Could not find set group ID***

Group ID specified in set command does not exist.

***Could not find specified fix gcmc group ID***

Self-explanatory.

***Could not find thermo compute ID***

Compute ID specified in thermo\_style command does not exist.

***Could not find thermo custom compute ID***

The compute ID needed by thermo style custom to compute a requested quantity does not exist.

***Could not find thermo custom fix ID***

The fix ID needed by thermo style custom to compute a requested quantity does not exist.

***Could not find thermo custom variable name***

Self-explanatory.

***Could not find thermo fix ID***

Fix ID specified in thermo\_style command does not exist.

***Could not find thermo variable name***

Self-explanatory.

***Could not find thermo\_modify pressure ID***

The compute ID needed by thermo style custom to compute pressure does not exist.

***Could not find thermo\_modify temperature ID***

The compute ID needed by thermo style custom to compute temperature does not exist.

***Could not find undump ID***

A dump ID used in the undump command does not exist.

***Could not find velocity group ID***

A group ID used in the velocity command does not exist.

***Could not find velocity temperature ID***

The compute ID needed by the velocity command to compute temperature does not exist.

***Could not find/initialize a specified accelerator device***

Could not initialize at least one of the devices specified for the gpu package

***Could not grab element entry from EIM potential file***

Self-explanatory

***Could not grab global entry from EIM potential file***

Self-explanatory.

***Could not grab pair entry from EIM potential file***

Self-explanatory.

***Could not initialize embedded Python***

The main module in Python was not accessible.

***Could not open Python file***

The specified file of Python code cannot be opened. Check that the path and name are correct.

***Could not process Python file***

The Python code in the specified file was not run successfully by Python, probably due to errors in the Python code.

***Could not process Python string***

The Python code in the here string was not run successfully by Python, probably due to errors in the Python code.

***Coulomb PPPMDisp order has been reduced below minorder***

The default minimum order is 2. This can be reset by the kspace\_modify minorder command.

***Coulombic cutoff not supported in pair\_style buck/long/coul/coul***

Must use long-range Coulombic interactions.

***Coulombic cutoff not supported in pair\_style lj/long/coul/long***

Must use long-range Coulombic interactions.

***Coulombic cutoff not supported in pair\_style lj/long/tip4p/long***

Must use long-range Coulombic interactions.

***Coulombic cutoffs of pair hybrid sub-styles do not match***

If using a Kspace solver, all Coulombic cutoffs of long pair styles must be the same.

***Coulombic cut not supported in pair\_style lj/long/dipole/long***

Must use long-range Coulombic interactions.

***Cound not find dump\_modify ID***

Self-explanatory.

***Create\_atoms command before simulation box is defined***

The create\_atoms command cannot be used before a read\_data, read\_restart, or create\_box command.

***Create\_atoms molecule has atom IDs, but system does not***

The atom\_style id command can be used to force atom IDs to be stored.

***Create\_atoms molecule must have atom types***

The defined molecule does not specify atom types.

***Create\_atoms molecule must have coordinates***

The defined molecule does not specify coordinates.

***Create\_atoms region ID does not exist***

A region ID used in the create\_atoms command does not exist.

***Create\_bonds command before simulation box is defined***

Self-explanatory.

***Create\_bonds command requires no kspace\_style be defined***

This is so that atom pairs that are already bonded to not appear in the neighbor list.

***Create\_bonds command requires special\_bonds 1-2 weights be 0.0***

This is so that atom pairs that are already bonded to not appear in the neighbor list.

***Create\_bonds max distance > neighbor cutoff***

Can only create bonds for atom pairs that will be in neighbor list.

***Create\_bonds requires a pair style be defined***

Self-explanatory.

***Create\_box region ID does not exist***

Self-explanatory.

***Create\_box region does not support a bounding box***

Not all regions represent bounded volumes. You cannot use such a region with the create\_box command.

***Custom floating point vector for fix store/state does not exist***

The command is accessing a vector added by the fix property/atom command, that does not exist.

***Custom integer vector for fix store/state does not exist***

The command is accessing a vector added by the fix property/atom command, that does not exist.

***Custom per-atom property ID is not floating point***

Self-explanatory.

***Custom per-atom property ID is not integer***

Self-explanatory.

***Cut-offs missing in pair\_style lj/long/dipole/long***

Self-explanatory.

***Cutoffs missing in pair\_style buck/long/coul/long***

Self-explanatory.

***Cutoffs missing in pair\_style lj/long/coul/long***

Self-explanatory.

***Cyclic loop in joint connections***

Fix poems cannot (yet) work with coupled bodies whose joints connect the bodies in a ring (or cycle).

***Degenerate lattice primitive vectors***

Invalid set of 3 lattice vectors for lattice command.

***Delete region ID does not exist***

Self-explanatory.

***Delete\_atoms command before simulation box is defined***

The delete\_atoms command cannot be used before a read\_data, read\_restart, or create\_box command.

***Delete\_atoms cutoff > max neighbor cutoff***

Can only delete atoms in atom pairs that will be in neighbor list.

***Delete\_atoms mol yes requires atom attribute molecule***

Cannot use this option with a non-molecular system.

***Delete\_atoms requires a pair style be defined***

This is because atom deletion within a cutoff uses a pairwise neighbor list.

***Delete\_bonds command before simulation box is defined***

The delete\_bonds command cannot be used before a read\_data, read\_restart, or create\_box command.

***Delete\_bonds command with no atoms existing***

No atoms are yet defined so the delete\_bonds command cannot be used.

***Deposition region extends outside simulation box***

Self-explanatory.

***Did not assign all atoms correctly***

Atoms read in from a data file were not assigned correctly to processors. This is likely due to some atom coordinates being outside a non-periodic simulation box.

***Did not assign all restart atoms correctly***

Atoms read in from the restart file were not assigned correctly to processors. This is likely due to some atom coordinates being outside a non-periodic simulation box. Normally this should not happen. You may wish to use the “remap” option on the read\_restart command to see if this helps.

***Did not find all elements in MEAM library file***

Some requested elements were not found in the MEAM file. Check spelling etc.

***Did not find fix shake partner info***

Could not find bond partners implied by fix shake command. This error can be triggered if the delete\_bonds command was used before fix shake, and it removed bonds without resetting the 1-2, 1-3, 1-4 weighting list via the special keyword.

***Did not find keyword in table file***

Keyword used in pair\_coeff command was not found in table file.

***Did not set pressure for fix rigid/nph***

The press keyword must be specified.

***Did not set temp for fix rigid/nvt/small***

Self-explanatory.

***Did not set temp or press for fix rigid/npt/small***

Self-explanatory.

***Did not set temperature for fix rigid/nvt***

The temp keyword must be specified.

***Did not set temperature or pressure for fix rigid/npt***

The temp and press keywords must be specified.

***Dihedral atom missing in delete\_bonds***

The delete\_bonds command cannot find one or more atoms in a particular dihedral on a particular processor. The pairwise cutoff is too short or the atoms are too far apart to make a valid dihedral.

***Dihedral atom missing in set command***

The set command cannot find one or more atoms in a particular dihedral on a particular processor. The pairwise cutoff is too short or the atoms are too far apart to make a valid dihedral.

***Dihedral atoms %d %d %d %d missing on proc %d at step %ld***

One or more of 4 atoms needed to compute a particular dihedral are missing on this processor. Typically this is because the pairwise cutoff is set too short or the dihedral has blown apart and an atom is too far away.

***Dihedral atoms missing on proc %d at step %ld***

One or more of 4 atoms needed to compute a particular dihedral are missing on this processor. Typically this is because the pairwise cutoff is set too short or the dihedral has blown apart and an atom is too far away.

***Dihedral charmm is incompatible with Pair style***

Dihedral style charmm must be used with a pair style charmm in order for the 1-4 epsilon/sigma parameters to be defined.

***Dihedral coeff for hybrid has invalid style***

Dihedral style hybrid uses another dihedral style as one of its coefficients. The dihedral style used in the dihedral\_coeff command or read from a restart file is not recognized.

***Dihedral coeffs are not set***

No dihedral coefficients have been assigned in the data file or via the dihedral\_coeff command.

***Dihedral style hybrid cannot have hybrid as an argument***

Self-explanatory.

***Dihedral style hybrid cannot have none as an argument***

Self-explanatory.

***Dihedral style hybrid cannot use same dihedral style twice***

Self-explanatory.

***Dihedral/improper extent > half of periodic box length***

This error was detected by the neigh\_modify check yes setting. It is an error because the dihedral atoms are so far apart it is ambiguous how it should be defined.

***Dihedral\_coeff command before dihedral\_style is defined***

Coefficients cannot be set in the data file or via the dihedral\_coeff command until an dihedral\_style has been assigned.

***Dihedral\_coeff command before simulation box is defined***

The dihedral\_coeff command cannot be used before a read\_data, read\_restart, or create\_box command.

***Dihedral\_coeff command when no divedrals allowed***

The chosen atom style does not allow for divedrals to be defined.

***Dihedral\_style command when no divedrals allowed***

The chosen atom style does not allow for divedrals to be defined.

***Dihedrals assigned incorrectly***

Dihedrals read in from the data file were not assigned correctly to atoms. This means there is something invalid about the topology definitions.

***Dihedrals defined but no dihedral types***

The data file header lists divedrals but no dihedral types.

***Dimension command after simulation box is defined***

The dimension command cannot be used after a read\_data, read\_restart, or create\_box command.

***Disk limit not supported by OS or illegal path***

Self-explanatory.

***Dispersion PPPMDisp order has been reduced below minorder***

The default minimum order is 2. This can be reset by the kspace\_modify minorder command.

***Displace\_atoms command before simulation box is defined***

The displace\_atoms command cannot be used before a read\_data, read\_restart, or create\_box command.

***Distance must be > 0 for compute event/displace***

Self-explanatory.

***Divide by 0 in influence function***

This should not normally occur. It is likely a problem with your model.

***Divide by 0 in influence function of pair peri/lps***

This should not normally occur. It is likely a problem with your model.

***Divide by 0 in variable formula***

Self-explanatory.

***Domain too large for neighbor bins***

The domain has become extremely large so that neighbor bins cannot be used. Most likely, one or more atoms have been blown out of the simulation box to a great distance.

***Double precision is not supported on this accelerator***

Self-explanatory

***Dump atom/gz only writes compressed files***

The dump atom/gz output file name must have a .gz suffix.

***Dump cfg arguments can not mix xs|ys|zs with xsu|ysu|zsu***

Self-explanatory.

***Dump cfg arguments must start with 'mass type xs ys zs' or 'mass type xsu ysu zsu'***

This is a requirement of the CFG output format. See the dump cfg doc page for more details.

***Dump cfg requires one snapshot per file***

Use the wildcard "\*" character in the filename.

***Dump cfg/gz only writes compressed files***

The dump cfg/gz output file name must have a .gz suffix.

***Dump custom and fix not computed at compatible times***

The fix must produce per-atom quantities on timesteps that dump custom needs them.

***Dump custom compute does not calculate per-atom array***

Self-explanatory.

***Dump custom compute does not calculate per-atom vector***

Self-explanatory.

***Dump custom compute does not compute per-atom info***

Self-explanatory.

***Dump custom compute vector is accessed out-of-range***

Self-explanatory.

***Dump custom fix does not compute per-atom array***

Self-explanatory.

***Dump custom fix does not compute per-atom info***

Self-explanatory.

***Dump custom fix does not compute per-atom vector***

Self-explanatory.

***Dump custom fix vector is accessed out-of-range***

Self-explanatory.

***Dump custom variable is not atom-style variable***

Only atom-style variables generate per-atom quantities, needed for dump output.

***Dump custom/gz only writes compressed files***

The dump custom/gz output file name must have a .gz suffix.

***Dump dcd of non-matching # of atoms***

Every snapshot written by dump dcd must contain the same # of atoms.

***Dump dcd requires sorting by atom ID***

Use the dump\_modify sort command to enable this.

***Dump every variable returned a bad timestep***

The variable must return a timestep greater than the current timestep.

***Dump file MPI-IO output not allowed with % in filename***

This is because a % signifies one file per processor and MPI-IO creates one large file for all processors.

***Dump file does not contain requested snapshot***

Self-explanatory.

***Dump file is incorrectly formatted***

Self-explanatory.

***Dump image body yes requires atom style body***

Self-explanatory.

***Dump image bond not allowed with no bond types***

Self-explanatory.

***Dump image cannot perform sorting***

Self-explanatory.

***Dump image line requires atom style line***

Self-explanatory.

***Dump image requires one snapshot per file***

Use a "\*" in the filename.

***Dump image tri requires atom style tri***

Self-explanatory.

***Dump local and fix not computed at compatible times***

The fix must produce per-atom quantities on timesteps that dump local needs them.

***Dump local attributes contain no compute or fix***

Self-explanatory.

***Dump local cannot sort by atom ID***

This is because dump local does not really dump per-atom info.

***Dump local compute does not calculate local array***

Self-explanatory.

***Dump local compute does not calculate local vector***

Self-explanatory.

***Dump local compute does not compute local info***

Self-explanatory.

***Dump local compute vector is accessed out-of-range***

Self-explanatory.

***Dump local count is not consistent across input fields***

Every column of output must be the same length.



***Dump local fix does not compute local array***

Self-explanatory.

***Dump local fix does not compute local info***

Self-explanatory.

***Dump local fix does not compute local vector***

Self-explanatory.

***Dump local fix vector is accessed out-of-range***

Self-explanatory.

***Dump modify bcolor not allowed with no bond types***

Self-explanatory.

***Dump modify bdiam not allowed with no bond types***

Self-explanatory.

***Dump modify compute ID does not compute per-atom array***

Self-explanatory.

***Dump modify compute ID does not compute per-atom info***

Self-explanatory.

***Dump modify compute ID does not compute per-atom vector***

Self-explanatory.

***Dump modify compute ID vector is not large enough***

Self-explanatory.

***Dump modify element names do not match atom types***

Number of element names must equal number of atom types.

***Dump modify fix ID does not compute per-atom array***

Self-explanatory.

***Dump modify fix ID does not compute per-atom info***

Self-explanatory.

***Dump modify fix ID does not compute per-atom vector***

Self-explanatory.

***Dump modify fix ID vector is not large enough***

Self-explanatory.

***Dump modify variable is not atom-style variable***

Self-explanatory.

***Dump sort column is invalid***

Self-explanatory.

***Dump xtc requires sorting by atom ID***

Use the dump\_modify sort command to enable this.

***Dump xyz/gz only writes compressed files***

The dump xyz/gz output file name must have a .gz suffix.

***Dump\_modify buffer yes not allowed for this style***

Self-explanatory.

***Dump\_modify format string is too short***

There are more fields to be dumped in a line of output than your format string specifies.

***Dump\_modify region ID does not exist***

Self-explanatory.

***Dumping an atom property that is not allocated***

The chosen atom style does not define the per-atom quantity being dumped.

***Duplicate atom IDs exist***

Self-explanatory.

***Duplicate fields in read\_dump command***

Self-explanatory.

***Duplicate particle in PeriDynamic bond - simulation box is too small***

This is likely because your box length is shorter than 2 times the bond length.

***Electronic temperature dropped below zero***

Something has gone wrong with the fix ttm electron temperature model.

***Element not defined in potential file***

The specified element is not in the potential file.

***Empty brackets in variable***

There is no variable syntax that uses empty brackets. Check the variable doc page.

***Energy was not tallied on needed timestep***

You are using a thermo keyword that requires potentials to have tallied energy, but they did not on this timestep. See the variable page for ideas on how to make this work.

***Epsilon or sigma reference not set by pair style in PPPMDisp***

Self-explanatory.

***Epsilon or sigma reference not set by pair style in ewald/n***

The pair style is not providing the needed epsilon or sigma values.

***Error in MEAM parameter file: keyword %s (further information)***

Self-explanatory. Check the parameter file.

***Error in vdw spline: inner radius > outer radius***

A pre-tabulated spline is invalid. Likely a problem with the potential parameters.

***Error writing averaged chunk data***

Something in the output to the file triggered an error.

***Error writing file header***

Something in the output to the file triggered an error.

***Error writing out correlation data***

Something in the output to the file triggered an error.

***Error writing out histogram data***

Something in the output to the file triggered an error.

***Error writing out time averaged data***

Something in the output to the file triggered an error.

***Failed to allocate %ld bytes for array %s***

Your LAMMPS simulation has run out of memory. You need to run a smaller simulation or on more processors.

***Failed to open FFmpeg pipeline to file %s***

The specified file cannot be opened. Check that the path and name are correct and writable and that the FFmpeg executable can be found and run.

***Failed to reallocate %ld bytes for array %s***

Your LAMMPS simulation has run out of memory. You need to run a smaller simulation or on more processors.

***Fewer SRD bins than processors in some dimension***

This is not allowed. Make your SRD bin size smaller.

***File variable could not read value***

Check the file assigned to the variable.

***Final box dimension due to fix deform is < 0.0***

Self-explanatory.

***Fix %s does not allow use of dynamic group***

Dynamic groups have not yet been enabled for this fix.

***Fix ID for compute chunk/atom does not exist***

Self-explanatory.

***Fix ID for compute erotate/rigid does not exist***

Self-explanatory.

***Fix ID for compute ke/rigid does not exist***

Self-explanatory.

***Fix ID for compute reduce does not exist***

Self-explanatory.

***Fix ID for compute slice does not exist***

Self-explanatory.

***Fix ID for fix ave/atom does not exist***

Self-explanatory.

***Fix ID for fix ave/chunk does not exist***

Self-explanatory.

***Fix ID for fix ave/correlate does not exist***

Self-explanatory.

***Fix ID for fix ave/histo does not exist***

Self-explanatory.

***Fix ID for fix ave/time does not exist***

Self-explanatory.

***Fix ID for fix store/state does not exist***

Self-explanatory

***Fix ID for fix vector does not exist***

Self-explanatory.

***Fix ID for read\_data does not exist***

Self-explanatory.

***Fix ID for velocity does not exist***

Self-explanatory.

***Fix ID must be alphanumeric or underscore characters***

Self-explanatory.

***Fix SRD: bad bin assignment for SRD advection***

Something has gone wrong in your SRD model; try using more conservative settings.

***Fix SRD: bad search bin assignment***

Something has gone wrong in your SRD model; try using more conservative settings.

***Fix SRD: bad stencil bin for big particle***

Something has gone wrong in your SRD model; try using more conservative settings.

***Fix SRD: too many big particles in bin***

Reset the ATOMPERBIN parameter at the top of fix\_srd.cpp to a larger value, and re-compile the code.

***Fix SRD: too many walls in bin***

This should not happen unless your system has been setup incorrectly.

***Fix adapt interface to this pair style not supported***

New coding for the pair style would need to be done.

***Fix adapt kspace style does not exist***

Self-explanatory.

***Fix adapt pair style does not exist***

Self-explanatory

***Fix adapt pair style param not supported***

The pair style does not know about the parameter you specified.

***Fix adapt requires atom attribute charge***

The atom style being used does not specify an atom charge.

***Fix adapt requires atom attribute diameter***

The atom style being used does not specify an atom diameter.

***Fix adapt type pair range is not valid for pair hybrid sub-style***

Self-explanatory.

***Fix append/atoms requires a lattice be defined***

Use the lattice command for this purpose.

***Fix ave/atom compute array is accessed out-of-range***

Self-explanatory.

***Fix ave/atom compute does not calculate a per-atom array***

Self-explanatory.

***Fix ave/atom compute does not calculate a per-atom vector***

A compute used by fix ave/atom must generate per-atom values.

***Fix ave/atom compute does not calculate per-atom values***

A compute used by fix ave/atom must generate per-atom values.

***Fix ave/atom fix array is accessed out-of-range***

Self-explanatory.

***Fix ave/atom fix does not calculate a per-atom array***

Self-explanatory.

***Fix ave/atom fix does not calculate a per-atom vector***

A fix used by fix ave/atom must generate per-atom values.

***Fix ave/atom fix does not calculate per-atom values***

A fix used by fix ave/atom must generate per-atom values.

***Fix ave/atom variable is not atom-style variable***

A variable used by fix ave/atom must generate per-atom values.

***Fix ave/chunk compute does not calculate a per-atom array***

Self-explanatory.

***Fix ave/chunk compute does not calculate a per-atom vector***

Self-explanatory.

***Fix ave/chunk compute does not calculate per-atom values***

Self-explanatory.

***Fix ave/chunk compute vector is accessed out-of-range***

Self-explanatory.

***Fix ave/chunk does not use chunk/atom compute***

The specified compute is not for a compute chunk/atom command.

***Fix ave/chunk fix does not calculate a per-atom array***

Self-explanatory.

***Fix ave/chunk fix does not calculate a per-atom vector***

Self-explanatory.

***Fix ave/chunk fix does not calculate per-atom values***

Self-explanatory.

***Fix ave/chunk fix vector is accessed out-of-range***

Self-explanatory.

***Fix ave/chunk variable is not atom-style variable***

Self-explanatory.

***Fix ave/correlate compute does not calculate a scalar***

Self-explanatory.

***Fix ave/correlate compute does not calculate a vector***

Self-explanatory.

***Fix ave/correlate compute vector is accessed out-of-range***

The index for the vector is out of bounds.

***Fix ave/correlate fix does not calculate a scalar***

Self-explanatory.

***Fix ave/correlate fix does not calculate a vector***

Self-explanatory.

***Fix ave/correlate fix vector is accessed out-of-range***

The index for the vector is out of bounds.

***Fix ave/correlate variable is not equal-style variable***

Self-explanatory.

***Fix ave/histo cannot input local values in scalar mode***

Self-explanatory.

***Fix ave/histo cannot input per-atom values in scalar mode***

Self-explanatory.

***Fix ave/histo compute array is accessed out-of-range***

Self-explanatory.

***Fix ave/histo compute does not calculate a global array***

Self-explanatory.

***Fix ave/histo compute does not calculate a global scalar***

Self-explanatory.

*Fix ave/histo compute does not calculate a global vector*

Self-explanatory.

*Fix ave/histo compute does not calculate a local array*

Self-explanatory.

*Fix ave/histo compute does not calculate a local vector*

Self-explanatory.

*Fix ave/histo compute does not calculate a per-atom array*

Self-explanatory.

*Fix ave/histo compute does not calculate a per-atom vector*

Self-explanatory.

*Fix ave/histo compute does not calculate local values*

Self-explanatory.

*Fix ave/histo compute does not calculate per-atom values*

Self-explanatory.

*Fix ave/histo compute vector is accessed out-of-range*

Self-explanatory.

*Fix ave/histo fix array is accessed out-of-range*

Self-explanatory.

*Fix ave/histo fix does not calculate a global array*

Self-explanatory.

*Fix ave/histo fix does not calculate a global scalar*

Self-explanatory.

*Fix ave/histo fix does not calculate a global vector*

Self-explanatory.

*Fix ave/histo fix does not calculate a local array*

Self-explanatory.

*Fix ave/histo fix does not calculate a local vector*

Self-explanatory.

*Fix ave/histo fix does not calculate a per-atom array*

Self-explanatory.

*Fix ave/histo fix does not calculate a per-atom vector*

Self-explanatory.

*Fix ave/histo fix does not calculate local values*

Self-explanatory.

*Fix ave/histo fix does not calculate per-atom values*

Self-explanatory.

*Fix ave/histo fix vector is accessed out-of-range*

Self-explanatory.

*Fix ave/histo input is invalid compute*

Self-explanatory.

*Fix ave/histo input is invalid fix*

Self-explanatory.

***Fix ave/histo input is invalid variable***

Self-explanatory.

***Fix ave/histo inputs are not all global, peratom, or local***

All inputs in a single fix ave/histo command must be of the same style.

***Fix ave/histo/weight value and weight vector lengths do not match***

Self-explanatory.

***Fix ave/time cannot set output array intensive/extensive from these inputs***

One or more of the vector inputs has individual elements which are flagged as intensive or extensive. Such an input cannot be flagged as all intensive/extensive when turned into an array by fix ave/time.

***Fix ave/time cannot use variable with vector mode***

Variables produce scalar values.

***Fix ave/time columns are inconsistent lengths***

Self-explanatory.

***Fix ave/time compute array is accessed out-of-range***

An index for the array is out of bounds.

***Fix ave/time compute does not calculate a scalar***

Self-explanatory.

***Fix ave/time compute does not calculate a vector***

Self-explanatory.

***Fix ave/time compute does not calculate an array***

Self-explanatory.

***Fix ave/time compute vector is accessed out-of-range***

The index for the vector is out of bounds.

***Fix ave/time fix array cannot be variable length***

Self-explanatory.

***Fix ave/time fix array is accessed out-of-range***

An index for the array is out of bounds.

***Fix ave/time fix does not calculate a scalar***

Self-explanatory.

***Fix ave/time fix does not calculate a vector***

Self-explanatory.

***Fix ave/time fix does not calculate an array***

Self-explanatory.

***Fix ave/time fix vector cannot be variable length***

Self-explanatory.

***Fix ave/time fix vector is accessed out-of-range***

The index for the vector is out of bounds.

***Fix ave/time variable is not equal-style variable***

Self-explanatory.

***Fix balance rcb cannot be used with comm\_style brick***

Comm\_style tiled must be used instead.

***Fix balance shift string is invalid***

The string can only contain the characters “x”, “y”, or “z”.

***Fix bond/break needs ghost atoms from further away***

This is because the fix needs to walk bonds to a certain distance to acquire needed info, The comm\_modify cutoff command can be used to extend the communication range.

***Fix bond/create angle type is invalid***

Self-explanatory.

***Fix bond/create cutoff is longer than pairwise cutoff***

This is not allowed because bond creation is done using the pairwise neighbor list.

***Fix bond/create dihedral type is invalid***

Self-explanatory.

***Fix bond/create improper type is invalid***

Self-explanatory.

***Fix bond/create induced too many angles/dihedrals/impropers per atom***

See the read\_data command for info on using the “extra/angle/per/atom”, (or dihedral, improper) keywords to allow for additional angles, dihedrals, and impropers to be formed.

***Fix bond/create needs ghost atoms from further away***

This is because the fix needs to walk bonds to a certain distance to acquire needed info, The comm\_modify cutoff command can be used to extend the communication range.

***Fix bond/react: Cannot use fix bond/react with non-molecular systems***

Only systems with bonds that can be changed can be used. Atom\_style template does not qualify.

***Fix bond/react: Invalid template atom ID in map file***

Atom IDs in molecule templates range from 1 to the number of atoms in the template.

***Fix bond/react: Rmax cutoff is longer than pairwise cutoff***

This is not allowed because bond creation is done using the pairwise neighbor list.

***Fix bond/react: Molecule template ID for fix bond/react does not exist***

A valid molecule template must have been created with the molecule command.

***Fix bond/react: Reaction templates must contain the same number of atoms***

There should be a one-to-one correspondence between atoms in the pre-reacted and post-reacted templates, as specified by the map file.

***Fix bond/react: Unknown section in map file***

Please ensure reaction map files are properly formatted.

***Fix bond/react: Atom/Bond type affected by reaction too close to template edge***

This means an atom which changes type or connectivity during the reaction is too close to an ‘edge’ atom defined in the map file. This could cause incorrect assignment of bonds, angle, etc. Generally, this means you must include more atoms in your templates, such that there are at least two atoms between each atom involved in the reaction and an edge atom.

***Fix bond/react: Fix bond/react needs ghost atoms from farther away***

This is because a processor needs to map the entire unreacted molecule template onto simulation atoms it knows about. The comm\_modify cutoff command can be used to extend the communication range.

***Fix bond/react: A deleted atom cannot remain bonded to an atom that is not deleted***

Self-explanatory.

***Fix bond/react: First neighbors of chiral atoms must be of mutually different types***

Self-explanatory.

***Fix bond/react: Chiral atoms must have exactly four first neighbors***

Self-explanatory.



***Fix bond/react: Molecule template 'Coords' section required for chiralIDs keyword***

The coordinates of atoms in the pre-reacted template are used to determine chirality.

***Fix bond/react special bond generation overflow***

The number of special bonds per-atom created by a reaction exceeds the system setting. See the read\_data or create\_box command for how to specify this value.

***Fix bond/react topology/atom exceed system topology/atom***

The number of bonds, angles etc per-atom created by a reaction exceeds the system setting. See the read\_data or create\_box command for how to specify this value.

***Fix bond/swap cannot use dihedral or improper styles***

These styles cannot be defined when using this fix.

***Fix bond/swap requires pair and bond styles***

Self-explanatory.

***Fix bond/swap requires special\_bonds = 0,1,1***

Self-explanatory.

***Fix box/relax generated negative box length***

The pressure being applied is likely too large. Try applying it incrementally, to build to the high pressure.

***Fix command before simulation box is defined***

The fix command cannot be used before a read\_data, read\_restart, or create\_box command.

***Fix deform cannot use yz variable with xy***

The yz setting cannot be a variable if xy deformation is also specified. This is because LAMMPS cannot determine if the yz setting will induce a box flip which would be invalid if xy is also changing.

***Fix deform is changing yz too much with xy***

When both yz and xy are changing, it induces changes in xz if the box must flip from one tilt extreme to another. Thus it is not allowed for yz to grow so much that a flip is induced.

***Fix deform tilt factors require triclinic box***

Cannot deform the tilt factors of a simulation box unless it is a triclinic (non-orthogonal) box.

***Fix deform volume setting is invalid***

Cannot use volume style unless other dimensions are being controlled.

***Fix deposit and fix rigid/small not using same molecule template ID***

Self-explanatory.

***Fix deposit and fix shake not using same molecule template ID***

Self-explanatory.

***Fix deposit molecule must have atom types***

The defined molecule does not specify atom types.

***Fix deposit molecule must have coordinates***

The defined molecule does not specify coordinates.

***Fix deposit molecule template ID must be same as atom\_style template ID***

When using atom\_style template, you cannot deposit molecules that are not in that template.

***Fix deposit region cannot be dynamic***

Only static regions can be used with fix deposit.

***Fix deposit region does not support a bounding box***

Not all regions represent bounded volumes. You cannot use such a region with the fix deposit command.

***Fix deposit shake fix does not exist***

Self-explanatory.

***Fix efield requires atom attribute q or mu***

The atom style defined does not have this attribute.

***Fix efield with dipoles cannot use atom-style variables***

This option is not supported.

***Fix evaporate molecule requires atom attribute molecule***

The atom style being used does not define a molecule ID.

***Fix external callback function not set***

This must be done by an external program in order to use this fix.

***Fix for fix ave/atom not computed at compatible time***

Fixes generate their values on specific timesteps. Fix ave/atom is requesting a value on a non-allowed timestep.

***Fix for fix ave/chunk not computed at compatible time***

Fixes generate their values on specific timesteps. Fix ave/chunk is requesting a value on a non-allowed timestep.

***Fix for fix ave/correlate not computed at compatible time***

Fixes generate their values on specific timesteps. Fix ave/correlate is requesting a value on a non-allowed timestep.

***Fix for fix ave/histo not computed at compatible time***

Fixes generate their values on specific timesteps. Fix ave/histo is requesting a value on a non-allowed timestep.

***Fix for fix ave/spatial not computed at compatible time***

Fixes generate their values on specific timesteps. Fix ave/spatial is requesting a value on a non-allowed timestep.

***Fix for fix ave/time not computed at compatible time***

Fixes generate their values on specific timesteps. Fix ave/time is requesting a value on a non-allowed timestep.

***Fix for fix store/state not computed at compatible time***

Fixes generate their values on specific timesteps. Fix store/state is requesting a value on a non-allowed timestep.

***Fix for fix vector not computed at compatible time***

Fixes generate their values on specific timesteps. Fix vector is requesting a value on a non-allowed timestep.

***Fix freeze requires atom attribute torque***

The atom style defined does not have this attribute.

***Fix gcmc and fix shake not using same molecule template ID***

Self-explanatory.

***Fix gcmc atom has charge, but atom style does not***

Self-explanatory.

***Fix gcmc cannot exchange individual atoms belonging to a molecule***

This is an error since you should not delete only one atom of a molecule. The user has specified atomic (non-molecular) gas exchanges, but an atom belonging to a molecule could be deleted.

***Fix gcmc does not (yet) work with atom\_style template***

Self-explanatory.

***Fix gcmc molecule command requires that atoms have molecule attributes***

Should not choose the gcmc molecule feature if no molecules are being simulated. The general molecule flag is off, but gcmc's molecule flag is on.

***Fix gcmc molecule has charges, but atom style does not***

Self-explanatory.

***Fix gcmc molecule must have atom types***

The defined molecule does not specify atom types.

***Fix gcmc molecule must have coordinates***

The defined molecule does not specify coordinates.

***Fix gcmc molecule template ID must be same as atom\_style template ID***

When using atom\_style template, you cannot insert molecules that are not in that template.

***Fix gcmc put atom outside box***

This should not normally happen. Contact the developers.

***Fix gcmc ran out of available atom IDs***

See the setting for tagint in the src/lmptype.h file.

***Fix gcmc ran out of available molecule IDs***

See the setting for tagint in the src/lmptype.h file.

***Fix gcmc region cannot be dynamic***

Only static regions can be used with fix gcmc.

***Fix gcmc region does not support a bounding box***

Not all regions represent bounded volumes. You cannot use such a region with the fix gcmc command.

***Fix gcmc region extends outside simulation box***

Self-explanatory.

***Fix gcmc shake fix does not exist***

Self-explanatory.

***Fix gld c coefficients must be  $\geq 0$*** 

Self-explanatory.

***Fix gld needs more prony series coefficients***

Self-explanatory.

***Fix gld prony terms must be  $> 0$*** 

Self-explanatory.

***Fix gld series type must be pprony for now***

Self-explanatory.

***Fix gld start temperature must be  $\geq 0$*** 

Self-explanatory.

***Fix gld stop temperature must be  $\geq 0$*** 

Self-explanatory.

***Fix gld tau coefficients must be  $> 0$*** 

Self-explanatory.

***Fix halt variable is not equal-style variable***

Self-explanatory.

***Fix heat group has no atoms***

Self-explanatory.

***Fix heat kinetic energy of an atom went negative***

This will cause the velocity rescaling about to be performed by fix heat to be invalid.

***Fix heat kinetic energy went negative***

This will cause the velocity rescaling about to be performed by fix heat to be invalid.

***Fix in variable not computed at compatible time***

Fixes generate their values on specific timesteps. The variable is requesting the values on a non-allowed timestep.

***Fix langevin angmom is not yet implemented with kokkos***

This option is not yet available.

***Fix langevin angmom requires atom style ellipsoid***

Self-explanatory.

***Fix langevin angmom requires extended particles***

This fix option cannot be used with point particles.

***Fix langevin gif and respa are not compatible***

Self-explanatory.

***Fix langevin gif cannot have period equal to dt/2***

If the period is equal to dt/2 then division by zero will happen.

***Fix langevin gif should come before fix nve***

Self-explanatory.

***Fix langevin gif with tbias is not yet implemented with kokkos***

This option is not yet available.

***Fix langevin omega is not yet implemented with kokkos***

This option is not yet available.

***Fix langevin omega requires atom style sphere***

Self-explanatory.

***Fix langevin omega requires extended particles***

One of the particles has radius 0.0.

***Fix langevin period must be > 0.0***

The time window for temperature relaxation must be > 0

***Fix langevin variable returned negative temperature***

Self-explanatory.

***Fix momentum group has no atoms***

Self-explanatory.

***Fix move cannot define z or vz variable for 2d problem***

Self-explanatory.

***Fix move cannot rotate around non z-axis for 2d problem***

Self-explanatory.

***Fix move cannot set linear z motion for 2d problem***

Self-explanatory.

***Fix move cannot set wiggle z motion for 2d problem***

Self-explanatory.

***Fix msst compute ID does not compute potential energy***

Self-explanatory.

***Fix msst compute ID does not compute pressure***

Self-explanatory.

***Fix msst compute ID does not compute temperature***

Self-explanatory.

***Fix msst requires a periodic box***

Self-explanatory.

***Fix msst tscale must satisfy  $0 \leq tscale < 1$*** 

Self-explanatory.

***Fix npt/nph has tilted box too far in one step - periodic cell is too far from equilibrium state***

Self-explanatory. The change in the box tilt is too extreme on a short timescale.

***Fix numdiff requires an atom map, see atom\_modify***

Self-explanatory. Efficient loop over all atoms for numerical difference requires an atom map.

***Fix numdiff requires consecutive atom IDs***

Self-explanatory. Efficient loop over all atoms for numerical difference requires consecutive atom IDs.

***Fix numdiff/virial must use group all***

Virial contributions computed by this fix are computed on all atoms.

***Fix nve/asphere requires extended particles***

This fix can only be used for particles with a shape setting.

***Fix nve/asphere/noforce requires atom style ellipsoid***

Self-explanatory.

***Fix nve/asphere/noforce requires extended particles***

One of the particles is not an ellipsoid.

***Fix nve/body requires atom style body***

Self-explanatory.

***Fix nve/body requires bodies***

This fix can only be used for particles that are bodies.

***Fix nve/line can only be used for 2d simulations***

Self-explanatory.

***Fix nve/line requires atom style line***

Self-explanatory.

***Fix nve/line requires line particles***

Self-explanatory.

***Fix nve/sphere dipole requires atom attribute mu***

An atom style with this attribute is needed.

***Fix nve/sphere requires atom style sphere***

Self-explanatory.

***Fix nve/sphere requires extended particles***

This fix can only be used for particles of a finite size.

***Fix nve/tri can only be used for 3d simulations***

Self-explanatory.

***Fix nve/tri requires atom style tri***

Self-explanatory.

***Fix nve/tri requires tri particles***

Self-explanatory.

***Fix nvt/nph/npt asphere requires extended particles***

The shape setting for a particle in the fix group has shape = 0.0, which means it is a point particle.

***Fix nvt/nph/npt body requires bodies***

Self-explanatory.

***Fix nvt/nph/npt sphere requires atom style sphere***

Self-explanatory.

***Fix nvt/npt/nph damping parameters must be > 0.0***

Self-explanatory.

***Fix nvt/npt/nph dilate group ID does not exist***

Self-explanatory.

***Fix nvt/sphere requires extended particles***

This fix can only be used for particles of a finite size.

***Fix orient/fcc file open failed***

The fix orient/fcc command could not open a specified file.

***Fix orient/fcc file read failed***

The fix orient/fcc command could not read the needed parameters from a specified file.

***Fix orient/fcc found self twice***

The neighbor lists used by fix orient/fcc are messed up. If this error occurs, it is likely a bug, so send an email to the [developers](#).

***Fix peri neigh does not exist***

Somehow a fix that the pair style defines has been deleted.

***Fix pour and fix rigid/small not using same molecule template ID***

Self-explanatory.

***Fix pour and fix shake not using same molecule template ID***

Self-explanatory.

***Fix pour insertion count per timestep is 0***

Self-explanatory.

***Fix pour molecule must have atom types***

The defined molecule does not specify atom types.

***Fix pour molecule must have coordinates***

The defined molecule does not specify coordinates.

***Fix pour molecule template ID must be same as atom style template ID***

When using atom\_style template, you cannot pour molecules that are not in that template.

***Fix pour polydisperse fractions do not sum to 1.0***

Self-explanatory.

***Fix pour region ID does not exist***

Self-explanatory.

***Fix pour region cannot be dynamic***

Only static regions can be used with fix pour.

***Fix pour region does not support a bounding box***

Not all regions represent bounded volumes. You cannot use such a region with the fix pour command.

***Fix pour requires atom attributes radius, rmass***

The atom style defined does not have these attributes.

***Fix pour rigid fix does not exist***

Self-explanatory.

***Fix pour shake fix does not exist***

Self-explanatory.

***Fix press/berendsen damping parameters must be > 0.0***

Self-explanatory.

***Fix property/atom cannot specify mol twice***

Self-explanatory.

***Fix property/atom cannot specify q twice***

Self-explanatory.

***Fix property/atom mol when atom\_style already has molecule attribute***

Self-explanatory.

***Fix property/atom q when atom\_style already has charge attribute***

Self-explanatory.

***Fix property/atom vector name already exists***

The name for an integer or floating-point vector must be unique.

***Fix qeq has negative upper Taper radius cutoff***

Self-explanatory.

***Fix qeq/comb group has no atoms***

Self-explanatory.

***Fix qeq/comb requires atom attribute q***

An atom style with charge must be used to perform charge equilibration.

***Fix qeq/dynamic group has no atoms***

Self-explanatory.

***Fix qeq/dynamic requires atom attribute q***

Self-explanatory.

***Fix qeq/fire group has no atoms***

Self-explanatory.

***Fix qeq/fire requires atom attribute q***

Self-explanatory.

***Fix qeq/point group has no atoms***

Self-explanatory.

***Fix qeq/point has insufficient QEq matrix size***

Occurs when number of neighbor atoms for an atom increased too much during a run. Increase SAFE\_ZONE and MIN\_CAP in fix\_qeq.h and re-compile.

***Fix qeq/point requires atom attribute q***

Self-explanatory.

***Fix qeq/shielded group has no atoms***

Self-explanatory.

***Fix qeq/shielded has insufficient QEq matrix size***

Occurs when number of neighbor atoms for an atom increased too much during a run. Increase SAFE\_ZONE and MIN\_CAP in fix\_qeq.h and re-compile.

***Fix qeq/shielded requires atom attribute q***

Self-explanatory.

***Fix qeq/slater could not extract params from pair coul/streitz***

This should not happen unless pair coul/streitz has been altered.

***Fix qeq/slater group has no atoms***

Self-explanatory.

***Fix qeq/slater has insufficient QEq matrix size***

Occurs when number of neighbor atoms for an atom increased too much during a run. Increase SAFE\_ZONE and MIN\_CAP in fix\_qeq.h and re-compile.

***Fix qeq/slater requires atom attribute q***

Self-explanatory.

***Fix reax/bonds numbonds > nsbmax\_most***

The limit of the number of bonds expected by the ReaxFF force field was exceeded.

***Fix recenter group has no atoms***

Self-explanatory.

***Fix restrain requires an atom map, see atom\_modify***

Self-explanatory.

***Fix rigid atom has non-zero image flag in a non-periodic dimension***

Image flags for non-periodic dimensions should not be set.

***Fix rigid file has no lines***

Self-explanatory.

***Fix rigid langevin period must be > 0.0***

Self-explanatory.

***Fix rigid molecule requires atom attribute molecule***

Self-explanatory.

***Fix rigid npt/nph dilate group ID does not exist***

Self-explanatory.

***Fix rigid npt/nph does not yet allow triclinic box***

This is a current restriction in LAMMPS.

***Fix rigid npt/nph period must be > 0.0***

Self-explanatory.

***Fix rigid npt/small t\_chain should not be less than 1***

Self-explanatory.

***Fix rigid npt/small t\_order must be 3 or 5***

Self-explanatory.

***Fix rigid nvt/npt/nph damping parameters must be > 0.0***

Self-explanatory.

***Fix rigid nvt/small t\_chain should not be less than 1***

Self-explanatory.

***Fix rigid nvt/small t\_iter should not be less than 1***

Self-explanatory.

***Fix rigid nvt/small t\_order must be 3 or 5***

Self-explanatory.

***Fix rigid xy torque cannot be on for 2d simulation***

Self-explanatory.

***Fix rigid z force cannot be on for 2d simulation***

Self-explanatory.

***Fix rigid/npt period must be > 0.0***

Self-explanatory.



**Fix rigid/npt temperature order must be 3 or 5**

Self-explanatory.

**Fix rigid/npt/small period must be > 0.0**

Self-explanatory.

**Fix rigid/nvt period must be > 0.0**

Self-explanatory.

**Fix rigid/nvt temperature order must be 3 or 5**

Self-explanatory.

**Fix rigid/nvt/small period must be > 0.0**

Self-explanatory.

**Fix rigid/small atom has non-zero image flag in a non-periodic dimension**

Image flags for non-periodic dimensions should not be set.

**Fix rigid/small langevin period must be > 0.0**

Self-explanatory.

**Fix rigid/small molecule must have atom types**

The defined molecule does not specify atom types.

**Fix rigid/small molecule must have coordinates**

The defined molecule does not specify coordinates.

**Fix rigid/small npt/nph period must be > 0.0**

Self-explanatory.

**Fix rigid/small nvt/npt/nph damping parameters must be > 0.0**

Self-explanatory.

**Fix rigid/small nvt/npt/nph dilate group ID does not exist**

Self-explanatory.

**Fix rigid/small requires an atom map, see atom\_modify**

Self-explanatory.

**Fix rigid/small requires atom attribute molecule**

Self-explanatory.

**Fix rigid: Bad principal moments**

The principal moments of inertia computed for a rigid body are not within the required tolerances.

**Fix shake cannot be used with minimization**

Cannot use fix shake while doing an energy minimization since it turns off bonds that should contribute to the energy.

**Fix shake molecule template must have shake info**

The defined molecule does not specify SHAKE information.

**Fix spring couple group ID does not exist**

Self-explanatory.

**Fix srd can only currently be used with comm\_style brick**

This is a current restriction in LAMMPS.

**Fix srd lamda must be >= 0.6 of SRD grid size**

This is a requirement for accuracy reasons.

**Fix srd no-slip requires atom attribute torque**

This is because the SRD collisions will impart torque to the solute particles.

***Fix srd requires SRD particles all have same mass***

Self-explanatory.

***Fix srd requires ghost atoms store velocity***

Use the comm\_modify vel yes command to enable this.

***Fix srd requires newton pair on***

Self-explanatory.

***Fix store/state compute array is accessed out-of-range***

Self-explanatory.

***Fix store/state compute does not calculate a per-atom array***

The compute calculates a per-atom vector.

***Fix store/state compute does not calculate a per-atom vector***

The compute calculates a per-atom vector.

***Fix store/state compute does not calculate per-atom values***

Computes that calculate global or local quantities cannot be used with fix store/state.

***Fix store/state fix array is accessed out-of-range***

Self-explanatory.

***Fix store/state fix does not calculate a per-atom array***

The fix calculates a per-atom vector.

***Fix store/state fix does not calculate a per-atom vector***

The fix calculates a per-atom array.

***Fix store/state fix does not calculate per-atom values***

Fixes that calculate global or local quantities cannot be used with fix store/state.

***Fix store/state for atom property that is not allocated***

Self-explanatory.

***Fix store/state variable is not atom-style variable***

Only atom-style variables calculate per-atom quantities.

***Fix temp/berendsen period must be > 0.0***

Self-explanatory.

***Fix temp/berendsen variable returned negative temperature***

Self-explanatory.

***Fix temp/csld is not compatible with fix rattle or fix shake***

These two commands cannot currently be used together with fix temp/csld.

***Fix temp/csld variable returned negative temperature***

Self-explanatory.

***Fix temp/csvr variable returned negative temperature***

Self-explanatory.

***Fix temp/rescale variable returned negative temperature***

Self-explanatory.

***Fix tfmc displacement length must be > 0***

Self-explanatory.

***Fix tfmc is not compatible with fix shake***

These two commands cannot currently be used together.

**Fix tfmc temperature must be > 0**

Self-explanatory.

**Fix thermal/conductivity swap value must be positive**

Self-explanatory.

**Fix tmd must come after integration fixes**

Any fix tmd command must appear in the input script after all time integration fixes (nve, nvt, npt). See the fix tmd documentation for details.

**Fix ttm electron temperatures must be > 0.0**

Self-explanatory.

**Fix ttm electronic\_density must be > 0.0**

Self-explanatory.

**Fix ttm electronic\_specific\_heat must be > 0.0**

Self-explanatory.

**Fix ttm electronic\_thermal\_conductivity must be >= 0.0**

Self-explanatory.

**Fix ttm gamma\_p must be > 0.0**

Self-explanatory.

**Fix ttm gamma\_s must be >= 0.0**

Self-explanatory.

**Fix ttm number of nodes must be > 0**

Self-explanatory.

**Fix ttm v\_0 must be >= 0.0**

Self-explanatory.

**Fix used in compute chunk/atom not computed at compatible time**

The chunk/atom compute cannot query the output of the fix on a timestep it is needed.

**Fix used in compute reduce not computed at compatible time**

Fixes generate their values on specific timesteps. Compute reduce is requesting a value on a non-allowed timestep.

**Fix used in compute slice not computed at compatible time**

Fixes generate their values on specific timesteps. Compute slice is requesting a value on a non-allowed timestep.

**Fix vector cannot set output array intensive/extensive from these inputs**

The inputs to the command have conflicting intensive/extensive attributes. You need to use more than one fix vector command.

**Fix vector compute does not calculate a scalar**

Self-explanatory.

**Fix vector compute does not calculate a vector**

Self-explanatory.

**Fix vector compute vector is accessed out-of-range**

Self-explanatory.

**Fix vector fix does not calculate a scalar**

Self-explanatory.

**Fix vector fix does not calculate a vector**

Self-explanatory.

**Fix vector fix vector is accessed out-of-range**

Self-explanatory.

***Fix vector variable is not equal-style variable***

Self-explanatory.

***Fix viscosity swap value must be positive***

Self-explanatory.

***Fix viscosity vtarget value must be positive***

Self-explanatory.

***Fix wall cutoff <= 0.0***

Self-explanatory.

***Fix wall/colloid requires atom style sphere***

Self-explanatory.

***Fix wall/colloid requires extended particles***

One of the particles has radius 0.0.

***Fix wall/gran is incompatible with Pair style***

Must use a granular pair style to define the parameters needed for this fix.

***Fix wall/gran requires atom style sphere***

Self-explanatory.

***Fix wall/piston command only available at zlo***

The face keyword must be zlo.

***Fix wall/region colloid requires atom style sphere***

Self-explanatory.

***Fix wall/region colloid requires extended particles***

One of the particles has radius 0.0.

***Fix wall/region cutoff <= 0.0***

Self-explanatory.

***Fix\_modify pressure ID does not compute pressure***

The compute ID assigned to the fix must compute pressure.

***Fix\_modify temperature ID does not compute temperature***

The compute ID assigned to the fix must compute temperature.

***For triclinic deformation, specified target stress must be hydrostatic***

Triclinic pressure control is allowed using the tri keyword, but non-hydrostatic pressure control can not be used in this case.

***Found no restart file matching pattern***

When using a "\*" in the restart file name, no matching file was found.

***GPU library not compiled for this accelerator***

Self-explanatory.

***GPU package does not (yet) work with atom\_style template***

Self-explanatory.

***GPU particle split must be set to 1 for this pair style.***

For this pair style, you cannot run part of the force calculation on the host. See the package command.

***GPUs are requested but Kokkos has not been compiled for CUDA***

Re-compile Kokkos with CUDA support to use GPUs.

***Ghost velocity forward comm not yet implemented with Kokkos***

This is a current restriction.

***Gmask function in equal-style variable formula***

Gmask is per-atom operation.

***Gravity changed since fix pour was created***

The gravity vector defined by fix gravity must be static.

***Gravity must point in -y to use with fix pour in 2d***

Self-explanatory.

***Gravity must point in -z to use with fix pour in 3d***

Self-explanatory.

***Gmask function in equal-style variable formula***

Gmask is per-atom operation.

***Group ID does not exist***

A group ID used in the group command does not exist.

***Group ID in variable formula does not exist***

Self-explanatory.

***Group all cannot be made dynamic***

This operation is not allowed.

***Group command before simulation box is defined***

The group command cannot be used before a read\_data, read\_restart, or create\_box command.

***Group dynamic cannot reference itself***

Self-explanatory.

***Group dynamic parent group cannot be dynamic***

Self-explanatory.

***Group dynamic parent group does not exist***

Self-explanatory.

***Group region ID does not exist***

A region ID used in the group command does not exist.

***If read\_dump purges it cannot replace or trim***

These operations are not compatible. See the read\_dump doc page for details.

***Illegal ... command***

Self-explanatory. Check the input script syntax and compare to the documentation for the command. You can use -echo screen as a command-line option when running LAMMPS to see the offending line.

***Illegal COMB parameter***

One or more of the coefficients defined in the potential file is invalid.

***Illegal COMB3 parameter***

One or more of the coefficients defined in the potential file is invalid.

***Illegal Stillinger-Weber parameter***

One or more of the coefficients defined in the potential file is invalid.

***Illegal Tersoff parameter***

One or more of the coefficients defined in the potential file is invalid.

***Illegal Vashishta parameter***

One or more of the coefficients defined in the potential file is invalid.

***Illegal compute voronoi/atom command (occupation and (surface or edges))***

Self-explanatory.

***Illegal coul/streitz parameter***

One or more of the coefficients defined in the potential file is invalid.

***Illegal dump\_modify sfactor value (must be > 0.0)***

Self-explanatory.

***Illegal dump\_modify tfactor value (must be > 0.0)***

Self-explanatory.

***Illegal fix gcmc gas mass <= 0***

The computed mass of the designated gas molecule or atom type was less than or equal to zero.

***Illegal fix tfmc random seed***

Seeds can only be nonzero positive integers.

***Illegal fix wall/piston velocity***

The piston velocity must be positive.

***Illegal integrate style***

Self-explanatory.

***Illegal nb3b/harmonic parameter***

One or more of the coefficients defined in the potential file is invalid.

***Illegal number of angle table entries***

There must be at least 2 table entries.

***Illegal number of bond table entries***

There must be at least 2 table entries.

***Illegal number of pair table entries***

There must be at least 2 table entries.

***Illegal or unset periodicity in restart***

This error should not normally occur unless the restart file is invalid.

***Illegal range increment value***

The increment must be  $\geq 1$ .

***Illegal simulation box***

The lower bound of the simulation box is greater than the upper bound.

***Illegal size double vector read requested***

This error should not normally occur unless the restart file is invalid.

***Illegal size integer vector read requested***

This error should not normally occur unless the restart file is invalid.

***Illegal size string or corrupt restart***

This error should not normally occur unless the restart file is invalid.

***Imageint setting in lmptype.h is invalid***

Imageint must be as large or larger than smallint.

***Imageint setting in lmptype.h is not compatible***

Format of imageint stored in restart file is not consistent with LAMMPS version you are running. See the settings in src/lmptype.h

***Improper atom missing in delete\_bonds***

The delete\_bonds command cannot find one or more atoms in a particular improper on a particular processor. The pairwise cutoff is too short or the atoms are too far apart to make a valid improper.

***Improper atom missing in set command***

The set command cannot find one or more atoms in a particular improper on a particular processor. The pairwise cutoff is too short or the atoms are too far apart to make a valid improper.

***Improper atoms %d %d %d %d missing on proc %d at step %ld***

One or more of 4 atoms needed to compute a particular improper are missing on this processor. Typically this is because the pairwise cutoff is set too short or the improper has blown apart and an atom is too far away.

***Improper atoms missing on proc %d at step %ld***

One or more of 4 atoms needed to compute a particular improper are missing on this processor. Typically this is because the pairwise cutoff is set too short or the improper has blown apart and an atom is too far away.

***Improper coeff for hybrid has invalid style***

Improper style hybrid uses another improper style as one of its coefficients. The improper style used in the improper\_coeff command or read from a restart file is not recognized.

***Improper coeffs are not set***

No improper coefficients have been assigned in the data file or via the improper\_coeff command.

***Improper style hybrid cannot have hybrid as an argument***

Self-explanatory.

***Improper style hybrid cannot have none as an argument***

Self-explanatory.

***Improper style hybrid cannot use same improper style twice***

Self-explanatory.

***Improper\_coeff command before improper\_style is defined***

Coefficients cannot be set in the data file or via the improper\_coeff command until an improper\_style has been assigned.

***Improper\_coeff command before simulation box is defined***

The improper\_coeff command cannot be used before a read\_data, read\_restart, or create\_box command.

***Improper\_coeff command when no impropers allowed***

The chosen atom style does not allow for impropers to be defined.

***Improper\_style command when no impropers allowed***

The chosen atom style does not allow for impropers to be defined.

***Impropers assigned incorrectly***

Impropers read in from the data file were not assigned correctly to atoms. This means there is something invalid about the topology definitions.

***Impropers defined but no improper types***

The data file header lists improper but no improper types.

***Incompatible KIM Simulator Model***

The requested KIM Simulator Model was defined for a different MD code and thus is not compatible with LAMMPS.

***Incompatible units for KIM Simulator Model***

The selected unit style is not compatible with the requested KIM Simulator Model.

***Incomplete use of variables in create\_atoms command***

The var and set options must be used together.

***Inconsistent iparam/jparam values in fix bond/create command***

If itype and jtype are the same, then their maxbond and newtype settings must also be the same.

***Inconsistent line segment in data file***

The end points of the line segment are not equal distances from the center point which is the atom coordinate.

***Inconsistent triangle in data file***

The centroid of the triangle as defined by the corner points is not the atom coordinate.

***Inconsistent use of finite-size particles by molecule template molecules***

Not all of the molecules define a radius for their constituent particles.

***Incorrect # of floating-point values in Bodies section of data file***

See page for body style.

***Incorrect # of integer values in Bodies section of data file***

See page for body style.

***Incorrect %s format in data file***

A section of the data file being read by fix property/atom does not have the correct number of values per line.

***Incorrect SNAP parameter file***

The file cannot be parsed correctly, check its internal syntax.

***Incorrect args for angle coefficients***

Self-explanatory. Check the input script or data file.

***Incorrect args for bond coefficients***

Self-explanatory. Check the input script or data file.

***Incorrect args for dihedral coefficients***

Self-explanatory. Check the input script or data file.

***Incorrect args for improper coefficients***

Self-explanatory. Check the input script or data file.

***Incorrect args for pair coefficients***

Self-explanatory. Check the input script or data file.

***Incorrect args in pair\_style command***

Self-explanatory.

***Incorrect atom format in data file***

Number of values per atom line in the data file is not consistent with the atom style.

***Incorrect atom format in neb file***

The number of fields per line is not what expected.

***Incorrect bonus data format in data file***

See the read\_data page for a description of how various kinds of bonus data must be formatted for certain atom styles.

***Incorrect boundaries with slab Ewald***

Must have periodic x,y dimensions and non-periodic z dimension to use 2d slab option with Ewald.

***Incorrect boundaries with slab EwaldDisp***

Must have periodic x,y dimensions and non-periodic z dimension to use 2d slab option with Ewald.

***Incorrect boundaries with slab PPPM***

Must have periodic x,y dimensions and non-periodic z dimension to use 2d slab option with PPPM.

***Incorrect boundaries with slab PPPMDisp***

Must have periodic x,y dimensions and non-periodic z dimension to use 2d slab option with ppm/dis.

***Incorrect conversion in format string***

A format style variable was not using either a %f, a %g, or a %e conversion. Or an immediate variable with format suffix was not using either a %f, a %g or a %e conversion in the format suffix.

***Incorrect element names in ADP potential file***

The element names in the ADP file do not match those requested.



***Incorrect element names in EAM potential file***

The element names in the EAM file do not match those requested.

***Incorrect format of ... section in data file***

Number or type of values per line in the given section of the data file is not consistent with the requirements for this section.

***Incorrect format in COMB potential file***

Incorrect number of words per line in the potential file.

***Incorrect format in COMB3 potential file***

Incorrect number of words per line in the potential file.

***Incorrect format in MEAM library file***

Incorrect number of words per line in the potential file.

***Incorrect format in SNAP coefficient file***

Incorrect number of words per line in the coefficient file.

***Incorrect format in SNAP parameter file***

Incorrect number of words per line in the parameter file.

***Incorrect format in Stillinger-Weber potential file***

Incorrect number of words per line in the potential file.

***Incorrect format in TMD target file***

Format of file read by fix tmd command is incorrect.

***Incorrect format in Tersoff potential file***

Incorrect number of words per line in the potential file.

***Incorrect format in Vashishta potential file***

Incorrect number of words per line in the potential file.

***Incorrect format in coul/streitz potential file***

Incorrect number of words per line in the potential file.

***Incorrect format in nb3b/harmonic potential file***

Incorrect number of words per line in the potential file.

***Incorrect integer value in Bodies section of data file***

See page for body style.

***Incorrect multiplicity arg for dihedral coefficients***

Self-explanatory. Check the input script or data file.

***Incorrect number of elements in potential file***

Self-explanatory.

***Incorrect rigid body format in fix rigid file***

The number of fields per line is not what expected.

***Incorrect rigid body format in fix rigid/small file***

The number of fields per line is not what expected.

***Incorrect sign arg for dihedral coefficients***

Self-explanatory. Check the input script or data file.

***Incorrect table format check for element types***

Self-explanatory.

***Incorrect velocity format in data file***

Each atom style defines a format for the Velocity section of the data file. The read-in lines do not match.

***Incorrect weight arg for dihedral coefficients***

Self-explanatory. Check the input script or data file.

***Index between variable brackets must be positive***

Self-explanatory.

***Indexed per-atom vector in variable formula without atom map***

Accessing a value from an atom vector requires the ability to lookup an atom index, which is provided by an atom map. An atom map does not exist (by default) for non-molecular problems. Using the atom\_modify map command will force an atom map to be created.

***Initial temperatures not all set in fix ttm***

Self-explanatory.

***Input line quote not followed by white-space***

An end quote must be followed by white-space.

***Insertion region extends outside simulation box***

Self-explanatory.

***Insufficient Jacobi rotations for POEMS body***

Eigensolve for rigid body was not sufficiently accurate.

***Insufficient Jacobi rotations for body nparticle***

Eigensolve for rigid body was not sufficiently accurate.

***Insufficient Jacobi rotations for rigid body***

Eigensolve for rigid body was not sufficiently accurate.

***Insufficient Jacobi rotations for rigid molecule***

Eigensolve for rigid body was not sufficiently accurate.

***Insufficient Jacobi rotations for triangle***

The calculation of the inertia tensor of the triangle failed. This should not happen if it is a reasonably shaped triangle.

***Insufficient memory on accelerator***

There is insufficient memory on one of the devices specified for the gpu package

***Internal error in atom\_style body***

This error should not occur. Contact the developers.

***Invalid -reorder N value***

Self-explanatory.

***Invalid Angles section in molecule file***

Self-explanatory.

***Invalid Bonds section in molecule file***

Self-explanatory.

***Invalid Boolean syntax in if command***

Self-explanatory.

***Invalid Charges section in molecule file***

Self-explanatory.

***Invalid Coords section in molecule file***

Self-explanatory.

***Invalid Diameters section in molecule file***

Self-explanatory.

***Invalid Dihedrals section in molecule file***

Self-explanatory.

***Invalid Improvers section in molecule file***

Self-explanatory.

***Invalid Kokkos command-line args***

Self-explanatory. See Section 2.7 of the manual for details.

***Invalid LAMMPS restart file***

The file does not appear to be a LAMMPS restart file since it does not contain the correct magic string at the beginning.

***Invalid Masses section in molecule file***

Self-explanatory.

***Invalid molecule ID in molecule file***

Molecule ID must be a non-zero positive integer.

***Invalid Molecules section in molecule file***

Self-explanatory.

***Invalid REAX atom type***

There is a mis-match between LAMMPS atom types and the elements listed in the ReaxFF force field file.

***Invalid Special Bond Counts section in molecule file***

Self-explanatory.

***Invalid Types section in molecule file***

Self-explanatory.

***Invalid angle count in molecule file***

Self-explanatory.

***Invalid angle table length***

Length must be 2 or greater.

***Invalid angle type in Angles section of data file***

Angle type must be positive integer and within range of specified angle types.

***Invalid angle type in Angles section of molecule file***

Self-explanatory.

***Invalid angle type index for fix shake***

Self-explanatory.

***Invalid args for non-hybrid pair coefficients***

“NULL” is only supported in pair\_coeff calls when using pair hybrid

***Invalid argument to factorial %d***

N must be  $\geq 0$  and  $\leq 167$ , otherwise the factorial result is too large.

***Invalid atom ID in %s section of data file***

An atom in a section of the data file being read by fix property/atom has an invalid atom ID that is  $\leq 0$  or  $>$  the maximum existing atom ID.

***Invalid atom ID in Angles section of data file***

Atom IDs must be positive integers and within range of defined atoms.

***Invalid atom ID in Angles section of molecule file***

Self-explanatory.

***Invalid atom ID in Atoms section of data file***

Atom IDs must be positive integers.

***Invalid atom ID in Bodies section of data file***

Atom IDs must be positive integers and within range of defined atoms.

***Invalid atom ID in Bonds section of data file***

Atom IDs must be positive integers and within range of defined atoms.

***Invalid atom ID in Bonds section of molecule file***

Self-explanatory.

***Invalid atom ID in Bonus section of data file***

Atom IDs must be positive integers and within range of defined atoms.

***Invalid atom ID in Dihedrals section of data file***

Atom IDs must be positive integers and within range of defined atoms.

***Invalid atom ID in Fragments section of molecule file***

Self-explanatory.

***Invalid atom ID in Improvers section of data file***

Atom IDs must be positive integers and within range of defined atoms.

***Invalid atom ID in Velocities section of data file***

Atom IDs must be positive integers and within range of defined atoms.

***Invalid atom ID in dihedrals section of molecule file***

Self-explanatory.

***Invalid atom ID in improvers section of molecule file***

Self-explanatory.

***Invalid atom ID in variable file***

Self-explanatory.

***Invalid atom IDs in neb file***

An ID in the file was not found in the system.

***Invalid atom diameter in molecule file***

Diameters must be  $\geq 0.0$ .

***Invalid atom mass for fix shake***

Mass specified in fix shake command must be  $> 0.0$ .

***Invalid atom mass in molecule file***

Masses must be  $> 0.0$ .

***Invalid atom type in Atoms section of data file***

Atom types must range from 1 to specified # of types.

***Invalid atom type in create\_atoms command***

The create\_box command specified the range of valid atom types. An invalid type is being requested.

***Invalid atom type in create\_atoms mol command***

The atom types in the defined molecule are added to the value specified in the create\_atoms command, as an offset. The final value for each atom must be between 1 to N, where N is the number of atom types.

***Invalid atom type in fix atom/swap command***

The atom type specified in the atom/swap command does not exist.

***Invalid atom type in fix bond/create command***

Self-explanatory.

***Invalid atom type in fix deposit command***

Self-explanatory.

***Invalid atom type in fix deposit mol command***

The atom types in the defined molecule are added to the value specified in the create\_atoms command, as an offset. The final value for each atom must be between 1 to N, where N is the number of atom types.

***Invalid atom type in fix gcmc command***

The atom type specified in the gcmc command does not exist.

***Invalid atom type in fix pour command***

Self-explanatory.

***Invalid atom type in fix pour mol command***

The atom types in the defined molecule are added to the value specified in the create\_atoms command, as an offset. The final value for each atom must be between 1 to N, where N is the number of atom types.

***Invalid atom type in molecule file***

Atom types must range from 1 to specified # of types.

***Invalid atom type in neighbor exclusion list***

Atom types must range from 1 to Ntypes inclusive.

***Invalid atom type index for fix shake***

Atom types must range from 1 to Ntypes inclusive.

***Invalid atom types in pair\_write command***

Atom types must range from 1 to Ntypes inclusive.

***Invalid atom vector in variable formula***

The atom vector is not recognized.

***Invalid atom\_style body command***

No body style argument was provided.

***Invalid atom\_style command***

Self-explanatory.

***Invalid attribute in dump custom command***

Self-explanatory.

***Invalid attribute in dump local command***

Self-explanatory.

***Invalid attribute in dump modify command***

Self-explanatory.

***Invalid basis setting in create\_atoms command***

The basis index must be between 1 to N where N is the number of basis atoms in the lattice. The type index must be between 1 to N where N is the number of atom types.

***Invalid basis setting in fix append/atoms command***

The basis index must be between 1 to N where N is the number of basis atoms in the lattice. The type index must be between 1 to N where N is the number of atom types.

***Invalid bin bounds in compute chunk/atom***

The lo/hi values are inconsistent.

***Invalid bin bounds in fix ave/spatial***

The lo/hi values are inconsistent.

***Invalid body nparticle command***

Arguments in atom-style command are not correct.

***Invalid bond count in molecule file***

Self-explanatory.

***Invalid bond table length***

Length must be 2 or greater.

***Invalid bond type in Bonds section of data file***

Bond type must be positive integer and within range of specified bond types.

***Invalid bond type in Bonds section of molecule file***

Self-explanatory.

***Invalid bond type in create\_bonds command***

Self-explanatory.

***Invalid bond type in fix bond/break command***

Self-explanatory.

***Invalid bond type in fix bond/create command***

Self-explanatory.

***Invalid bond type index for fix shake***

Self-explanatory. Check the fix shake command in the input script.

***Invalid coeffs for this dihedral style***

Cannot set class 2 coeffs in data file for this dihedral style.

***Invalid color in dump\_modify command***

The specified color name was not in the list of recognized colors. See the dump\_modify doc page.

***Invalid color map min/max values***

The min/max values are not consistent with either each other or with values in the color map.

***Invalid command-line argument***

One or more command-line arguments is invalid. Check the syntax of the command you are using to launch LAMMPS.

***Invalid compute ID in variable formula***

The compute is not recognized.

***Invalid create\_atoms rotation vector for 2d model***

The rotation vector can only have a z component.

***Invalid custom OpenCL parameter string.***

There are not enough or too many parameters in the custom string for package GPU.

***Invalid cutoff in comm\_modify command***

Specified cutoff must be  $\geq 0.0$ .

***Invalid cutoffs in pair\_write command***

Inner cutoff must be larger than 0.0 and less than outer cutoff.

***Invalid d1 or d2 value for pair colloid coeff***

Neither d1 or d2 can be  $< 0$ .

***Invalid data file section: Angle Coeffs***

Atom style does not allow angles.

***Invalid data file section: AngleAngle Coeffs***

Atom style does not allow impropers.

***Invalid data file section: AngleAngleTorsion Coeffs***

Atom style does not allow dihedrals.

***Invalid data file section: AngleTorsion Coeffs***

Atom style does not allow dihedrals.

***Invalid data file section: Angles***

Atom style does not allow angles.

***Invalid data file section: Bodies***

Atom style does not allow bodies.

***Invalid data file section: Bond Coeffs***

Atom style does not allow bonds.

***Invalid data file section: BondAngle Coeffs***

Atom style does not allow angles.

***Invalid data file section: BondBond Coeffs***

Atom style does not allow angles.

***Invalid data file section: BondBond13 Coeffs***

Atom style does not allow dihedrals.

***Invalid data file section: Bonds***

Atom style does not allow bonds.

***Invalid data file section: Dihedral Coeffs***

Atom style does not allow dihedrals.

***Invalid data file section: Dihedrals***

Atom style does not allow dihedrals.

***Invalid data file section: Ellipsoids***

Atom style does not allow ellipsoids.

***Invalid data file section: EndBondTorsion Coeffs***

Atom style does not allow dihedrals.

***Invalid data file section: Improper Coeffs***

Atom style does not allow impropers.

***Invalid data file section: Impropers***

Atom style does not allow impropers.

***Invalid data file section: Lines***

Atom style does not allow lines.

***Invalid data file section: MiddleBondTorsion Coeffs***

Atom style does not allow dihedrals.

***Invalid data file section: Triangles***

Atom style does not allow triangles.

***Invalid delta\_conf in tad command***

The value must be between 0 and 1 inclusive.

***Invalid density in Atoms section of data file***

Density value cannot be  $\leq 0.0$ .

***Invalid density in set command***

Density must be  $> 0.0$ .

***Invalid diameter in set command***

Self-explanatory.

***Invalid dihedral count in molecule file***

Self-explanatory.

***Invalid dihedral type in Dihedrals section of data file***

Dihedral type must be positive integer and within range of specified dihedral types.

***Invalid dihedral type in dihedrals section of molecule file***

Self-explanatory.

***Invalid dipole length in set command***

Self-explanatory.

***Invalid displace\_atoms rotate axis for 2d***

Axis must be in z direction.

***Invalid dump dcd filename***

Filenames used with the dump dcd style cannot be binary or compressed or cause multiple files to be written.

***Invalid dump frequency***

Dump frequency must be 1 or greater.

***Invalid dump image element name***

The specified element name was not in the standard list of elements. See the dump\_modify doc page.

***Invalid dump image filename***

The file produced by dump image cannot be binary and must be for a single processor.

***Invalid dump image theta value***

Theta must be between 0.0 and 180.0 inclusive.

***Invalid dump image zoom value***

Zoom value must be > 0.0.

***Invalid dump movie filename***

The file produced by dump movie cannot be binary or compressed and must be a single file for a single processor.

***Invalid dump xtc filename***

Filenames used with the dump xtc style cannot be binary or compressed or cause multiple files to be written.

***Invalid dump xyz filename***

Filenames used with the dump xyz style cannot be binary or cause files to be written by each processor.

***Invalid dump\_modify threshold operator***

Operator keyword used for threshold specification in not recognized.

***Invalid entry in -reorder file***

Self-explanatory.

***Invalid fix ID in variable formula***

The fix is not recognized.

***Invalid fix ave/time off column***

Self-explanatory.

***Invalid fix box/relax command for a 2d simulation***

Fix box/relax styles involving the z dimension cannot be used in a 2d simulation.

***Invalid fix box/relax command pressure settings***

If multiple dimensions are coupled, those dimensions must be specified.

***Invalid fix box/relax pressure settings***

Settings for coupled dimensions must be the same.

***Invalid fix halt attribute***

Self-explanatory.



***Invalid fix halt operator***

Self-explanatory.

***Invalid fix nvt/npt/nph command for a 2d simulation***

Cannot control z dimension in a 2d model.

***Invalid fix nvt/npt/nph command pressure settings***

If multiple dimensions are coupled, those dimensions must be specified.

***Invalid fix nvt/npt/nph pressure settings***

Settings for coupled dimensions must be the same.

***Invalid fix press/berendsen for a 2d simulation***

The z component of pressure cannot be controlled for a 2d model.

***Invalid fix press/berendsen pressure settings***

Settings for coupled dimensions must be the same.

***Invalid fix qeq parameter file***

Element index > number of atom types.

***Invalid fix rigid npt/nph command for a 2d simulation***

Cannot control z dimension in a 2d model.

***Invalid fix rigid npt/nph command pressure settings***

If multiple dimensions are coupled, those dimensions must be specified.

***Invalid fix rigid/small npt/nph command for a 2d simulation***

Cannot control z dimension in a 2d model.

***Invalid fix rigid/small npt/nph command pressure settings***

If multiple dimensions are coupled, those dimensions must be specified.

***Invalid flag in force field section of restart file***

Unrecognized entry in restart file.

***Invalid flag in header section of restart file***

Unrecognized entry in restart file.

***Invalid flag in peratom section of restart file***

The format of this section of the file is not correct.

***Invalid flag in type arrays section of restart file***

Unrecognized entry in restart file.

***Invalid frequency in temper command***

Nevery must be > 0.

***Invalid group ID in neigh\_modify command***

A group ID used in the neigh\_modify command does not exist.

***Invalid group function in variable formula***

Group function is not recognized.

***Invalid group in comm\_modify command***

Self-explanatory.

***Invalid image up vector***

Up vector cannot be (0,0,0).

***Invalid immediate variable***

Syntax of immediate value is incorrect.

***Invalid improper count in molecule file***

Self-explanatory.

***Invalid improper type in Improvers section of data file***

Improper type must be positive integer and within range of specified improper types.

***Invalid improper type in improvers section of molecule file***

Self-explanatory.

***Invalid index for non-body particles in compute body/local command***

Only indices 1,2,3 can be used for non-body particles.

***Invalid index in compute body/local command***

Self-explanatory.

***Invalid is\_active() function in variable formula***

Self-explanatory.

***Invalid is\_available() function in variable formula***

Self-explanatory.

***Invalid is\_defined() function in variable formula***

Self-explanatory.

***Invalid keyword in angle table parameters***

Self-explanatory.

***Invalid keyword in bond table parameters***

Self-explanatory.

***Invalid keyword in compute angle/local command***

Self-explanatory.

***Invalid keyword in compute bond/local command***

Self-explanatory.

***Invalid keyword in compute dihedral/local command***

Self-explanatory.

***Invalid keyword in compute improper/local command***

Self-explanatory.

***Invalid keyword in compute pair/local command***

Self-explanatory.

***Invalid keyword in compute property/atom command***

Self-explanatory.

***Invalid keyword in compute property/chunk command***

Self-explanatory.

***Invalid keyword in compute property/local command***

Self-explanatory.

***Invalid keyword in dump cfg command***

Self-explanatory.

***Invalid keyword in pair table parameters***

Keyword used in list of table parameters is not recognized.

***Invalid length in set command***

Self-explanatory.

***Invalid mass in set command***

Self-explanatory.

***Invalid mass line in data file***

Self-explanatory.

***Invalid mass value***

Self-explanatory.

***Invalid math function in variable formula***

Self-explanatory.

***Invalid math/group/special function in variable formula***

Self-explanatory.

***Invalid option in lattice command for non-custom style***

Certain lattice keywords are not supported unless the lattice style is “custom”.

***Invalid order of forces within respa levels***

For respa, ordering of force computations within respa levels must obey certain rules. E.g. bonds cannot be compute less frequently than angles, pairwise forces cannot be computed less frequently than kspace, etc.

***Invalid pair table cutoff***

Cutoffs in pair\_coeff command are not valid with read-in pair table.

***Invalid pair table length***

Length of read-in pair table is invalid

***Invalid param file for fix qeq/shielded***

Invalid value of gamma.

***Invalid param file for fix qeq/slater***

Zeta value is 0.0.

***Invalid partitions in processors part command***

Valid partitions are numbered 1 to N and the sender and receiver cannot be the same partition.

***Invalid python command***

Self-explanatory. Check the input script syntax and compare to the documentation for the command. You can use -echo screen as a command-line option when running LAMMPS to see the offending line.

***Invalid radius in Atoms section of data file***

Radius must be  $\geq 0.0$ .

***Invalid random number seed in fix ttm command***

Random number seed must be  $> 0$ .

***Invalid random number seed in set command***

Random number seed must be  $> 0$ .

***Invalid replace values in compute reduce***

Self-explanatory.

***Invalid rigid body ID in fix rigid file***

The ID does not match the number of an existing ID of rigid bodies that are defined by the fix rigid command.

***Invalid rigid body ID in fix rigid/small file***

The ID does not match the number of an existing ID of rigid bodies that are defined by the fix rigid/small command.

***Invalid run command N value***

The number of timesteps must fit in a 32-bit integer. If you want to run for more steps than this, perform multiple shorter runs.

***Invalid run command start/stop value***

Self-explanatory.

***Invalid run command upto value***

Self-explanatory.

***Invalid seed for Marsaglia random # generator***

The initial seed for this random number generator must be a positive integer less than or equal to 900 million.

***Invalid seed for Park random # generator***

The initial seed for this random number generator must be a positive integer.

***Invalid shake angle type in molecule file***

Self-explanatory.

***Invalid shake atom in molecule file***

Self-explanatory.

***Invalid shake bond type in molecule file***

Self-explanatory.

***Invalid shake flag in molecule file***

Self-explanatory.

***Invalid shape in Ellipsoids section of data file***

Self-explanatory.

***Invalid shape in Triangles section of data file***

Two or more of the triangle corners are duplicate points.

***Invalid shape in set command***

Self-explanatory.

***Invalid shear direction for fix wall/gran***

Self-explanatory.

***Invalid special atom index in molecule file***

Self-explanatory.

***Invalid special function in variable formula***

Self-explanatory.

***Invalid style in pair\_write command***

Self-explanatory. Check the input script.

***Invalid syntax in variable formula***

Self-explanatory.

***Invalid t\_event in prd command***

Self-explanatory.

***Invalid t\_event in tad command***

The value must be greater than 0.

***Invalid template atom in Atoms section of data file***

The atom indices must be between 1 to N, where N is the number of atoms in the template molecule the atom belongs to.

***Invalid template index in Atoms section of data file***

The template indices must be between 1 to N, where N is the number of molecules in the template.

***Invalid thermo keyword in variable formula***

The keyword is not recognized.

***Invalid threads\_per\_atom specified.***

For 3-body potentials on the GPU, the threads\_per\_atom setting cannot be greater than 4 for NVIDIA GPUs.

***Invalid timestep reset for fix ave/atom***

Resetting the timestep has invalidated the sequence of timesteps this fix needs to process.

***Invalid timestep reset for fix ave/chunk***

Resetting the timestep has invalidated the sequence of timesteps this fix needs to process.

***Invalid timestep reset for fix ave/correlate***

Resetting the timestep has invalidated the sequence of timesteps this fix needs to process.

***Invalid timestep reset for fix ave/histo***

Resetting the timestep has invalidated the sequence of timesteps this fix needs to process.

***Invalid timestep reset for fix ave/spatial***

Resetting the timestep has invalidated the sequence of timesteps this fix needs to process.

***Invalid timestep reset for fix ave/time***

Resetting the timestep has invalidated the sequence of timesteps this fix needs to process.

***Invalid tmax in tad command***

The value must be greater than 0.0.

***Invalid type for mass set***

Mass command must set a type from 1-N where N is the number of atom types.

***Invalid label2type() function syntax in variable formula***

The first argument must be a label map kind (atom, bond, angle, dihedral, or improper) and the second argument must be a valid type label that has been assigned to a numeric type.

***Invalid use of library file() function***

This function is called through the library interface. This error should not occur. Contact the developers if it does.

***Invalid value in set command***

The value specified for the setting is invalid, likely because it is too small or too large.

***Invalid variable evaluation in variable formula***

A variable used in a formula could not be evaluated.

***Invalid variable in next command***

Self-explanatory.

***Invalid variable name***

Variable name used in an input script line is invalid.

***Invalid variable name in variable formula***

Variable name is not recognized.

***Invalid variable style in special function next***

Only file-style or atomfile-style variables can be used with next().

***Invalid variable style with next command***

Variable styles *equal* and *world* cannot be used in a next command.

***Invalid volume in set command***

Volume must be > 0.0.

***Invalid wiggle direction for fix wall/gran***

Self-explanatory.

***Invoked angle equil angle on angle style none***

Self-explanatory.

***Invoked angle single on angle style none***

Self-explanatory.

***Invoked bond equil distance on bond style none***

Self-explanatory.

***Invoked bond single on bond style none***

Self-explanatory.

***Invoked pair single on pair style none***

A command (e.g. a dump) attempted to invoke the single() function on a pair style none, which is illegal. You are probably attempting to compute per-atom quantities with an undefined pair style.

***Invoking coulombic in pair style lj/coul requires atom attribute q***

The atom style defined does not have this attribute.

***Invoking coulombic in pair style lj/long/dipole/long requires atom attribute q***

The atom style defined does not have these attributes.

***KIM Simulator Model has no Model definition***

There is no model definition (key: model-defn) in the KIM Simulator Model. Please contact the OpenKIM database maintainers to verify and potentially correct this.

***KOKKOS package does not yet support comm\_style tiled***

Self-explanatory.

***KOKKOS package requires a kokkos enabled atom\_style***

Self-explanatory.

***KSpace accuracy must be > 0***

The kspace accuracy designated in the input must be greater than zero.

***KSpace accuracy too large to estimate G vector***

Reduce the accuracy request or specify gewald explicitly via the kspace\_modify command.

***KSpace accuracy too low***

Requested accuracy must be less than 1.0.

***KSpace solver requires a pair style***

No pair style is defined.

***KSpace style does not yet support triclinic geometries***

The specified kspace style does not allow for non-orthogonal simulation boxes.

***KSpace style has not yet been set***

Cannot use kspace\_modify command until a kspace style is set.

***KSpace style is incompatible with Pair style***

Setting a kspace style requires that a pair style with matching long-range Coulombic or dispersion components be used.

***Keyword %s in MEAM parameter file not recognized***

Self-explanatory.

***Kokkos has been compiled for CUDA but no GPUs are requested***

One or more GPUs must be used when Kokkos is compiled for CUDA.

***Kspace\_modify mesh parameter must be all zero or all positive***

Valid kspace mesh parameters are >0. The code will try to auto-detect suitable values when all three mesh sizes are set to zero (the default).

***Kspace\_modify mesh/disp parameter must be all zero or all positive***

Valid kspace mesh/disp parameters are >0. The code will try to auto-detect suitable values when all three mesh sizes are set to zero **and** the required accuracy via *force/disp/real* as well as *force/disp/kspace* is set.

***Kspace style does not support compute group/group***

Self-explanatory.

***Kspace style ppm/disp/tip4p requires newton on***

Self-explanatory.

***Kspace style ppm/tip4p requires newton on***

Self-explanatory.

***Kspace style requires atom attribute q***

The atom style defined does not have these attributes.

***Kspace\_modify eigtol must be smaller than one***

Self-explanatory.

***LAMMPS is not built with Python embedded***

This is done by including the PYTHON package before LAMMPS is built. This is required to use python-style variables.

***LAMMPS unit\_style lj not supported by KIM models***

Self-explanatory. Check the input script or data file.

***LJ6 off not supported in pair\_style buck/long/coul/long***

Self-explanatory.

***Label map is incomplete: all types must be assigned a unique type label***

For a given type-kind (atom types, bond types, etc.) to be written to the data file, all associated types must be assigned a type label, and each type label can be assigned to only one numeric type.

***Label wasn't found in input script***

Self-explanatory.

***Labelmap command before simulation box is defined***

The labelmap command cannot be used before a read\_data, read\_restart, or create\_box command.

***Lattice orient vectors are not orthogonal***

The three specified lattice orientation vectors must be mutually orthogonal.

***Lattice orient vectors are not right-handed***

The three specified lattice orientation vectors must create a right-handed coordinate system such that  $a_1 \times a_2 = a_3$ .

***Lattice primitive vectors are collinear***

The specified lattice primitive vectors do not form a unit cell with non-zero volume.

***Lattice settings are not compatible with 2d simulation***

One or more of the specified lattice vectors has a non-zero z component.

***Lattice spacings are invalid***

Each x,y,z spacing must be > 0.

***Lattice style incompatible with simulation dimension***

2d simulation can use sq, sq2, or hex lattice. 3d simulation can use sc, bcc, or fcc lattice.

***Log of zero/negative value in variable formula***

Self-explanatory.

***Lost atoms via balance: original %ld current %ld***

This should not occur. Report the problem to the developers.

***Lost atoms: original %ld current %ld***

Lost atoms are checked for each time thermo output is done. See the thermo\_modify lost command for options. Lost atoms usually indicate bad dynamics, e.g. atoms have been blown far out of the simulation box, or moved further than one processor's sub-domain away before reneighboring.

***MEAM library error %d***

A call to the MEAM Fortran library returned an error.

***MPI\_LMP\_BIGINT and bigint in lmptype.h are not compatible***

The size of the MPI datatype does not match the size of a bigint.

***MPI\_LMP\_TAGINT and tagint in lmptype.h are not compatible***

The size of the MPI datatype does not match the size of a tagint.

***MSM can only currently be used with comm\_style brick***

This is a current restriction in LAMMPS.

***MSM grid is too large***

The global MSM grid is larger than OFFSET in one or more dimensions. OFFSET is currently set to 16384. You likely need to decrease the requested accuracy.

***MSM order must be 4, 6, 8, or 10***

This is a limitation of the MSM implementation in LAMMPS: the MSM order can only be 4, 6, 8, or 10.

***Mass command before simulation box is defined***

The mass command cannot be used before a read\_data, read\_restart, or create\_box command.

***Matrix factorization to split dispersion coefficients failed***

This should not normally happen. Contact the developers.

***Min\_style command before simulation box is defined***

The min\_style command cannot be used before a read\_data, read\_restart, or create\_box command.

***Minimization could not find thermo\_pe compute***

This compute is created by the thermo command. It must have been explicitly deleted by a uncompute command.

***Minimize command before simulation box is defined***

The minimize command cannot be used before a read\_data, read\_restart, or create\_box command.

***Mismatched brackets in variable***

Self-explanatory.

***Mismatched compute in variable formula***

A compute is referenced incorrectly or a compute that produces per-atom values is used in an equal-style variable formula.

***Mismatched fix in variable formula***

A fix is referenced incorrectly or a fix that produces per-atom values is used in an equal-style variable formula.

***Mismatched parameter in MEAM library file: z!=lat***

The coordination number and lattice do not match, check that consistent values are given.

***Mismatched variable in variable formula***

A variable is referenced incorrectly or an atom-style variable that produces per-atom values is used in an equal-style variable formula.

***Modulo 0 in variable formula***

Self-explanatory.

***Molecule IDs too large for compute chunk/atom***

The IDs must not be larger than can be stored in a 32-bit integer since chunk IDs are 32-bit integers.



***Molecule auto special bond generation overflow***

Counts exceed maxspecial setting for other atoms in system.

***Molecule file has angles but no nangles setting***

Self-explanatory.

***Molecule file has body params but no setting for them***

Self-explanatory.

***Molecule file has bonds but no nbonds setting***

Self-explanatory.

***Molecule file has dihedrals but no ndihedrals setting***

Self-explanatory.

***Molecule file has fragments but no nfragments setting***

Self-explanatory.

***Molecule file has impropers but no nimpropers setting***

Self-explanatory.

***Molecule file has no Body Doubles section***

Self-explanatory.

***Molecule file has no Body Integers section***

Self-explanatory.

***Molecule file has no Fragments section***

Self-explanatory.

***Molecule file has special flags but no bonds***

Self-explanatory.

***Molecule file needs both Special Bond sections***

Self-explanatory.

***Molecule file requires atom style body***

Self-explanatory.

***Molecule file shake flags not before shake atoms***

The order of the two sections is important.

***Molecule file shake flags not before shake bonds***

The order of the two sections is important.

***Molecule file shake info is incomplete***

All 3 SHAKE sections are needed.

***Molecule file special list does not match special count***

The number of values in an atom's special list does not match count.

***Molecule file z center-of-mass must be 0.0 for 2d***

Self-explanatory.

***Molecule file z coord must be 0.0 for 2d***

Self-explanatory.

***Molecule natoms must be 1 for body particle***

Self-explanatory.

***Molecule sizescale must be 1.0 for body particle***

Self-explanatory.

***Molecule template ID for atom\_style template does not exist***

Self-explanatory.

***Molecule template ID for create\_atoms does not exist***

Self-explanatory.

***Molecule template ID for fix deposit does not exist***

Self-explanatory.

***Molecule template ID for fix gcmc does not exist***

Self-explanatory.

***Molecule template ID for fix pour does not exist***

Self-explanatory.

***Molecule template ID for fix rigid/small does not exist***

Self-explanatory.

***Molecule template ID for fix shake does not exist***

Self-explanatory.

***Molecule template ID must be alphanumeric or underscore characters***

Self-explanatory.

***Molecule topology/atom exceeds system topology/atom***

The number of bonds, angles, etc per-atom in the molecule exceeds the system setting. See the create\_box command for how to specify these values.

***Molecule topology type exceeds system topology type***

The number of bond, angle, etc types in the molecule exceeds the system setting. See the create\_box command for how to specify these values.

***More than one fix deform***

Only one fix deform can be defined at a time.

***More than one fix freeze***

Only one of these fixes can be defined, since the granular pair potentials access it.

***More than one fix shake***

Only one fix shake can be defined.

***Mu not allowed when not using semi-grand in fix atom/swap command***

Self-explanatory.

***Must define angle\_style before Angle Coeffs***

Must use an angle\_style command before reading a data file that defines Angle Coeffs.

***Must define angle\_style before BondAngle Coeffs***

Must use an angle\_style command before reading a data file that defines Angle Coeffs.

***Must define angle\_style before BondBond Coeffs***

Must use an angle\_style command before reading a data file that defines Angle Coeffs.

***Must define bond\_style before Bond Coeffs***

Must use a bond\_style command before reading a data file that defines Bond Coeffs.

***Must define dihedral\_style before AngleAngleTorsion Coeffs***

Must use a dihedral\_style command before reading a data file that defines AngleAngleTorsion Coeffs.

***Must define dihedral\_style before AngleTorsion Coeffs***

Must use a dihedral\_style command before reading a data file that defines AngleTorsion Coeffs.

***Must define dihedral\_style before BondBond13 Coeffs***

Must use a dihedral\_style command before reading a data file that defines BondBond13 Coeffs.

***Must define dihedral\_style before Dihedral Coeffs***

Must use a dihedral\_style command before reading a data file that defines Dihedral Coeffs.

***Must define dihedral\_style before EndBondTorsion Coeffs***

Must use a dihedral\_style command before reading a data file that defines EndBondTorsion Coeffs.

***Must define dihedral\_style before MiddleBondTorsion Coeffs***

Must use a dihedral\_style command before reading a data file that defines MiddleBondTorsion Coeffs.

***Must define improper\_style before AngleAngle Coeffs***

Must use an improper\_style command before reading a data file that defines AngleAngle Coeffs.

***Must define improper\_style before Improper Coeffs***

Must use an improper\_style command before reading a data file that defines Improper Coeffs.

***Must define pair\_style before Pair Coeffs***

Must use a pair\_style command before reading a data file that defines Pair Coeffs.

***Must define pair\_style before PairIJ Coeffs***

Must use a pair\_style command before reading a data file that defines PairIJ Coeffs.

***Must have more than one processor partition to temper***

Cannot use the temper command with only one processor partition. Use the -partition command-line option.

***Must not have multiple fixes change box parameter ...***

Self-explanatory.

***Must read Angle Type Labels before Angles***

An Angle Type Labels section of a data file must come before the Angles section.

***Must read Atom Type Labels before Atoms***

An Atom Type Labels section of a data file must come before the Atoms section.

***Must read Atoms before Angles***

The Atoms section of a data file must come before an Angles section.

***Must read Atoms before Bodies***

The Atoms section of a data file must come before a Bodies section.

***Must read Atoms before Bonds***

The Atoms section of a data file must come before a Bonds section.

***Must read Atoms before Dihedrals***

The Atoms section of a data file must come before a Dihedrals section.

***Must read Atoms before Ellipsoids***

The Atoms section of a data file must come before a Ellipsoids section.

***Must read Atoms before Improvers***

The Atoms section of a data file must come before an Improvers section.

***Must read Atoms before Lines***

The Atoms section of a data file must come before a Lines section.

***Must read Atoms before Triangles***

The Atoms section of a data file must come before a Triangles section.

***Must read Atoms before Velocities***

The Atoms section of a data file must come before a Velocities section.

***Must read Bond Type Labels before Bonds***

A Bond Type Labels section of a data file must come before the Bonds section.

***Must read Dihedral Type Labels before Dihedrals***

An Dihedral Type Labels section of a data file must come before the Dihedrals section.

***Must read Improper Type Labels before Improvers***

An Improper Type Labels section of a data file must come before the Improvers section.

***Must re-specify non-restarted pair style (xxx) after read\_restart***

For pair styles, that do not store their settings in a restart file, it must be defined with a new 'pair\_style' command after read\_restart.

***Must set both respa inner and outer***

Cannot use just the inner or outer option with respa without using the other.

***Must set number of threads via package omp command***

Because you are using the OPENMP package, set the number of threads via its settings, not by the pair\_style snap nthreads setting.

***Must shrink-wrap piston boundary***

The boundary style of the face where the piston is applied must be of type s (shrink-wrapped).

***Must specify a region in fix deposit***

The region keyword must be specified with this fix.

***Must specify a region in fix pour***

Self-explanatory.

***Must specify at least 2 types in fix atom/swap command***

Self-explanatory.

***Must use 'kim\_style init' command before simulation box is defined***

Self-explanatory.

***Must use 'kim\_style define' command after simulation box is defined***

Self-explanatory.

***Must use 'kim\_style init' command before 'kim\_style define'***

Self-explanatory.

***Must use 'kspace\_modify pressure/scalar no' for rRESPA with kspace\_style MSM***

The kspace scalar pressure option cannot (yet) be used with rRESPA.

***Must use 'kspace\_modify pressure/scalar no' for tensor components with kspace\_style msm***

Otherwise MSM will compute only a scalar pressure. See the kspace\_modify command for details on this setting.

***Must use 'kspace\_modify pressure/scalar no' to obtain per-atom virial with kspace\_style MSM***

The kspace scalar pressure option cannot be used to obtain per-atom virial.

***Must use 'kspace\_modify pressure/scalar no' with GPU MSM Pair styles***

The kspace scalar pressure option is not (yet) compatible with GPU MSM Pair styles.

***Must use 'kspace\_modify pressure/scalar no' with kspace\_style msm/cg***

The kspace scalar pressure option is not compatible with kspace\_style msm/cg.

***Must use -in switch with multiple partitions***

A multi-partition simulation cannot read the input script from stdin. The -in command-line option must be used to specify a file.

***Must use Kokkos half/thread or full neighbor list with threads or GPUs***

Using Kokkos half-neighbor lists with threading is not allowed.

***Must use a block or cylinder region with fix pour***

Self-explanatory.

***Must use a block region with fix pour for 2d simulations***

Self-explanatory.

***Must use a bond style with TIP4P potential***

TIP4P potentials assume bond lengths in water are constrained by a fix shake command.

***Must use a molecular atom style with fix poems molecule***

Self-explanatory.

***Must use a z-axis cylinder region with fix pour***

Self-explanatory.

***Must use an angle style with TIP4P potential***

TIP4P potentials assume angles in water are constrained by a fix shake command.

***Must use atom map style array with Kokkos***

See the atom\_modify map command.

***Must use atom style with molecule IDs with fix bond/swap***

Self-explanatory.

***Must use pair\_style comb or comb3 with fix qeq/comb***

Self-explanatory.

***Must use variable energy with fix addforce***

Must define an energy variable when applying a dynamic force during minimization.

***Must use variable energy with fix efield***

You must define an energy when performing a minimization with a variable E-field.

***NEB command before simulation box is defined***

Self-explanatory.

***NEB requires damped dynamics minimizer***

Use a different minimization style.

***NEB requires use of fix neb***

Self-explanatory.

***NL ramp in wall/piston only implemented in zlo for now***

The ramp keyword can only be used for piston applied to face zlo.

***Need nswaptypes mu values in fix atom/swap command***

Self-explanatory.

***Needed bonus data not in data file***

Some atom styles require bonus data. See the read\_data page for details.

***Needed molecular topology not in data file***

The header of the data file indicated bonds, angles, etc would be included, but they are not present.

***Neigh\_modify exclude molecule requires atom attribute molecule***

Self-explanatory.

***Neigh\_modify include group != atom\_modify first group***

Self-explanatory.

***Neighbor delay must be 0 or multiple of every setting***

The delay and every parameters set via the neigh\_modify command are inconsistent. If the delay setting is non-zero, then it must be a multiple of the every setting.

***Neighbor include group not allowed with ghost neighbors***

This is a current restriction within LAMMPS.

***Neighbor list overflow, boost neigh\_modify one***

There are too many neighbors of a single atom. Use the neigh\_modify command to increase the max number of neighbors allowed for one atom. You may also want to boost the page size.

***Neighbor multi not yet enabled for ghost neighbors***

This is a current restriction within LAMMPS.

***Neighbor multi not yet enabled for granular***

Self-explanatory.

***Neighbor multi not yet enabled for rRESPA***

Self-explanatory.

***Neighbor page size must be >= 10x the one atom setting***

This is required to prevent wasting too much memory.

***New atom IDs exceed maximum allowed ID***

See the setting for tagint in the src/lmptype.h file.

***New bond exceeded bonds per atom in create\_bonds*** See the read\_data command for info on using the “extra/bond/per/atom” keyword to allow for additional bonds to be formed

***New bond exceeded bonds per atom in fix bond/create***

See the read\_data command for info on using the “extra/bond/per/atom” keyword to allow for additional bonds to be formed

***New bond exceeded special list size in fix bond/create***

See the “read\_data extra/special/per/atom” command (or the “create\_box extra/special/per/atom” command) for info on how to leave space in the special bonds list to allow for additional bonds to be formed.

***Newton bond change after simulation box is defined***

The newton command cannot be used to change the newton bond value after a read\_data, read\_restart, or create\_box command.

***Next command must list all universe and uloop variables***

This is to insure they stay in sync.

***No Kspace style defined for compute group/group***

Self-explanatory.

***No OpenMP support compiled in***

An OpenMP flag is set, but LAMMPS was not built with OpenMP support.

***No angle style is defined for compute angle/local***

Self-explanatory.

***No angles allowed with this atom style***

Self-explanatory.

***No atoms in data file***

The header of the data file indicated that atoms would be included, but they are not present.

***No basis atoms in lattice***

Basis atoms must be defined for lattice style user.

***No bodies allowed with this atom style***

Self-explanatory. Check data file.

***No bond style is defined for compute bond/local***

Self-explanatory.

***No bonds allowed with this atom style***

Self-explanatory.

***No box information in dump. You have to use 'box no'***

Self-explanatory.

***No count or invalid atom count in molecule file***

The number of atoms must be specified.

***No dihedral style is defined for compute dihedral/local***

Self-explanatory.

***No dihedrals allowed with this atom style***

Self-explanatory.

***No dump custom arguments specified***

The dump custom command requires that atom quantities be specified to output to dump file.

***No dump local arguments specified***

Self-explanatory.

***No ellipsoids allowed with this atom style***

Self-explanatory. Check data file.

***No fix gravity defined for fix pour***

Gravity is required to use fix pour.

***No improper style is defined for compute improper/local***

Self-explanatory.

***No impropers allowed with this atom style***

Self-explanatory.

***No input values for fix ave/spatial***

Self-explanatory.

***No lines allowed with this atom style***

Self-explanatory. Check data file.

***No matching element in ADP potential file***

The ADP potential file does not contain elements that match the requested elements.

***No matching element in EAM potential file***

The EAM potential file does not contain elements that match the requested elements.

***No molecule topology allowed with atom style template***

The data file cannot specify the number of bonds, angles, etc, because this info is inferred from the molecule templates.

***No overlap of box and region for create\_atoms***

Self-explanatory.

***No pair coul/streitz, for fix qeq/slater***

These commands must be used together.

***No pair hbond/dreiding coefficients set***

Self-explanatory.

***No pair style defined for compute group/group***

Cannot calculate group interactions without a pair style defined.

***No pair style is defined for compute pair/local***

Self-explanatory.

***No pair style is defined for compute property/local***

Self-explanatory.

***No rigid bodies defined***

The fix specification did not end up defining any rigid bodies.

***No triangles allowed with this atom style***

Self-explanatory. Check data file.

***No values in fix ave/chunk command***

Self-explanatory.

***No values in fix ave/time command***

Self-explanatory.

***Non digit character between brackets in variable***

Self-explanatory.

***Non integer # of swaps in temper command***

Swap frequency in temper command must evenly divide the total # of timesteps.

***Non-numeric box dimensions - simulation unstable***

The box size has apparently blown up.

***Non-zero atom IDs with atom\_modify id = no***

Self-explanatory.

***Non-zero read\_data shift z value for 2d simulation***

Self-explanatory.

***Nprocs not a multiple of N for -reorder***

Self-explanatory.

***Number of core atoms != number of shell atoms***

There must be a one-to-one pairing of core and shell atoms.

***Numeric index is out of bounds***

A command with an argument that specifies an integer or range of integers is using a value that is less than 1 or greater than the maximum allowed limit.

***One or more Atom IDs is negative***

Atom IDs must be positive integers.

***One or more atom IDs is too big***

The limit on atom IDs is set by the SMALLBIG, BIGBIG, SMALLSMALL setting in your LAMMPS build. See the [Build settings](#) page for more info.

***One or more atom IDs is zero***

Either all atoms IDs must be zero or none of them.

***One or more atoms belong to multiple rigid bodies***

Two or more rigid bodies defined by the fix rigid command cannot contain the same atom.

***One or more rigid bodies are a single particle***

Self-explanatory.

***One or zero atoms in rigid body***

Any rigid body defined by the fix rigid command must contain 2 or more atoms.

***Only 2 types allowed when not using semi-grand in fix atom/swap command***

Self-explanatory.

***Only one cut-off allowed when requesting all long***

Self-explanatory.

***Only one cutoff allowed when requesting all long***

Self-explanatory.



**Only zhi currently implemented for fix append/atoms**

Self-explanatory.

**Out of range atoms - cannot compute MSM**

One or more atoms are attempting to map their charge to a MSM grid point that is not owned by a processor. This is likely for one of two reasons, both of them bad. First, it may mean that an atom near the boundary of a processor's sub-domain has moved more than 1/2 the *neighbor skin distance* without neighbor lists being rebuilt and atoms being migrated to new processors. This also means you may be missing pairwise interactions that need to be computed. The solution is to change the re-neighboring criteria via the *neigh\_modify* command. The safest settings are "delay 0 every 1 check yes". Second, it may mean that an atom has moved far outside a processor's sub-domain or even the entire simulation box. This indicates bad physics, e.g. due to highly overlapping atoms, too large a timestep, etc.

**Out of range atoms - cannot compute PPPM**

One or more atoms are attempting to map their charge to a PPPM grid point that is not owned by a processor. This is likely for one of two reasons, both of them bad. First, it may mean that an atom near the boundary of a processor's sub-domain has moved more than 1/2 the *neighbor skin distance* without neighbor lists being rebuilt and atoms being migrated to new processors. This also means you may be missing pairwise interactions that need to be computed. The solution is to change the re-neighboring criteria via the *neigh\_modify* command. The safest settings are "delay 0 every 1 check yes". Second, it may mean that an atom has moved far outside a processor's sub-domain or even the entire simulation box. This indicates bad physics, e.g. due to highly overlapping atoms, too large a timestep, etc.

**Out of range atoms - cannot compute PPPMDisp**

One or more atoms are attempting to map their charge to a PPPM grid point that is not owned by a processor. This is likely for one of two reasons, both of them bad. First, it may mean that an atom near the boundary of a processor's sub-domain has moved more than 1/2 the *neighbor skin distance* without neighbor lists being rebuilt and atoms being migrated to new processors. This also means you may be missing pairwise interactions that need to be computed. The solution is to change the re-neighboring criteria via the *neigh\_modify* command. The safest settings are "delay 0 every 1 check yes". Second, it may mean that an atom has moved far outside a processor's sub-domain or even the entire simulation box. This indicates bad physics, e.g. due to highly overlapping atoms, too large a timestep, etc.

**Overflow of allocated fix vector storage**

This should not normally happen if the fix correctly calculated how long the vector will grow to. Contact the developers.

**Overlapping large/large in pair colloid**

This potential is infinite when there is an overlap.

**Overlapping small/large in pair colloid**

This potential is infinite when there is an overlap.

**POEMS fix must come before NPT/NPH fix**

NPT/NPH fix must be defined in input script after all poems fixes, else the fix contribution to the pressure virial is incorrect.

**PPPM can only currently be used with comm\_style brick**

This is a current restriction in LAMMPS.

**PPPM grid is too large**

The global PPPM grid is larger than OFFSET in one or more dimensions. OFFSET is currently set to 4096. You likely need to decrease the requested accuracy.

**PPPM grid stencil extends beyond nearest neighbor processor**

This is not allowed if the kspace\_modify overlap setting is no.

**PPPM order < minimum allowed order**

The default minimum order is 2. This can be reset by the kspace\_modify minorder command.

***PPPM order cannot be < 2 or > than %d***

This is a limitation of the PPPM implementation in LAMMPS.

***PPPMDisp Coulomb grid is too large***

The global PPPM grid is larger than OFFSET in one or more dimensions. OFFSET is currently set to 4096. You likely need to decrease the requested accuracy.

***PPPMDisp Dispersion grid is too large***

The global PPPM grid is larger than OFFSET in one or more dimensions. OFFSET is currently set to 4096. You likely need to decrease the requested accuracy.

***PPPMDisp can only currently be used with comm\_style brick***

This is a current restriction in LAMMPS.

***PPPMDisp coulomb order cannot be greater than %d***

This is a limitation of the PPPM implementation in LAMMPS.

***PPPMDisp used but no parameters set, for further information please see the pppm/disp documentation***

An efficient and accurate usage of the pppm/disp requires settings via the kspace\_modify command. Please see the pppm/disp documentation for further instructions.

***PRD command before simulation box is defined***

The prd command cannot be used before a read\_data, read\_restart, or create\_box command.

***PRD nsteps must be multiple of t\_event***

Self-explanatory.

***PRD t\_corr must be multiple of t\_event***

Self-explanatory.

***Package command after simulation box is defined***

The package command cannot be used after a read\_data, read\_restart, or create\_box command.

***Package gpu command without GPU package installed***

The GPU package must be installed via “make yes-gpu” before LAMMPS is built.

***Package intel command without INTEL package installed***

The INTEL package must be installed via “make yes-intel” before LAMMPS is built.

***Package kokkos command without KOKKOS package enabled***

The KOKKOS package must be installed via “make yes-kokkos” before LAMMPS is built, and the “-k on” must be used to enable the package.

***Package omp command without OPENMP package installed***

The OPENMP package must be installed via “make yes-openmp” before LAMMPS is built.

***Pair body requires atom style body***

Self-explanatory.

***Pair body requires body style nparticle***

This pair style is specific to the nparticle body style.

***Pair brownian requires atom style sphere***

Self-explanatory.

***Pair brownian requires extended particles***

One of the particles has radius 0.0.

***Pair brownian requires monodisperse particles***

All particles must be the same finite size.

***Pair brownian/poly requires atom style sphere***

Self-explanatory.

***Pair brownian/poly requires extended particles***

One of the particles has radius 0.0.

***Pair brownian/poly requires newton pair off***

Self-explanatory.

***Pair coeff for hybrid has invalid style***

Style in pair coeff must have been listed in pair\_style command.

***Pair coul/wolf requires atom attribute q***

The atom style defined does not have this attribute.

***Pair cutoff < Respa interior cutoff***

One or more pairwise cutoffs are too short to use with the specified rRESPA cutoffs.

***Pair dipole/cut requires atom attributes q, mu, torque***

The atom style defined does not have these attributes.

***Pair dipole/cut/gpu requires atom attributes q, mu, torque***

The atom style defined does not have this attribute.

***Pair dipole/long requires atom attributes q, mu, torque***

The atom style defined does not have these attributes.

***Pair dipole/sf/gpu requires atom attributes q, mu, torque***

The atom style defined does not one or more of these attributes.

***Pair distance < table inner cutoff***

Two atoms are closer together than the pairwise table allows.

***Pair distance > table outer cutoff***

Two atoms are further apart than the pairwise table allows.

***Pair dpd requires ghost atoms store velocity***

Use the comm\_modify vel yes command to enable this.

***Pair gayberne epsilon a,b,c coeffs are not all set***

Each atom type involved in pair\_style gayberne must have these 3 coefficients set at least once.

***Pair gayberne requires atom style ellipsoid***

Self-explanatory.

***Pair gayberne requires atoms with same type have same shape***

Self-explanatory.

***Pair gayberne/gpu requires atom style ellipsoid***

Self-explanatory.

***Pair gayberne/gpu requires atoms with same type have same shape***

Self-explanatory.

***Pair granular requires atom attributes radius, rmass***

The atom style defined does not have these attributes.

***Pair granular requires ghost atoms store velocity***

Use the comm\_modify vel yes command to enable this.

***Pair granular with shear history requires newton pair off***

This is a current restriction of the implementation of pair granular styles with history.

***Pair hybrid single calls do not support per sub-style special bond values***

Self-explanatory.

***Pair hybrid sub-style does not support single call***

You are attempting to invoke a single() call on a pair style that does not support it.

***Pair hybrid sub-style is not used***

No pair\_coeff command used a sub-style specified in the pair\_style command.

***Pair inner cutoff < Respa interior cutoff***

One or more pairwise cutoffs are too short to use with the specified rRESPA cutoffs.

***Pair inner cutoff >= Pair outer cutoff***

The specified cutoffs for the pair style are inconsistent.

***Pair line/lj requires atom style line***

Self-explanatory.

***Pair lj/long/dipole/long requires atom attributes mu, torque***

The atom style defined does not have these attributes.

***Pair lubricate requires atom style sphere***

Self-explanatory.

***Pair lubricate requires ghost atoms store velocity***

Use the comm\_modify vel yes command to enable this.

***Pair lubricate requires monodisperse particles***

All particles must be the same finite size.

***Pair lubricate/poly requires atom style sphere***

Self-explanatory.

***Pair lubricate/poly requires extended particles***

One of the particles has radius 0.0.

***Pair lubricate/poly requires ghost atoms store velocity***

Use the comm\_modify vel yes command to enable this.

***Pair lubricate/poly requires newton pair off***

Self-explanatory.

***Pair lubricateU requires atom style sphere***

Self-explanatory.

***Pair lubricateU requires ghost atoms store velocity***

Use the comm\_modify vel yes command to enable this.

***Pair lubricateU requires monodisperse particles***

All particles must be the same finite size.

***Pair lubricateU/poly requires ghost atoms store velocity***

Use the comm\_modify vel yes command to enable this.

***Pair lubricateU/poly requires newton pair off***

Self-explanatory.

***Pair peri lattice is not identical in x, y, and z***

The lattice defined by the lattice command must be cubic.

***Pair peri requires a lattice be defined***

Use the lattice command for this purpose.

***Pair peri requires an atom map, see atom\_modify***

Even for atomic systems, an atom map is required to find Peridynamic bonds. Use the atom\_modify command to define one.

***Pair resquared epsilon a,b,c coeffs are not all set***

Self-explanatory.

***Pair resquared epsilon and sigma coeffs are not all set***

Self-explanatory.

***Pair resquared requires atom style ellipsoid***

Self-explanatory.

***Pair resquared requires atoms with same type have same shape***

Self-explanatory.

***Pair resquared/gpu requires atom style ellipsoid***

Self-explanatory.

***Pair resquared/gpu requires atoms with same type have same shape***

Self-explanatory.

***Pair style AIREBO requires atom IDs***

This is a requirement to use the AIREBO potential.

***Pair style AIREBO requires newton pair on***

See the newton command. This is a restriction to use the AIREBO potential.

***Pair style BOP requires atom IDs***

This is a requirement to use the BOP potential.

***Pair style BOP requires newton pair on***

See the newton command. This is a restriction to use the BOP potential.

***Pair style COMB requires atom IDs***

This is a requirement to use the AIREBO potential.

***Pair style COMB requires atom attribute q***

Self-explanatory.

***Pair style COMB requires newton pair on***

See the newton command. This is a restriction to use the COMB potential.

***Pair style COMB3 requires atom IDs***

This is a requirement to use the COMB3 potential.

***Pair style COMB3 requires atom attribute q***

Self-explanatory.

***Pair style COMB3 requires newton pair on***

See the newton command. This is a restriction to use the COMB3 potential.

***Pair style LCBOP requires atom IDs***

This is a requirement to use the LCBOP potential.

***Pair style LCBOP requires newton pair on***

See the newton command. This is a restriction to use the Tersoff potential.

***Pair style MEAM requires newton pair on***

See the newton command. This is a restriction to use the MEAM potential.

***Pair style SNAP requires newton pair on***

See the newton command. This is a restriction to use the SNAP potential.

***Pair style Stillinger-Weber requires atom IDs***

This is a requirement to use the SW potential.

***Pair style Stillinger-Weber requires newton pair on***

See the newton command. This is a restriction to use the SW potential.

***Pair style Tersoff requires atom IDs***

This is a requirement to use the Tersoff potential.

***Pair style Tersoff requires newton pair on***

See the newton command. This is a restriction to use the Tersoff potential.

***Pair style Vashishta requires atom IDs***

This is a requirement to use the Vashishta potential.

***Pair style Vashishta requires newton pair on***

See the newton command. This is a restriction to use the Vashishta potential.

***Pair style bop requires comm ghost cutoff at least 3x larger than %g***

Use the communicate ghost command to set this. See the pair bop page for more details.

***Pair style born/coul/long requires atom attribute q***

An atom style that defines this attribute must be used.

***Pair style born/coul/long/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style born/coul/wolf requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style buck/coul/cut requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style buck/coul/long requires atom attribute q***

The atom style defined does not have these attributes.

***Pair style buck/coul/long/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style buck/long/coul/long requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style coul/cut requires atom attribute q***

The atom style defined does not have these attributes.

***Pair style coul/cut/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style coul/debye/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style coul/dsf requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style coul/dsf/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style coul/long/gpu requires atom attribute q***

The atom style defined does not have these attributes.

***Pair style coul/streitz requires atom attribute q***

Self-explanatory.

***Pair style does not have extra field requested by compute pair/local***

The pair style does not support the pN value requested by the compute pair/local command.

***Pair style does not support bond\_style quartic***

The pair style does not have a single() function, so it can not be invoked by bond\_style quartic.

***Pair style does not support compute group/group***

The pair\_style does not have a single() function, so it cannot be invoked by the compute group/group command.

***Pair style does not support compute pair/local***

The pair style does not have a single() function, so it can not be invoked by compute pair/local.

***Pair style does not support compute property/local***

The pair style does not have a single() function, so it can not be invoked by fix bond/swap.

***Pair style does not support fix bond/swap***

The pair style does not have a single() function, so it can not be invoked by fix bond/swap.

***Pair style does not support pair\_write***

The pair style does not have a single() function, so it can not be invoked by pair write.

***Pair style does not support rRESPA inner/middle/outer***

You are attempting to use rRESPA options with a pair style that does not support them.

***Pair style granular with history requires atoms have IDs***

Atoms in the simulation do not have IDs, so history effects cannot be tracked by the granular pair potential.

***Pair style hbond/dreiding requires an atom map, see atom\_modify***

Self-explanatory.

***Pair style hbond/dreiding requires atom IDs***

Self-explanatory.

***Pair style hbond/dreiding requires molecular system***

Self-explanatory.

***Pair style hbond/dreiding requires newton pair on***

See the newton command for details.

***Pair style hybrid cannot have hybrid as an argument***

Self-explanatory.

***Pair style hybrid cannot have none as an argument***

Self-explanatory.

***Pair style is incompatible with KSpace style***

If a pair style with a long-range Coulombic component is selected, then a kspace style must also be used.

***Pair style is incompatible with TIP4P KSpace style***

The pair style does not have the requires TIP4P settings.

***Pair style lj/charmm/coul/charmm requires atom attribute q***

The atom style defined does not have these attributes.

***Pair style lj/charmm/coul/long requires atom attribute q***

The atom style defined does not have these attributes.

***Pair style lj/charmm/coul/long/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/class2/coul/cut requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/class2/coul/long requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/class2/coul/long/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/cut/coul/cut requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/cut/coul/cut/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/cut/coul/debye/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/cut/coul/dsf requires atom attribute q***

The atom style defined does not have these attributes.

***Pair style lj/cut/coul/dsf/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/cut/coul/long requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/cut/coul/long/gpu requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/cut/tip4p/cut requires atom IDs***

This is a requirement to use this potential.

***Pair style lj/cut/tip4p/cut requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style lj/cut/tip4p/cut requires newton pair on***

See the newton command. This is a restriction to use this potential.

***Pair style lj/cut/tip4p/long requires atom IDs***

There are no atom IDs defined in the system and the TIP4P potential requires them to find O,H atoms with a water molecule.

***Pair style lj/cut/tip4p/long requires atom attribute q***

The atom style defined does not have these attributes.

***Pair style lj/cut/tip4p/long requires newton pair on***

This is because the computation of constraint forces within a water molecule adds forces to atoms owned by other processors.

***Pair style lj/gromacs/coul/gromacs requires atom attribute q***

An atom\_style with this attribute is needed.

***Pair style lj/long/dipole/long does not currently support respa***

This feature is not yet supported.

***Pair style lj/long/tip4p/long requires atom IDs***

There are no atom IDs defined in the system and the TIP4P potential requires them to find O,H atoms with a water molecule.

***Pair style lj/long/tip4p/long requires atom attribute q***

The atom style defined does not have these attributes.

***Pair style lj/long/tip4p/long requires newton pair on***

This is because the computation of constraint forces within a water molecule adds forces to atoms owned by other processors.

***Pair style lj/spica/coul/long/gpu requires atom attribute q***

The atom style defined does not have this attribute.



***Pair style nb3b/harmonic requires atom IDs***

This is a requirement to use this potential.

***Pair style nb3b/harmonic requires newton pair on***

See the newton command. This is a restriction to use this potential.

***Pair style nm/cut/coul/cut requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style nm/cut/coul/long requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style peri requires atom style peri***

Self-explanatory.

***Pair style polymorphic requires atom IDs***

This is a requirement to use the polymorphic potential.

***Pair style polymorphic requires newton pair on***

See the newton command. This is a restriction to use the polymorphic potential.

***Pair style reax requires atom IDs***

This is a requirement to use the ReaxFF potential.

***Pair style reax requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style reax requires newton pair on***

This is a requirement to use the ReaxFF potential.

***Pair style requires a KSpace style***

No kspace style is defined.

***Pair style requires use of kspace\_style ewald/disp***

Self-explanatory.

***Pair style sw/gpu requires atom IDs***

This is a requirement to use this potential.

***Pair style sw/gpu requires newton pair off***

See the newton command. This is a restriction to use this potential.

***Pair style vashishta/gpu requires atom IDs***

This is a requirement to use this potential.

***Pair style vashishta/gpu requires newton pair off***

See the newton command. This is a restriction to use this potential.

***Pair style tersoff/gpu requires atom IDs***

This is a requirement to use the tersoff/gpu potential.

***Pair style tersoff/gpu requires newton pair off***

See the newton command. This is a restriction to use this pair style.

***Pair style tip4p/cut requires atom IDs***

This is a requirement to use this potential.

***Pair style tip4p/cut requires atom attribute q***

The atom style defined does not have this attribute.

***Pair style tip4p/cut requires newton pair on***

See the newton command. This is a restriction to use this potential.

***Pair style tip4p/long requires atom IDs***

There are no atom IDs defined in the system and the TIP4P potential requires them to find O,H atoms with a water molecule.

***Pair style tip4p/long requires atom attribute q***

The atom style defined does not have these attributes.

***Pair style tip4p/long requires newton pair on***

This is because the computation of constraint forces within a water molecule adds forces to atoms owned by other processors.

***Pair table cutoffs must all be equal to use with KSpace***

When using pair style table with a long-range KSpace solver, the cutoffs for all atom type pairs must all be the same, since the long-range solver starts at that cutoff.

***Pair table parameters did not set N***

List of pair table parameters must include N setting.

***Pair tersoff/zbl requires metal or real units***

This is a current restriction of this pair potential.

***Pair tersoff/zbl/kk requires metal or real units***

This is a current restriction of this pair potential.

***Pair tri/lj requires atom style tri***

Self-explanatory.

***Pair yukawa/colloid requires atom style sphere***

Self-explanatory.

***Pair yukawa/colloid requires atoms with same type have same radius***

Self-explanatory.

***Pair yukawa/colloid/gpu requires atom style sphere***

Self-explanatory.

***PairKIM only works with 3D problems***

This is a current limitation.

***Pair\_coeff command before pair\_style is defined***

Self-explanatory.

***Pair\_coeff command before simulation box is defined***

The pair\_coeff command cannot be used before a read\_data, read\_restart, or create\_box command.

***Pair\_modify command before pair\_style is defined***

Self-explanatory.

***Pair\_modify special setting for pair hybrid incompatible with global special\_bonds setting***

Cannot override a setting of 0.0 or 1.0 or change a setting between 0.0 and 1.0.

***Pair\_write command before pair\_style is defined***

Self-explanatory.

***Particle on or inside fix wall surface***

Particles must be “exterior” to the wall in order for energy/force to be calculated.

***Particle outside surface of region used in fix wall/region***

Particles must be inside the region for energy/force to be calculated. A particle outside the region generates an error.

***Per-atom compute in equal-style variable formula***

Equal-style variables cannot use per-atom quantities.

***Per-atom energy was not tallied on needed timestep***

You are using a thermo keyword that requires potentials to have tallied energy, but they did not on this timestep. See the variable page for ideas on how to make this work.

***Per-atom fix in equal-style variable formula***

Equal-style variables cannot use per-atom quantities.

***Per-atom virial was not tallied on needed timestep***

You are using a thermo keyword that requires potentials to have tallied the virial, but they did not on this timestep. See the variable page for ideas on how to make this work.

***Per-processor system is too big***

The number of owned atoms plus ghost atoms on a single processor must fit in 32-bit integer.

***Potential energy ID for fix neb does not exist***

Self-explanatory.

***Potential energy ID for fix nvt/nph/npt does not exist***

A compute for potential energy must be defined.

***Potential file has duplicate entry***

The potential file has more than one entry for the same element.

***Potential file is missing an entry***

The potential file does not have a needed entry.

***Power by 0 in variable formula***

Self-explanatory.

***Pressure ID for fix box/relax does not exist***

The compute ID needed to compute pressure for the fix does not exist.

***Pressure ID for fix modify does not exist***

Self-explanatory.

***Pressure ID for fix npt/nph does not exist***

Self-explanatory.

***Pressure ID for fix press/berendsen does not exist***

The compute ID needed to compute pressure for the fix does not exist.

***Pressure ID for fix rigid npt/nph does not exist***

Self-explanatory.

***Pressure ID for thermo does not exist***

The compute ID needed to compute pressure for thermodynamics does not exist.

***Pressure control can not be used with fix nvt***

Self-explanatory.

***Pressure control can not be used with fix nvt/asphere***

Self-explanatory.

***Pressure control can not be used with fix nvt/body***

Self-explanatory.

***Pressure control can not be used with fix nvt/sllod***

Self-explanatory.

***Pressure control can not be used with fix nvt/sphere***

Self-explanatory.

***Pressure control must be used with fix nph***

Self-explanatory.

***Pressure control must be used with fix nph/asphere***

Self-explanatory.

***Pressure control must be used with fix nph/body***

Self-explanatory.

***Pressure control must be used with fix nph/small***

Self-explanatory.

***Pressure control must be used with fix nph/sphere***

Self-explanatory.

***Pressure control must be used with fix nphug***

A pressure control keyword (iso, aniso, tri, x, y, or z) must be provided.

***Pressure control must be used with fix npt***

Self-explanatory.

***Pressure control must be used with fix npt/asphere***

Self-explanatory.

***Pressure control must be used with fix npt/body***

Self-explanatory.

***Pressure control must be used with fix npt/sphere***

Self-explanatory.

***Processor count in z must be 1 for 2d simulation***

Self-explanatory.

***Processor partitions do not match number of allocated processors***

The total number of processors in all partitions must match the number of processors LAMMPS is running on.

***Processors command after simulation box is defined***

The processors command cannot be used after a read\_data, read\_restart, or create\_box command.

***Processors custom grid file is inconsistent***

The vales in the custom file are not consistent with the number of processors you are running on or the Px,Py,Pz settings of the processors command. Or there was not a setting for every processor.

***Processors grid numa and map style are incompatible***

Using numa for gstyle in the processors command requires using cart for the map option.

***Processors part option and grid style are incompatible***

Cannot use gstyle numa or custom with the part option.

***Processors twogrid requires proc count be a multiple of core count***

Self-explanatory.

***Pstart and Pstop must have the same value***

Self-explanatory.

***Python function evaluation failed***

The Python function did not run successfully and/or did not return a value (if it is supposed to return a value). This is probably due to some error condition in the function.

***Python function is not callable***

The provided Python code was run successfully, but it not define a callable function with the required name.

***Python invoke of undefined function***

Cannot invoke a function that has not been previously defined.

***Python variable does not match Python function***

This matching is defined by the python-style variable and the python command.

***Python variable has no function***

No python command was used to define the function associated with the python-style variable.

***QEQ with 'newton pair off' not supported***

See the newton command. This is a restriction to use the QEQ fixes.

***R0 < 0 for fix spring command***

Equilibrium spring length is invalid.

***RATTLE coordinate constraints are not satisfied up to desired tolerance***

Self-explanatory.

***RATTLE determinant = 0.0***

The determinant of the matrix being solved for a single cluster specified by the fix rattle command is numerically invalid.

***RATTLE failed***

Certain constraints were not satisfied.

***RATTLE velocity constraints are not satisfied up to desired tolerance***

Self-explanatory.

***Read data add offset is too big***

It cannot be larger than the size of atom IDs, e.g. the maximum 32-bit integer.

***Read dump of atom property that is not allocated***

Self-explanatory.

***Read rerun dump file timestep > specified stop***

Self-explanatory.

***Read restart MPI-IO input not allowed with % in filename***

This is because a % signifies one file per processor and MPI-IO creates one large file for all processors.

***Read\_data shrink wrap did not assign all atoms correctly***

This is typically because the box-size specified in the data file is large compared to the actual extent of atoms in a shrink-wrapped dimension. When LAMMPS shrink-wraps the box atoms will be lost if the processor they are re-assigned to is too far away. Choose a box size closer to the actual extent of the atoms.

***Read\_dump command before simulation box is defined***

The read\_dump command cannot be used before a read\_data, read\_restart, or create\_box command.

***Read\_dump field not found in dump file***

Self-explanatory.

***Read\_dump triclinic status does not match simulation***

Both the dump snapshot and the current LAMMPS simulation must be using either an orthogonal or triclinic box.

***Read\_dump xyz fields do not have consistent scaling/wrapping***

Self-explanatory.

***Reading from MPI-IO filename when MPIIO package is not installed***

Self-explanatory.

***Reax\_defs.h setting for NATDEF is too small***

Edit the setting in the ReaxFF library and re-compile the library and re-build LAMMPS.

***Reax\_defs.h setting for NNEIGHMAXDEF is too small***

Edit the setting in the ReaxFF library and re-compile the library and re-build LAMMPS.

***Receiving partition in processors part command is already a receiver***

Cannot specify a partition to be a receiver twice.

***Region ID for compute chunk/atom does not exist***

Self-explanatory.

***Region ID for compute reduce/region does not exist***

Self-explanatory.

***Region ID for compute temp/region does not exist***

Self-explanatory.

***Region ID for dump custom does not exist***

Self-explanatory.

***Region ID for fix addforce does not exist***

Self-explanatory.

***Region ID for fix atom/swap does not exist***

Self-explanatory.

***Region ID for fix ave/spatial does not exist***

Self-explanatory.

***Region ID for fix aveforce does not exist***

Self-explanatory.

***Region ID for fix deposit does not exist***

Self-explanatory.

***Region ID for fix efield does not exist***

Self-explanatory.

***Region ID for fix evaporate does not exist***

Self-explanatory.

***Region ID for fix gcmc does not exist***

Self-explanatory.

***Region ID for fix heat does not exist***

Self-explanatory.

***Region ID for fix setforce does not exist***

Self-explanatory.

***Region ID for fix wall/region does not exist***

Self-explanatory.

***Region ID for group dynamic does not exist***

Self-explanatory.

***Region ID in variable formula does not exist***

Self-explanatory.

***Region cannot have 0 length rotation vector***

Self-explanatory.

***Region for fix oneway does not exist***

Self-explanatory.

***Region intersect region ID does not exist***

Self-explanatory.

***Region union or intersect cannot be dynamic***

The sub-regions can be dynamic, but not the combined region.

***Region union region ID does not exist***

One or more of the region IDs specified by the region union command does not exist.

***Replacing a fix, but new style != old style***

A fix ID can be used a second time, but only if the style matches the previous fix. In this case it is assumed you wish to reset a fix's parameters. This error may mean you are mistakenly re-using a fix ID when you do not intend to.

***Replicate command before simulation box is defined***

The replicate command cannot be used before a read\_data, read\_restart, or create\_box command.

***Replicate did not assign all atoms correctly***

Atoms replicated by the replicate command were not assigned correctly to processors. This is likely due to some atom coordinates being outside a non-periodic simulation box.

***Replicated system atom IDs are too big***

See the setting for tagint in the src/lmptype.h file.

***Replicated system is too big***

See the setting for bigint in the src/lmptype.h file.

***Required border comm not yet implemented with Kokkos***

There are various limitations in the communication options supported by Kokkos.

***Rerun command before simulation box is defined***

The rerun command cannot be used before a read\_data, read\_restart, or create\_box command.

***Rerun dump file does not contain requested snapshot***

Self-explanatory.

***Resetting timestep size is not allowed with fix move***

This is because fix move is moving atoms based on elapsed time.

***Respa inner cutoffs are invalid***

The first cutoff must be  $\leq$  the second cutoff.

***Respa levels must be  $\geq 1$*** 

Self-explanatory.

***Respa middle cutoffs are invalid***

The first cutoff must be  $\leq$  the second cutoff.

***Restart file MPI-IO output not allowed with % in filename***

This is because a % signifies one file per processor and MPI-IO creates one large file for all processors.

***Restart file byte ordering is not recognized***

The file does not appear to be a LAMMPS restart file since it does not contain a recognized byte-ordering flag at the beginning.

***Restart file byte ordering is swapped***

The file was written on a machine with different byte-ordering than the machine you are reading it on. Convert it to a text data file instead, on the machine you wrote it on.

***Restart file incompatible with current version***

This is probably because you are trying to read a file created with a version of LAMMPS that is too old compared to the current version. Use your older version of LAMMPS and convert the restart file to a data file.

***Restart file is a MPI-IO file***

The file is inconsistent with the filename you specified for it.

***Restart file is a multi-proc file***

The file is inconsistent with the filename you specified for it.

***Restart file is not a MPI-IO file***

The file is inconsistent with the filename you specified for it.

***Restart file is not a multi-proc file***

The file is inconsistent with the filename you specified for it.

***Restart variable returned a bad timestep***

The variable must return a timestep greater than the current timestep.

***Restrain atoms %d %d %d %d missing on proc %d at step %ld***

The 4 atoms in a restrain dihedral specified by the fix restrain command are not all accessible to a processor. This probably means an atom has moved too far.

***Restrain atoms %d %d %d missing on proc %d at step %ld***

The 3 atoms in a restrain angle specified by the fix restrain command are not all accessible to a processor. This probably means an atom has moved too far.

***Restrain atoms %d %d missing on proc %d at step %ld***

The 2 atoms in a restrain bond specified by the fix restrain command are not all accessible to a processor. This probably means an atom has moved too far.

***Reuse of compute ID***

A compute ID cannot be used twice.

***Reuse of dump ID***

A dump ID cannot be used twice.

***Reuse of molecule template ID***

The template IDs must be unique.

***Reuse of region ID***

A region ID cannot be used twice.

***Rigid body atoms %d %d missing on proc %d at step %ld***

This means that an atom cannot find the atom that owns the rigid body it is part of, or vice versa. The solution is to use the communicate cutoff command to insure ghost atoms are acquired from far enough away to encompass the max distance printed when the fix rigid/small command was invoked.

***Rigid body has degenerate moment of inertia***

Fix poems will only work with bodies (collections of atoms) that have non-zero principal moments of inertia. This means they must be 3 or more non-collinear atoms, even with joint atoms removed.

***Rigid fix must come before NPT/NPH fix***

NPT/NPH fix must be defined in input script after all rigid fixes, else the rigid fix contribution to the pressure virial is incorrect.

***Rmask function in equal-style variable formula***

Rmask is per-atom operation.

***Run command before simulation box is defined***

The run command cannot be used before a read\_data, read\_restart, or create\_box command.

***Run command start value is after start of run***

Self-explanatory.

***Run command stop value is before end of run***

Self-explanatory.

***Run\_style command before simulation box is defined***

The run\_style command cannot be used before a read\_data, read\_restart, or create\_box command.



***SRD bin size for fix srd differs from user request***

Fix SRD had to adjust the bin size to fit the simulation box. See the cubic keyword if you want this message to be an error vs warning.

***SRD bins for fix srd are not cubic enough***

The bin shape is not within tolerance of cubic. See the cubic keyword if you want this message to be an error vs warning.

***SRD particle %d started inside big particle %d on step %ld bounce %d***

See the inside keyword if you want this message to be an error vs warning.

***SRD particle %d started inside wall %d on step %ld bounce %d***

See the inside keyword if you want this message to be an error vs warning.

***Same dimension twice in fix ave/spatial***

Self-explanatory.

***Sending partition in processors part command is already a sender***

Cannot specify a partition to be a sender twice.

***Set command before simulation box is defined***

The set command cannot be used before a read\_data, read\_restart, or create\_box command.

***Set command floating point vector does not exist***

Self-explanatory.

***Set command integer vector does not exist***

Self-explanatory.

***Set command with no atoms existing***

No atoms are yet defined so the set command cannot be used.

***Set region ID does not exist***

Region ID specified in set command does not exist.

***Shake angles have different bond types***

All 3-atom angle-constrained SHAKE clusters specified by the fix shake command that are the same angle type, must also have the same bond types for the 2 bonds in the angle.

***Shake atoms %d %d %d %d missing on proc %d at step %ld***

The 4 atoms in a single shake cluster specified by the fix shake command are not all accessible to a processor. This probably means an atom has moved too far.

***Shake atoms %d %d %d missing on proc %d at step %ld***

The 3 atoms in a single shake cluster specified by the fix shake command are not all accessible to a processor. This probably means an atom has moved too far.

***Shake atoms %d %d missing on proc %d at step %ld***

The 2 atoms in a single shake cluster specified by the fix shake command are not all accessible to a processor. This probably means an atom has moved too far.

***Shake cluster of more than 4 atoms***

A single cluster specified by the fix shake command can have no more than 4 atoms.

***Shake clusters are connected***

A single cluster specified by the fix shake command must have a single central atom with up to 3 other atoms bonded to it.

***Shake determinant = 0.0***

The determinant of the matrix being solved for a single cluster specified by the fix shake command is numerically invalid.

***Shake fix must come before NPT/NPH fix***

NPT fix must be defined in input script after SHAKE fix, else the SHAKE fix contribution to the pressure virial is incorrect.

***Shear history overflow, boost neigh\_modify one***

There are too many neighbors of a single atom. Use the neigh\_modify command to increase the max number of neighbors allowed for one atom. You may also want to boost the page size.

***Small to big integers are not sized correctly***

This error occurs when the sizes of smallint, imageint, tagint, bigint, as defined in src/lmptype.h are not what is expected. Contact the developers if this occurs.

***Smallint setting in lmptype.h is invalid***

It has to be the size of an integer.

***Smallint setting in lmptype.h is not compatible***

Smallint stored in restart file is not consistent with LAMMPS version you are running.

***Special list size exceeded in fix bond/create***

See the “read\_data extra/special/per/atom” command (or the “create\_box extra/special/per/atom” command) for info on how to leave space in the special bonds list to allow for additional bonds to be formed.

***Species XXX is not supported by this KIM Simulator Model***

The kim\_style define command was referencing a species that is not present in the requested KIM Simulator Model.

***Specified processors != physical processors***

The 3d grid of processors defined by the processors command does not match the number of processors LAMMPS is being run on.

***Specified target stress must be uniaxial or hydrostatic***

Self-explanatory.

***Sqrt of negative value in variable formula***

Self-explanatory.

***Subsequent read data induced too many angles per atom***

See the extra/angle/per/atom keyword for the create\_box or the read\_data command to set this limit larger

***Subsequent read data induced too many bonds per atom***

See the extra/bond/per/atom keyword for the create\_box or the read\_data command to set this limit larger

***Subsequent read data induced too many dihedrals per atom***

See the extra/dihedral/per/atom keyword for the create\_box or the read\_data command to set this limit larger

***Subsequent read data induced too many improper per atom***

See the extra/improper/per/atom keyword for the create\_box or the read\_data command to set this limit larger

***Substitution for illegal variable***

Input script line contained a variable that could not be substituted for.

***Support for writing images in JPEG format not included***

LAMMPS was not built with the -DLAMMPS\_JPEG switch in the Makefile.

***Support for writing images in PNG format not included***

LAMMPS was not built with the -DLAMMPS\_PNG switch in the Makefile.

***Support for writing movies not included***

LAMMPS was not built with the -DLAMMPS\_FFmpeg switch in the Makefile

***System in data file is too big***

See the setting for bigint in the src/lmptype.h file.

***System is not charge neutral, net charge = %g***

The total charge on all atoms on the system is not 0.0. For some KSpace solvers this is an error.

***TAD nsteps must be multiple of t\_event***

Self-explanatory.

***TIP4P hydrogen has incorrect atom type***

The TIP4P pairwise computation found an H atom whose type does not agree with the specified H type.

***TIP4P hydrogen is missing***

The TIP4P pairwise computation failed to find the correct H atom within a water molecule.

***TMD target file did not list all group atoms***

The target file for the fix tmd command did not list all atoms in the fix group.

***Tad command before simulation box is defined***

Self-explanatory.

***Tagint setting in lmptype.h is invalid***

Tagint must be as large or larger than smallint.

***Tagint setting in lmptype.h is not compatible***

Format of tagint stored in restart file is not consistent with LAMMPS version you are running. See the settings in src/lmptype.h

***Target pressure for fix rigid/nph cannot be < 0.0***

Self-explanatory.

***Target pressure for fix rigid/npt/small cannot be < 0.0***

Self-explanatory.

***Target temperature for fix nvt/npt/nph cannot be 0.0***

Self-explanatory.

***Target temperature for fix rigid/npt cannot be 0.0***

Self-explanatory.

***Target temperature for fix rigid/npt/small cannot be 0.0***

Self-explanatory.

***Target temperature for fix rigid/nvt cannot be 0.0***

Self-explanatory.

***Target temperature for fix rigid/nvt/small cannot be 0.0***

Self-explanatory.

***Temper command before simulation box is defined***

The temper command cannot be used before a read\_data, read\_restart, or create\_box command.

***Temperature ID for fix bond/swap does not exist***

Self-explanatory.

***Temperature ID for fix box/relax does not exist***

Self-explanatory.

***Temperature ID for fix nvt/npt does not exist***

Self-explanatory.

***Temperature ID for fix press/berendsen does not exist***

Self-explanatory.

***Temperature ID for fix rigid nvt/npt/nph does not exist***

Self-explanatory.

***Temperature ID for fix temp/berendsen does not exist***

Self-explanatory.

***Temperature ID for fix temp/csld does not exist***

Self-explanatory.

***Temperature ID for fix temp/csvr does not exist***

Self-explanatory.

***Temperature ID for fix temp/rescale does not exist***

Self-explanatory.

***Temperature compute degrees of freedom < 0***

This should not happen if you are calculating the temperature on a valid set of atoms.

***Temperature control can not be used with fix nph***

Self-explanatory.

***Temperature control can not be used with fix nph/asphere***

Self-explanatory.

***Temperature control can not be used with fix nph/body***

Self-explanatory.

***Temperature control can not be used with fix nph/sphere***

Self-explanatory.

***Temperature control must be used with fix nphug***

The temp keyword must be provided.

***Temperature control must be used with fix npt***

Self-explanatory.

***Temperature control must be used with fix npt/asphere***

Self-explanatory.

***Temperature control must be used with fix npt/body***

Self-explanatory.

***Temperature control must be used with fix npt/sphere***

Self-explanatory.

***Temperature control must be used with fix nvt***

Self-explanatory.

***Temperature control must be used with fix nvt/asphere***

Self-explanatory.

***Temperature control must be used with fix nvt/body***

Self-explanatory.

***Temperature control must be used with fix nvt/sllod***

Self-explanatory.

***Temperature control must be used with fix nvt/sphere***

Self-explanatory.

***Temperature control must not be used with fix nph/small***

Self-explanatory.

***Temperature for fix nvt/sllod does not have a bias***

The specified compute must compute temperature with a bias.

***Tempering could not find thermo\_pe compute***

This compute is created by the thermo command. It must have been explicitly deleted by a uncompute command.

***Tempering fix ID is not defined***

The fix ID specified by the temper command does not exist.

***Tempering temperature fix is not valid***

The fix specified by the temper command is not one that controls temperature (nvt or langevin).

***Test\_descriptor\_string already allocated***

This is an internal error. Contact the developers.

***The package gpu command is required for gpu styles***

Self-explanatory.

***Thermo and fix not computed at compatible times***

Fixes generate values on specific timesteps. The thermo output does not match these timesteps.

***Thermo compute array is accessed out-of-range***

Self-explanatory.

***Thermo compute does not compute array***

Self-explanatory.

***Thermo compute does not compute scalar***

Self-explanatory.

***Thermo compute does not compute vector***

Self-explanatory.

***Thermo compute vector is accessed out-of-range***

Self-explanatory.

***Thermo custom variable cannot be indexed***

Self-explanatory.

***Thermo custom variable is not equal-style variable***

Only equal-style variables can be output with thermodynamics, not atom-style variables.

***Thermo every variable returned a bad timestep***

The variable must return a timestep greater than the current timestep.

***Thermo fix array is accessed out-of-range***

Self-explanatory.

***Thermo fix does not compute array***

Self-explanatory.

***Thermo fix does not compute scalar***

Self-explanatory.

***Thermo fix does not compute vector***

Self-explanatory.

***Thermo fix vector is accessed out-of-range***

Self-explanatory.

***Thermo keyword in variable requires thermo to use/init pe***

You are using a thermo keyword in a variable that requires potential energy to be calculated, but your thermo output does not use it. Add it to your thermo output.

***Thermo keyword in variable requires thermo to use/init press***

You are using a thermo keyword in a variable that requires pressure to be calculated, but your thermo output does not use it. Add it to your thermo output.

***Thermo keyword in variable requires thermo to use/init temp***

You are using a thermo keyword in a variable that requires temperature to be calculated, but your thermo output does not use it. Add it to your thermo output.

***Thermo style does not use press***

Cannot use thermo\_modify to set this parameter since the thermo\_style is not computing this quantity.

***Thermo style does not use temp***

Cannot use thermo\_modify to set this parameter since the thermo\_style is not computing this quantity.

***Thermo\_modify every variable returned a bad timestep***

The returned timestep is less than or equal to the current timestep.

***Thermo\_modify int format does not contain d character***

Self-explanatory.

***Thermo\_modify pressure ID does not compute pressure***

The specified compute ID does not compute pressure.

***Thermo\_modify temperature ID does not compute temperature***

The specified compute ID does not compute temperature.

***Thermo\_style command before simulation box is defined***

The thermo\_style command cannot be used before a read\_data, read\_restart, or create\_box command.

***This variable thermo keyword cannot be used between runs***

Keywords that refer to time (such as cpu, elapsed) do not make sense in between runs.

***Threshold for an atom property that is not allocated***

A dump threshold has been requested on a quantity that is not defined by the atom style used in this simulation.

***Timestep must be >= 0***

Specified timestep is invalid.

***Too big a problem to use velocity create loop all***

The system size must fit in a 32-bit integer to use this option.

***Too big a timestep for dump dcd***

The timestep must fit in a 32-bit integer to use this dump style.

***Too big a timestep for dump xtc***

The timestep must fit in a 32-bit integer to use this dump style.

***Too few bits for lookup table***

Table size specified via pair\_modify command does not work with your machine's floating point representation.

***Too few lines in %s section of data file***

Self-explanatory.

***Too few values in body lines in data file***

Self-explanatory.

***Too few values in body section of molecule file***

Self-explanatory.

***Too many -pk arguments in command line***

The string formed by concatenating the arguments is too long. Use a package command in the input script instead.

***Too many MSM grid levels***

The max number of MSM grid levels is hardwired to 10.

***Too many args in variable function***

More args are used than any variable function allows.

***Too many atom pairs for pair bop***

The number of atomic pairs exceeds the expected number. Check your atomic structure to ensure that it is realistic.

***Too many atom sorting bins***

This is likely due to an immense simulation box that has blown up to a large size.

***Too many atom triplets for pair bop***

The number of three atom groups for angle determinations exceeds the expected number. Check your atomic structure to ensure that it is realistic.

***Too many atoms for dump dcd***

The system size must fit in a 32-bit integer to use this dump style.

***Too many atoms for dump xtc***

The system size must fit in a 32-bit integer to use this dump style.

***Too many elements extracted from MEAM library.***

Increase 'maxelt' in meam.h and recompile.

***Too many exponent bits for lookup table***

Table size specified via pair\_modify command does not work with your machine's floating point representation.

***Too many groups***

The maximum number of atom groups (including the "all" group) is given by MAX\_GROUP in group.cpp and is 32.

***Too many iterations***

You must use a number of iterations that fit in a 32-bit integer for minimization.

***Too many lines in one body in data file - boost MAXBODY***

MAXBODY is a setting at the top of the src/read\_data.cpp file. Set it larger and re-compile the code.

***Too many local+ghost atoms for neighbor list***

The number of nlocal + nghost atoms on a processor is limited by the size of a 32-bit integer with 2 bits removed for masking 1-2, 1-3, 1-4 neighbors.

***Too many mantissa bits for lookup table***

Table size specified via pair\_modify command does not work with your machine's floating point representation.

***Too many masses for fix shake***

The fix shake command cannot list more masses than there are atom types.

***Too many molecules for fix poems***

The limit is  $2^{31} = \sim 2$  billion molecules.

***Too many molecules for fix rigid***

The limit is  $2^{31} = \sim 2$  billion molecules.

***Too many neighbor bins***

This is likely due to an immense simulation box that has blown up to a large size.

***Too many timesteps***

The cumulative timesteps must fit in a 64-bit integer.

***Too many timesteps for NEB***

You must use a number of timesteps that fit in a 32-bit integer for NEB.

***Too many total atoms***

See the setting for bigint in the src/lmptype.h file.

***Too many total bits for bitmapped lookup table***

Table size specified via pair\_modify command is too large. Note that a value of N generates a  $2^N$  size table.

***Too many values in body lines in data file***

Self-explanatory.

***Too many values in body section of molecule file***

Self-explanatory.

***Too much buffered per-proc info for dump***

The size of the buffered string must fit in a 32-bit integer for a dump.

***Too much per-proc info for dump***

Number of local atoms times number of columns must fit in a 32-bit integer for dump.

***Topology type exceeds system topology type***

The number of bond, angle, etc types exceeds the system setting. See the create\_box or read\_data command for how to specify these values.

***Tree structure in joint connections***

Fix poems cannot (yet) work with coupled bodies whose joints connect the bodies in a tree structure.

***Triclinic box skew is too large***

The displacement in a skewed direction must be less than half the box length in that dimension. E.g. the xy tilt must be between -half and +half of the x box length. This constraint can be relaxed by using the box tilt command.

***Tried to convert a double to int, but input\_double > INT\_MAX***

Self-explanatory.

***Trying to build an occasional neighbor list before initialization completed***

This is not allowed. Source code caller needs to be modified.

***Two fix ave commands using same compute chunk/atom command in incompatible ways***

They are both attempting to “lock” the chunk/atom command so that the chunk assignments persist for some number of timesteps, but are doing it in different ways.

***Two groups cannot be the same in fix spring couple***

Self-explanatory.

***The %s type label %s is already in use for type %s***

For a given type-kind (atom types, bond types, etc.), a given type label can be assigned to only one numeric type.

***Type label string %s for %s type %s is invalid***

See the labelmap command documentation for valid type labels.

***Unable to initialize accelerator for use***

There was a problem initializing an accelerator for the gpu package

***Unbalanced quotes in input line***

No matching end double quote was found following a leading double quote.

***Unexpected end of -reorder file***

Self-explanatory.

***Unexpected empty line in Angle Coeffs section***

Read a blank line where there should be coefficient data.

***Unexpected empty line in Bond Coeffs section***

Read a blank line where there should be coefficient data.

***Unexpected empty line in Dihedral Coeffs section***

Read a blank line where there should be coefficient data.

***Unexpected empty line in Improper Coeffs section***

Read a blank line where there should be coefficient data.



***Unexpected empty line in Pair Coeffs section***

Read a blank line where there should be coefficient data.

***Unexpected end of custom file***

Self-explanatory.

***Unexpected end of data file***

LAMMPS hit the end of the data file while attempting to read a section. Something is wrong with the format of the data file.

***Unexpected end of dump file***

A read operation from the file failed.

***Unexpected end of fix rigid file***

A read operation from the file failed.

***Unexpected end of fix rigid/small file***

A read operation from the file failed.

***Unexpected end of molecule file***

Self-explanatory.

***Unexpected end of neb file***

A read operation from the file failed.

***Units command after simulation box is defined***

The units command cannot be used after a read\_data, read\_restart, or create\_box command.

***Universe/uloop variable count < # of partitions***

A universe or uloop style variable must specify a number of values  $\geq$  to the number of processor partitions.

***Unrecognized angle style***

The choice of angle style is unknown.

***Unrecognized atom style***

The choice of atom style is unknown.

***Unrecognized body style***

The choice of body style is unknown.

***Unrecognized bond style***

The choice of bond style is unknown.

***Unknown category for info is\_active()***

Self-explanatory.

***Unknown category for info is\_available()***

Self-explanatory.

***Unknown category for info is\_defined()***

Self-explanatory.

***Unrecognized command: %s***

The command is not known to LAMMPS. Check the input script.

***Unrecognized compute style***

The choice of compute style is unknown.

***Unrecognized dihedral style***

The choice of dihedral style is unknown.

***Unrecognized dump reader style***

The choice of dump reader style via the format keyword is unknown.

***Unrecognized dump style***

The choice of dump style is unknown.

***Unknown error in GPU library***

Self-explanatory.

***Unrecognized fix style***

The choice of fix style is unknown.

***Unknown identifier in data file: %s***

A section of the data file cannot be read by LAMMPS.

***Unrecognized improper style***

The choice of improper style is unknown.

***Unknown keyword in thermo\_style custom command***

One or more specified keywords are not recognized.

***Unrecognized kspace style***

The choice of kspace style is unknown.

***Unknown name for info newton category***

Self-explanatory.

***Unknown name for info package category***

Self-explanatory.

***Unknown name for info pair category***

Self-explanatory.

***Unrecognized pair style***

The choice of pair style is unknown.

***Unknown pair\_modify hybrid sub-style***

The choice of sub-style is unknown.

***Unrecognized region style***

The choice of region style is unknown.

***Unknown section in molecule file***

Self-explanatory.

***Unknown table style in angle style table***

Self-explanatory.

***Unknown table style in bond style table***

Self-explanatory.

***Unknown table style in pair\_style command***

Style of table is invalid for use with pair\_style table command.

***Unknown unit\_style***

Self-explanatory. Check the input script or data file.

***Unrecognized lattice type in MEAM library file***

The lattice type in an entry of the MEAM library file is not valid.

***Unrecognized lattice type in MEAM parameter file***

The lattice type in an entry of the MEAM parameter file is not valid.

***Unrecognized pair style in compute pair command***

Self-explanatory.

***Unsupported mixing rule in kspace\_style ewald/disp***

Only geometric mixing is supported.

***Unsupported order in kspace\_style ewald/disp***

Only  $1/r^6$  dispersion or dipole terms are supported.

***Unsupported order in kspace\_style ppm/disp, pair\_style %s***

Only pair styles with  $1/r$  and  $1/r^6$  dependence are currently supported.

***Unsupported parameter in MEAM library file***

Self-explanatory.

***Use cutoff keyword to set cutoff in single mode***

Mode is single so cutoff/multi keyword cannot be used.

***Use cutoff/multi keyword to set cutoff in multi mode***

Mode is multi so cutoff keyword cannot be used.

***Using fix nvt/sllod with inconsistent fix deform remap option***

Fix nvt/sllod requires that deforming atoms have a velocity profile provided by “remap v” as a fix deform option.

***Using fix nvt/sllod with no fix deform defined***

Self-explanatory.

***Using fix srd with inconsistent fix deform remap option***

When shearing the box in an SRD simulation, the remap v option for fix deform needs to be used.

***Using pair lubricate with inconsistent fix deform remap option***

Must use remap v option with fix deform with this pair style.

***Using pair lubricate/poly with inconsistent fix deform remap option***

If fix deform is used, the remap v option is required.

***Using suffix gpu without GPU package installed***

Self-explanatory.

***Using suffix intel without INTEL package installed***

Self-explanatory.

***Using suffix kk without KOKKOS package enabled***

Self-explanatory.

***Using suffix omp without OPENMP package installed***

Self-explanatory.

***Using update dipole flag requires atom attribute mu***

Self-explanatory.

***Using update dipole flag requires atom style sphere***

Self-explanatory.

***Variable ID in variable formula does not exist***

Self-explanatory.

***Variable atom ID is too large***

Specified ID is larger than the maximum allowed atom ID.

***Variable evaluation before simulation box is defined***

Cannot evaluate a compute or fix or atom-based value in a variable before the simulation has been setup.

***Variable evaluation in fix wall gave bad value***

The returned value for epsilon or sigma < 0.0.

***Variable evaluation in region gave bad value***

Variable returned a radius < 0.0.

***Variable for compute ti is invalid style***

Self-explanatory.

***Variable for create\_atoms is invalid style***

The variables must be equal-style variables.

***Variable for displace\_atoms is invalid style***

It must be an equal-style or atom-style variable.

***Variable for dump every is invalid style***

Only equal-style variables can be used.

***Variable for dump image center is invalid style***

Must be an equal-style variable.

***Variable for dump image phi is invalid style***

Must be an equal-style variable.

***Variable for dump image theta is invalid style***

Must be an equal-style variable.

***Variable for dump image zoom is invalid style***

Must be an equal-style variable.

***Variable for fix adapt is invalid style***

Only equal-style variables can be used.

***Variable for fix addforce is invalid style***

Self-explanatory.

***Variable for fix aveforce is invalid style***

Only equal-style variables can be used.

***Variable for fix deform is invalid style***

The variable must be an equal-style variable.

***Variable for fix efield is invalid style***

The variable must be an equal- or atom-style variable.

***Variable for fix gravity is invalid style***

Only equal-style variables can be used.

***Variable for fix heat is invalid style***

Only equal-style or atom-style variables can be used.

***Variable for fix indent is invalid style***

Only equal-style variables can be used.

***Variable for fix indent is not equal style***

Only equal-style variables can be used.

***Variable for fix langevin is invalid style***

It must be an equal-style variable.

***Variable for fix move is invalid style***

Only equal-style variables can be used.

***Variable for fix setforce is invalid style***

Only equal-style variables can be used.

***Variable for fix temp/berendsen is invalid style***

Only equal-style variables can be used.

***Variable for fix temp/csld is invalid style***

Only equal-style variables can be used.

***Variable for fix temp/csvr is invalid style***

Only equal-style variables can be used.

***Variable for fix temp/rescale is invalid style***

Only equal-style variables can be used.

***Variable for fix wall is invalid style***

Only equal-style variables can be used.

***Variable for fix wall/reflect is invalid style***

Only equal-style variables can be used.

***Variable for fix wall/srd is invalid style***

Only equal-style variables can be used.

***Variable for group dynamic is invalid style***

The variable must be an atom-style variable.

***Variable for group is invalid style***

Only atom-style variables can be used.

***Variable for region cylinder is invalid style***

Only equal-style variables are allowed.

***Variable for region is invalid style***

Only equal-style variables can be used.

***Variable for region is not equal style***

Self-explanatory.

***Variable for region sphere is invalid style***

Only equal-style variables are allowed.

***Variable for restart is invalid style***

Only equal-style variables can be used.

***Variable for set command is invalid style***

Only atom-style variables can be used.

***Variable for thermo every is invalid style***

Only equal-style variables can be used.

***Variable for velocity set is invalid style***

Only atom-style variables can be used.

***Variable for voronoi radius is not atom style***

Self-explanatory.

***Variable formula compute array is accessed out-of-range***

Self-explanatory.

***Variable formula compute vector is accessed out-of-range***

Self-explanatory.

***Variable formula fix array is accessed out-of-range***

Self-explanatory.

***Variable formula fix vector is accessed out-of-range***

Self-explanatory.

***Variable has circular dependency***

A circular dependency is when variable “a” is used by variable “b” and variable “b” is also used by variable “a”. Circular dependencies with longer chains of dependence are also not allowed.

***Variable name between brackets must be alphanumeric or underscore characters***

Self-explanatory.

***Variable name for compute chunk/atom does not exist***

Self-explanatory.

***Variable name for compute reduce does not exist***

Self-explanatory.

***Variable name for compute ti does not exist***

Self-explanatory.

***Variable name for create\_atoms does not exist***

Self-explanatory.

***Variable name for displace\_atoms does not exist***

Self-explanatory.

***Variable name for dump every does not exist***

Self-explanatory.

***Variable name for dump image center does not exist***

Self-explanatory.

***Variable name for dump image phi does not exist***

Self-explanatory.

***Variable name for dump image theta does not exist***

Self-explanatory.

***Variable name for dump image zoom does not exist***

Self-explanatory.

***Variable name for fix adapt does not exist***

Self-explanatory.

***Variable name for fix addforce does not exist***

Self-explanatory.

***Variable name for fix ave/atom does not exist***

Self-explanatory.

***Variable name for fix ave/chunk does not exist***

Self-explanatory.

***Variable name for fix ave/correlate does not exist***

Self-explanatory.

***Variable name for fix ave/histo does not exist***

Self-explanatory.

***Variable name for fix ave/spatial does not exist***

Self-explanatory.

***Variable name for fix ave/time does not exist***

Self-explanatory.

*Variable name for fix aveforce does not exist*

Self-explanatory.

*Variable name for fix deform does not exist*

Self-explanatory.

*Variable name for fix efield does not exist*

Self-explanatory.

*Variable name for fix gravity does not exist*

Self-explanatory.

*Variable name for fix heat does not exist*

Self-explanatory.

*Variable name for fix indent does not exist*

Self-explanatory.

*Variable name for fix langevin does not exist*

Self-explanatory.

*Variable name for fix move does not exist*

Self-explanatory.

*Variable name for fix setforce does not exist*

Self-explanatory.

*Variable name for fix store/state does not exist*

Self-explanatory.

*Variable name for fix temp/berendsen does not exist*

Self-explanatory.

*Variable name for fix temp/csld does not exist*

Self-explanatory.

*Variable name for fix temp/csvr does not exist*

Self-explanatory.

*Variable name for fix temp/rescale does not exist*

Self-explanatory.

*Variable name for fix vector does not exist*

Self-explanatory.

*Variable name for fix wall does not exist*

Self-explanatory.

*Variable name for fix wall/reflect does not exist*

Self-explanatory.

*Variable name for fix wall/srd does not exist*

Self-explanatory.

*Variable name for group does not exist*

Self-explanatory.

*Variable name for group dynamic does not exist*

Self-explanatory.

*Variable name for region cylinder does not exist*

Self-explanatory.

***Variable name for region does not exist***

Self-explanatory.

***Variable name for region sphere does not exist***

Self-explanatory.

***Variable name for restart does not exist***

Self-explanatory.

***Variable name for set command does not exist***

Self-explanatory.

***Variable name for thermo every does not exist***

Self-explanatory.

***Variable name for velocity set does not exist***

Self-explanatory.

***Variable name for voronoi radius does not exist***

Self-explanatory.

***Variable name must be alphanumeric or underscore characters***

Self-explanatory.

***Variable uses atom property that is not allocated***

Self-explanatory.

***Velocity command before simulation box is defined***

The velocity command cannot be used before a read\_data, read\_restart, or create\_box command.

***Velocity command with no atoms existing***

A velocity command has been used, but no atoms yet exist.

***Velocity ramp in z for a 2d problem***

Self-explanatory.

***Velocity rigid used with non-rigid fix-ID***

Self-explanatory.

***Velocity temperature ID does calculate a velocity bias***

The specified compute must compute a bias for temperature.

***Velocity temperature ID does not compute temperature***

The compute ID given to the velocity command must compute temperature.

***Verlet/split can only currently be used with comm\_style brick***

This is a current restriction in LAMMPS.

***Verlet/split does not yet support TIP4P***

This is a current limitation.

***Verlet/split requires 2 partitions***

See the -partition command-line switch.

***Verlet/split requires Rspace partition layout be multiple of Kspace partition layout in each dim***

This is controlled by the processors command.

***Verlet/split requires Rspace partition size be multiple of Kspace partition size***

This is so there is an equal number of Rspace processors for every Kspace processor.

***Virial was not tallied on needed timestep***

You are using a thermo keyword that requires potentials to have tallied the virial, but they did not on this timestep. See the variable page for ideas on how to make this work.



**Voro++ error: narea and neigh have a different size**

This error is returned by the Voro++ library.

**Wall defined twice in fix wall command**

Self-explanatory.

**Wall defined twice in fix wall/reflect command**

Self-explanatory.

**Wall defined twice in fix wall/srd command**

Self-explanatory.

**Water H epsilon must be 0.0 for pair style lj/cut/tip4p/cut**

This is because LAMMPS does not compute the Lennard-Jones interactions with these particles for efficiency reasons.

**Water H epsilon must be 0.0 for pair style lj/cut/tip4p/long**

This is because LAMMPS does not compute the Lennard-Jones interactions with these particles for efficiency reasons.

**Water H epsilon must be 0.0 for pair style lj/long/tip4p/long**

This is because LAMMPS does not compute the Lennard-Jones interactions with these particles for efficiency reasons.

**World variable count does not match # of partitions**

A world-style variable must specify a number of values equal to the number of processor partitions.

**Write \_data command before simulation box is defined**

Self-explanatory.

**Write \_restart command before simulation box is defined**

The write\_restart command cannot be used before a read\_data, read\_restart, or create\_box command.

**Writing to MPI-IO filename when MPIIO package is not installed**

Self-explanatory.

**Zero length rotation vector with displace\_atoms**

Self-explanatory.

**Zero length rotation vector with fix move**

Self-explanatory.

**Zero-length lattice orient vector**

Self-explanatory.

## 11.6 Warning messages

This is an alphabetic list of the WARNING messages LAMMPS prints out and the reason why. If the explanation here is not sufficient, the documentation for the offending command may help. Warning messages also list the source file and line number where the warning was generated. For example, a message like this:

```
WARNING: Bond atom missing in box size check (domain.cpp:187)
```

means that line #187 in the file src/domain.cpp generated the error. Looking in the source code may help you figure out what went wrong.

Doc page with [ERROR messages](#)

***Adjusting Coulombic cutoff for MSM, new cutoff = %cg***

The adjust/cutoff command is turned on and the Coulombic cutoff has been adjusted to match the user-specified accuracy.

***Angle atoms missing at step %ld***

One or more of 3 atoms needed to compute a particular angle are missing on this processor. Typically this is because the pairwise cutoff is set too short or the angle has blown apart and an atom is too far away.

***Angle style in data file differs from currently defined angle style***

Self-explanatory.

***Angles are defined but no angle style is set***

The topology contains angles, but there are no angle forces computed since there was no angle\_style command.

***Atom style in data file differs from currently defined atom style***

Self-explanatory.

***Bond atom missing in box size check***

The second atom needed to compute a particular bond is missing on this processor. Typically this is because the pairwise cutoff is set too short or the bond has blown apart and an atom is too far away.

***Bond atom missing in image check***

The second atom in a particular bond is missing on this processor. Typically this is because the pairwise cutoff is set too short or the bond has blown apart and an atom is too far away.

***Bond atoms missing at step %ld***

The second atom needed to compute a particular bond is missing on this processor. Typically this is because the pairwise cutoff is set too short or the bond has blown apart and an atom is too far away.

***Bond style in data file differs from currently defined bond style***

Self-explanatory.

***Bonds are defined but no bond style is set***

The topology contains bonds, but there are no bond forces computed since there was no bond\_style command.

***Bond/angle/dihedral extent > half of periodic box length***

This is a restriction because LAMMPS can be confused about which image of an atom in the bonded interaction is the correct one to use. “Extent” in this context means the maximum end-to-end length of the bond/angle/dihedral. LAMMPS computes this by taking the maximum bond length, multiplying by the number of bonds in the interaction (e.g. 3 for a dihedral) and adding a small amount of stretch.

***Both groups in compute group/group have a net charge; the Kspace boundary correction to energy will be non-zero***

Self-explanatory.

***Calling write\_dump before a full system init.***

The write\_dump command is used before the system has been fully initialized as part of a ‘run’ or ‘minimize’ command. Not all dump styles and features are fully supported at this point and thus the command may fail or produce incomplete or incorrect output. Insert a “run 0” command, if a full system init is required.

***Cannot count rigid body degrees-of-freedom before bodies are fully initialized***

This means the temperature associated with the rigid bodies may be incorrect on this timestep.

***Cannot count rigid body degrees-of-freedom before bodies are initialized***

This means the temperature associated with the rigid bodies may be incorrect on this timestep.

***Cannot include log terms without 1/r terms; setting flagHI to 1***

Self-explanatory.

***Cannot include log terms without 1/r terms; setting flagHI to 1.***

Self-explanatory.

**Charges are set, but coulombic solver is not used**

Self-explanatory.

**Charges did not converge at step %ld: %lg**

Self-explanatory.

**Communication cutoff is 0.0. No ghost atoms will be generated. Atoms may get lost**

The communication cutoff defaults to the maximum of what is inferred from pair and bond styles (will be zero, if none are defined) and what is specified via `comm_modify cutoff` (defaults to 0.0). If this results to 0.0, no ghost atoms will be generated and LAMMPS may lose atoms or use incorrect periodic images of atoms in interaction lists. To avoid, either use `pair style zero` with a suitable cutoff or use `comm_modify cutoff`.

**Communication cutoff is shorter than a bond length based estimate. This may lead to errors.**

Since LAMMPS stores topology data with individual atoms, all atoms comprising a bond, angle, dihedral or improper must be present on any sub-domain that “owns” the atom with the information, either as a local or a ghost atom. The communication cutoff is what determines up to what distance from a sub-domain boundary ghost atoms are created. The communication cutoff is by default the largest non-bonded cutoff plus the neighbor skin distance, but for short or non-bonded cutoffs and/or long bonds, this may not be sufficient. This warning indicates that there is an increased risk of a simulation stopping unexpectedly because of Bond/Angle/Dihedral/Improper atoms missing. It can be silenced by manually setting the communication cutoff via `comm_modify cutoff`. However, since the heuristic used to determine the estimate is not always accurate, it is not changed automatically and the warning may be ignored depending on the specific system being simulated.

**Communication cutoff is too small for SNAP micro load balancing, increased to %lf**

Self-explanatory.

**Compute cna/atom cutoff may be too large to find ghost atom neighbors**

The neighbor cutoff used may not encompass enough ghost atoms to perform this operation correctly.

**Computing temperature of portions of rigid bodies**

The group defined by the temperature compute does not encompass all the atoms in one or more rigid bodies, so the change in degrees-of-freedom for the atoms in those partial rigid bodies will not be accounted for.

**Create\_bonds max distance > minimum neighbor cutoff**

This means atom pairs for some atom types may not be in the neighbor list and thus no bond can be created between them.

**Delete\_atoms cutoff > minimum neighbor cutoff**

This means atom pairs for some atom types may not be in the neighbor list and thus an atom in that pair cannot be deleted.

**Dihedral atoms missing at step %ld**

One or more of 4 atoms needed to compute a particular dihedral are missing on this processor. Typically this is because the pairwise cutoff is set too short or the dihedral has blown apart and an atom is too far away.

**Dihedral problem**

Conformation of the 4 listed dihedral atoms is extreme; you may want to check your simulation geometry.

**Dihedral problem: %d %ld %d %d %d %d**

Conformation of the 4 listed dihedral atoms is extreme; you may want to check your simulation geometry.

**Dihedral style in data file differs from currently defined dihedral style**

Self-explanatory.

**Dihedrals are defined but no dihedral style is set**

The topology contains dihedrals, but there are no dihedral forces computed since there was no `dihedral_style` command.

**Dump dcd/xtc timestamp may be wrong with fix dt/reset**

If the fix changes the timestep, the dump dcd file will not reflect the change.

***Energy due to X extra global DOFs will be included in minimizer energies***

When using fixes like box/relax, the potential energy used by the minimizer is augmented by an additional energy provided by the fix. Thus the printed converged energy may be different from the total potential energy.

***Estimated error in splitting of dispersion coeffs is %g***

Error is greater than 0.0001 percent.

***Ewald/disp Newton solver failed, using old method to estimate g\_ewald***

Self-explanatory. Choosing a different cutoff value may help.

***FENE bond too long***

A FENE bond has stretched dangerously far. It's interaction strength will be truncated to attempt to prevent the bond from blowing up.

***FENE bond too long: %ld %d %d %g***

A FENE bond has stretched dangerously far. It's interaction strength will be truncated to attempt to prevent the bond from blowing up.

***FENE bond too long: %ld %g***

A FENE bond has stretched dangerously far. It's interaction strength will be truncated to attempt to prevent the bond from blowing up.

***Fix halt condition for fix-id %s met on step %ld with value %g***

Self explanatory.

***Fix SRD walls overlap but fix srd overlap not set***

You likely want to set this in your input script.

***Fix bond/create is used multiple times or with fix bond/break - may not work as expected***

When using fix bond/create multiple times or in combination with fix bond/break, the individual fix instances do not share information about changes they made at the same time step and thus it may result in unexpected behavior.

***Fix bond/react: Atom affected by reaction too close to template edge***

This means an atom which changes type or connectivity during the reaction is too close to an 'edge' atom defined in the superimpose file. This could cause incorrect assignment of bonds, angle, etc. Generally, this means you must include more atoms in your templates, such that there are at least two atoms between each atom involved in the reaction and an edge atom.

***Fix bond/swap will ignore defined angles***

See the page for fix bond/swap for more info on this restriction.

***Fix deposit near setting < possible overlap separation %g***

This test is performed for finite size particles with a diameter, not for point particles. The near setting is smaller than the particle diameter which can lead to overlaps.

***Fix evaporate may delete atom with non-zero molecule ID***

This is probably an error, since you should not delete only one atom of a molecule.

***Fix gcmc using full\_energy option***

Fix gcmc has automatically turned on the full\_energy option since it is required for systems like the one specified by the user. User input included one or more of the following: kspace, triclinic, a hybrid pair style, an eam pair style, or no "single" function for the pair style.

***Fix langevin gif using random gaussians is not implemented with kokkos*** This will most likely cause errors in kinetic fluctuations.

***Fix property/atom mol or charge w/out ghost communication***

A model typically needs these properties defined for ghost atoms.

***Fix qeq CG convergence failed (%g) after %d iterations at %ld step***

Self-explanatory.

***Fix qeq has non-zero lower Taper radius cutoff***

Absolute value must be  $\leq 0.01$ .

***Fix qeq has very low Taper radius cutoff***

Value should typically be  $\geq 5.0$ .

***Fix qeq/dynamic tolerance may be too small for damped dynamics***

Self-explanatory.

***Fix qeq/fire tolerance may be too small for damped fires***

Self-explanatory.

***Fix rattle should come after all other integration fixes***

This fix is designed to work after all other integration fixes change atom positions. Thus it should be the last integration fix specified. If not, it will not satisfy the desired constraints as well as it otherwise would.

***Fix recenter should come after all other integration fixes***

Other fixes may change the position of the center-of-mass, so fix recenter should come last.

***Fix srd SRD moves may trigger frequent reneighboring***

This is because the SRD particles may move long distances.

***Fix srd grid size > 1/4 of big particle diameter***

This may cause accuracy problems.

***Fix srd particle moved outside valid domain***

This may indicate a problem with your simulation parameters.

***Fix srd particles may move > big particle diameter***

This may cause accuracy problems.

***Fix srd viscosity < 0.0 due to low SRD density***

This may cause accuracy problems.

***Fixes cannot send data in Kokkos communication, switching to classic communication***

This is current restriction with Kokkos.

***For better accuracy use 'pair\_modify table 0'***

The user-specified force accuracy cannot be achieved unless the table feature is disabled by using 'pair\_modify table 0'.

***Geometric mixing assumed for  $1/r^6$  coefficients***

Self-explanatory.

***Group for fix\_modify temp != fix group***

The fix\_modify command is specifying a temperature computation that computes a temperature on a different group of atoms than the fix itself operates on. This is probably not what you want to do.

***H matrix size has been exceeded: m\_fill=%d H.m=%dn***

This is the size of the matrix.

***Ignoring unknown or incorrect info command flag***

Self-explanatory. An unknown argument was given to the info command. Compare your input with the documentation.

***Improper atoms missing at step %ld***

One or more of 4 atoms needed to compute a particular improper are missing on this processor. Typically this is because the pairwise cutoff is set too short or the improper has blown apart and an atom is too far away.

***Improper problem: %d %ld %d %d %d %d***

Conformation of the 4 listed improper atoms is extreme; you may want to check your simulation geometry.

***Improper style in data file differs from currently defined improper style***

Self-explanatory.

***Improper style is defined but no improper style is set***

The topology contains improper style, but there are no improper forces computed since there was no `improper_style` command.

***Inconsistent image flags***

The image flags for a pair on bonded atoms appear to be inconsistent. Inconsistent means that when the coordinates of the two atoms are unwrapped using the image flags, the two atoms are far apart. Specifically they are further apart than half a periodic box length. Or they are more than a box length apart in a non-periodic dimension. This is usually due to the initial data file not having correct image flags for the 2 atoms in a bond that straddles a periodic boundary. They should be different by 1 in that case. This is a warning because inconsistent image flags will not cause problems for dynamics or most LAMMPS simulations. However they can cause problems when such atoms are used with the `fix rigid` or `replicate` commands. Note that if you have an infinite periodic crystal with bonds then it is impossible to have fully consistent image flags, since some bonds will cross periodic boundaries and connect two atoms with the same image flag.

***Increasing communication cutoff for GPU style***

The pair style has increased the communication cutoff to be consistent with the communication cutoff requirements for this pair style when run on the GPU.

***KIM Model does not provide 'energy'; Potential energy will be zero***

Self-explanatory.

***KIM Model does not provide 'forces'; Forces will be zero***

Self-explanatory.

***KIM Model does not provide 'particleEnergy'; energy per atom will be zero***

Self-explanatory.

***KIM Model does not provide 'particleVirial'; virial per atom will be zero***

Self-explanatory.

***Kspace\_modify slab param < 2.0 may cause unphysical behavior***

The `kspace_modify slab` parameter should be larger to insure periodic grids padded with empty space do not overlap.

***Less insertions than requested***

The `fix pour` command was unsuccessful at finding open space for as many particles as it tried to insert.

***Library error in lammmps\_gather\_atoms***

This library function cannot be used if atom IDs are not defined or are not consecutively numbered.

***Library error in lammmps\_scatter\_atoms***

This library function cannot be used if atom IDs are not defined or are not consecutively numbered, or if no atom map is defined. See the `atom_modify` command for details about atom maps.

***Likewise 1-2 special neighbor interactions != 1.0***

The topology contains bonds, but there is no bond style defined and a 1-2 special neighbor scaling factor was not 1.0. This means that pair style interactions may have scaled or missing pairs in the neighbor list in expectation of interactions for those pairs being computed from the bond style.

***Likewise 1-3 special neighbor interactions != 1.0***

The topology contains angles, but there is no angle style defined and a 1-3 special neighbor scaling factor was not 1.0. This means that pair style interactions may have scaled or missing pairs in the neighbor list in expectation of interactions for those pairs being computed from the angle style.

***Likewise 1-4 special neighbor interactions != 1.0***

The topology contains dihedrals, but there is no dihedral style defined and a 1-4 special neighbor scaling factor

was not 1.0. This means that pair style interactions may have scaled or missing pairs in the neighbor list in expectation of interactions for those pairs being computed from the dihedral style.

***Lost atoms via change\_box: original %ld current %ld***

The command options you have used caused atoms to be lost.

***Lost atoms via displace\_atoms: original %ld current %ld***

The command options you have used caused atoms to be lost.

***Lost atoms: original %ld current %ld***

Lost atoms are checked for each time thermo output is done. See the thermo\_modify lost command for options. Lost atoms usually indicate bad dynamics, e.g. atoms have been blown far out of the simulation box, or moved further than one processor's sub-domain away before reneighboring.

***MSM mesh too small, increasing to 2 points in each direction***

Self-explanatory.

***Mismatch between velocity and compute groups***

The temperature computation used by the velocity command will not be on the same group of atoms that velocities are being set for.

***Mixing forced for lj coefficients***

Self-explanatory.

***Molecule attributes do not match system attributes***

An attribute is specified (e.g. diameter, charge) that is not defined for the specified atom style.

***Molecule has bond topology but no special bond settings***

This means the bonded atoms will not be excluded in pairwise interactions.

***Molecule template for create\_atoms has multiple molecules***

The create\_atoms command will only create molecules of a single type, i.e. the first molecule in the template.

***Molecule template for fix gcmc has multiple molecules***

The fix gcmc command will only create molecules of a single type, i.e. the first molecule in the template.

***Molecule template for fix shake has multiple molecules***

The fix shake command will only recognize molecules of a single type, i.e. the first molecule in the template.

***More than one compute centro/atom***

It is not efficient to use compute centro/atom more than once.

***More than one compute cluster/atom***

It is not efficient to use compute cluster/atom more than once.

***More than one compute cna/atom defined***

It is not efficient to use compute cna/atom more than once.

***More than one compute contact/atom***

It is not efficient to use compute contact/atom more than once.

***More than one compute coord/atom***

It is not efficient to use compute coord/atom more than once.

***More than one compute damage/atom***

It is not efficient to use compute ke/atom more than once.

***More than one compute dilatation/atom***

Self-explanatory.

***More than one compute erotate/sphere/atom***

It is not efficient to use compute erotate/sphere/atom more than once.

***More than one compute hexorder/atom***

It is not efficient to use compute hexorder/atom more than once.

***More than one compute ke/atom***

It is not efficient to use compute ke/atom more than once.

***More than one compute orientorder/atom***

It is not efficient to use compute orientorder/atom more than once.

***More than one compute plasticity/atom***

Self-explanatory.

***More than one compute sna/atom***

Self-explanatory.

***More than one compute sna/grid***

Self-explanatory.

***More than one compute sna/grid/local***

Self-explanatory.

***More than one compute snad/atom***

Self-explanatory.

***More than one compute snav/atom***

Self-explanatory.

***More than one fix poems***

It is not efficient to use fix poems more than once.

***More than one fix rigid***

It is not efficient to use fix rigid more than once.

***Neighbor exclusions used with KSpace solver may give inconsistent Coulombic energies***

This is because excluding specific pair interactions also excludes them from long-range interactions which may not be the desired effect. The special\_bonds command handles this consistently by insuring excluded (or weighted) 1-2, 1-3, 1-4 interactions are treated consistently by both the short-range pair style and the long-range solver. This is not done for exclusions of charged atom pairs via the neigh\_modify exclude command.

***New thermo\_style command, previous thermo\_modify settings will be lost***

If a thermo\_style command is used after a thermo\_modify command, the settings changed by the thermo\_modify command will be reset to their default values. This is because the thermo\_modify command acts on the currently defined thermo style, and a thermo\_style command creates a new style.

***No Kspace calculation with verlet/split***

The second partition performs a kspace calculation so the kspace\_style command must be used.

***No automatic unit conversion to XTC file format conventions possible for units lj***

This means no scaling will be performed.

***No fixes defined, atoms won't move***

If you are not using a fix like nve, nvt, npt then atom velocities and coordinates will not be updated during timestepping.

***No joints between rigid bodies, use fix rigid instead***

The bodies defined by fix poems are not connected by joints. POEMS will integrate the body motion, but it would be more efficient to use fix rigid.

***Not using real units with pair reaxff***

This is most likely an error, unless you have created your own ReaxFF parameter file in a different set of units.



***Number of MSM mesh points changed to be a multiple of 2***

MSM requires that the number of grid points in each direction be a multiple of two and the number of grid points in one or more directions have been adjusted to meet this requirement.

***OMP\_NUM\_THREADS environment is not set.***

This environment variable must be set appropriately to use the OPENMP package.

***One or more atoms are time integrated more than once***

This is probably an error since you typically do not want to advance the positions or velocities of an atom more than once per timestep.

***One or more chunks do not contain all atoms in molecule***

This may not be what you intended.

***One or more dynamic groups may not be updated at correct point in timestep***

If there are other fixes that act immediately after the initial stage of time integration within a timestep (i.e. after atoms move), then the command that sets up the dynamic group should appear after those fixes. This will insure that dynamic group assignments are made after all atoms have moved.

***One or more respa levels compute no forces***

This is computationally inefficient.

***Pair COMB charge %.10f with force %.10f hit max barrier***

Something is possibly wrong with your model.

***Pair COMB charge %.10f with force %.10f hit min barrier***

Something is possibly wrong with your model.

***Pair brownian needs newton pair on for momentum conservation***

Self-explanatory.

***Pair dpd needs newton pair on for momentum conservation***

Self-explanatory.

***Pair dsmc: num\_of\_collisions > number\_of\_A***

Collision model in DSMC is breaking down.

***Pair dsmc: num\_of\_collisions > number\_of\_B***

Collision model in DSMC is breaking down.

***Pair style in data file differs from currently defined pair style***

Self-explanatory.

***Pair style restartinfo set but has no restart support***

This pair style has a bug, where it does not support reading and writing information to a restart file, but does not set the member variable “restartinfo” to 0 as required in that case.

***Particle deposition was unsuccessful***

The fix deposit command was not able to insert as many atoms as needed. The requested volume fraction may be too high, or other atoms may be in the insertion region.

***Proc sub-domain size < neighbor skin, could lead to lost atoms***

The decomposition of the physical domain (likely due to load balancing) has led to a processor’s sub-domain being smaller than the neighbor skin in one or more dimensions. Since reneighboring is triggered by atoms moving the skin distance, this may lead to lost atoms, if an atom moves all the way across a neighboring processor’s sub-domain before reneighboring is triggered.

***Reducing PPPM order b/c stencil extends beyond nearest neighbor processor***

This may lead to a larger grid than desired. See the kspace\_modify overlap command to prevent changing of the PPPM order.

***Reducing PPPMDisp Coulomb order b/c stencil extends beyond neighbor processor***

This may lead to a larger grid than desired. See the `kpace_modify overlap` command to prevent changing of the PPPM order.

***Reducing PPPMDisp dispersion order b/c stencil extends beyond neighbor processor***

This may lead to a larger grid than desired. See the `kpace_modify overlap` command to prevent changing of the PPPM order.

***Replacing a fix, but new group != old group***

The ID and style of a fix match for a fix you are changing with a fix command, but the new group you are specifying does not match the old group.

***Replicating in a non-periodic dimension***

The parameters for a replicate command will cause a non-periodic dimension to be replicated; this may cause unwanted behavior.

***Resetting reneighboring criteria during PRD***

A PRD simulation requires that `neigh_modify` settings be `delay = 0`, `every = 1`, `check = yes`. Since these settings were not in place, LAMMPS changed them and will restore them to their original values after the PRD simulation.

***Resetting reneighboring criteria during TAD***

A TAD simulation requires that `neigh_modify` settings be `delay = 0`, `every = 1`, `check = yes`. Since these settings were not in place, LAMMPS changed them and will restore them to their original values after the PRD simulation.

***Resetting reneighboring criteria during minimization***

Minimization requires that `neigh_modify` settings be `delay = 0`, `every = 1`, `check = yes`. Since these settings were not in place, LAMMPS changed them and will restore them to their original values after the minimization.

***Restart file used different # of processors***

The restart file was written out by a LAMMPS simulation running on a different number of processors. Due to round-off, the trajectories of your restarted simulation may diverge a little more quickly than if you ran on the same # of processors.

***Restart file used different 3d processor grid***

The restart file was written out by a LAMMPS simulation running on a different 3d grid of processors. Due to round-off, the trajectories of your restarted simulation may diverge a little more quickly than if you ran on the same # of processors.

***Restart file used different boundary settings, using restart file values***

Your input script cannot change these restart file settings.

***Restart file used different newton bond setting, using restart file value***

The restart file value will override the setting in the input script.

***Restart file used different newton pair setting, using input script value***

The input script value will override the setting in the restart file.

***Restrain problem: %d %ld %d %d %d %d***

Conformation of the 4 listed dihedral atoms is extreme; you may want to check your simulation geometry.

***Running PRD with only one replica***

This is allowed, but you will get no parallel speed-up.

***SRD bin shifting turned on due to small lamda***

This is done to try to preserve accuracy.

***SRD bin size for fix srd differs from user request***

Fix SRD had to adjust the bin size to fit the simulation box. See the `cubic` keyword if you want this message to be an error vs warning.

***SRD bins for fix srd are not cubic enough***

The bin shape is not within tolerance of cubic. See the cubic keyword if you want this message to be an error vs warning.

***SRD particle %d started inside big particle %d on step %ld bounce %d***

See the inside keyword if you want this message to be an error vs warning.

***SRD particle %d started inside wall %d on step %ld bounce %d***

See the inside keyword if you want this message to be an error vs warning.

***Shake determinant < 0.0***

The determinant of the quadratic equation being solved for a single cluster specified by the fix shake command is numerically suspect. LAMMPS will set it to 0.0 and continue.

***Shell command '%s' failed with error '%s'***

Self-explanatory.

***Shell command returned with non-zero status***

This may indicate the shell command did not operate as expected.

***Should not allow rigid bodies to bounce off reflecting walls***

LAMMPS allows this, but their dynamics are not computed correctly.

***Should not use fix nve/limit with fix shake or fix rattle***

This will lead to invalid constraint forces in the SHAKE/RATTLE computation.

***Simulations might be very slow because of large number of structure factors***

Self-explanatory.

***Slab correction not needed for MSM***

Slab correction is intended to be used with Ewald or PPPM and is not needed by MSM.

***Specifying an 'subset' value of '0' is equivalent to no 'subset' keyword***

Self-explanatory.

***System is not charge neutral, net charge = %g***

The total charge on all atoms on the system is not 0.0. For some KSpace solvers this is only a warning.

***Table inner cutoff >= outer cutoff***

You specified an inner cutoff for a Coulombic table that is longer than the global cutoff. Probably not what you wanted.

***Temperature for MSST is not for group all***

User-assigned temperature to MSST fix does not compute temperature for all atoms. Since MSST computes a global pressure, the kinetic energy contribution from the temperature is assumed to also be for all atoms. Thus the pressure used by MSST could be inaccurate.

***Temperature for NPT is not for group all***

User-assigned temperature to NPT fix does not compute temperature for all atoms. Since NPT computes a global pressure, the kinetic energy contribution from the temperature is assumed to also be for all atoms. Thus the pressure used by NPT could be inaccurate.

***Temperature for fix modify is not for group all***

The temperature compute is being used with a pressure calculation which does operate on group all, so this may be inconsistent.

***Temperature for thermo pressure is not for group all***

User-assigned temperature to thermo via the thermo\_modify command does not compute temperature for all atoms. Since thermo computes a global pressure, the kinetic energy contribution from the temperature is assumed to also be for all atoms. Thus the pressure printed by thermo could be inaccurate.

***The fix ave/spatial command has been replaced by the more flexible fix ave/chunk and compute chunk/atom commands – fix ave/spatial will be removed in the summer of 2015***

Self-explanatory.

***The minimizer does not re-orient dipoles when using fix efield***

This means that only the atom coordinates will be minimized, not the orientation of the dipoles.

***Too many common neighbors in CNA %d times***

More than the maximum # of neighbors was found multiple times. This was unexpected.

***Too many inner timesteps in fix ttm***

Self-explanatory.

***Too many neighbors in CNA for %d atoms***

More than the maximum # of neighbors was found multiple times. This was unexpected.

***Triclinic box skew is large***

The displacement in a skewed direction is normally required to be less than half the box length in that dimension. E.g. the xy tilt must be between -half and +half of the x box length. You have relaxed the constraint using the box tilt command, but the warning means that a LAMMPS simulation may be inefficient as a result.

***Use special bonds = 0,1,1 with bond style fene***

Most FENE models need this setting for the special\_bonds command.

***Use special bonds = 0,1,1 with bond style fene/expand***

Most FENE models need this setting for the special\_bonds command.

***Using a many-body potential with bonds/angles/dihedrals and special\_bond exclusions***

This is likely not what you want to do. The exclusion settings will eliminate neighbors in the neighbor list, which the many-body potential needs to calculate its terms correctly.

***Using compute temp/deform with inconsistent fix deform remap option***

Fix nvt/sllod assumes deforming atoms have a velocity profile provided by “remap v” or “remap none” as a fix deform option.

***Using compute temp/deform with no fix deform defined***

This is probably an error, since it makes little sense to use compute temp/deform in this case.

***Using fix srd with box deformation but no SRD thermostat***

The deformation will heat the SRD particles so this can be dangerous.

***Using kspace solver on system with no charge***

Self-explanatory.

***Using largest cut-off for lj/long/dipole/long long long***

Self-explanatory.

***Using largest cutoff for buck/long/coul/long***

Self-explanatory.

***Using largest cutoff for lj/long/coul/long***

Self-explanatory.

***Using largest cutoff for pair\_style lj/long/tip4p/long***

Self-explanatory.

***Using package gpu without any pair style defined***

Self-explanatory.

***Using pair potential shift with pair\_modify compute no***

The shift effects will thus not be computed.

*Using pair tail corrections with nonperiodic system*

This is probably a bogus thing to do, since tail corrections are computed by integrating the density of a periodic system out to infinity.

*Using pair tail corrections with pair\_modify compute no*

The tail corrections will thus not be computed.



**Part II**

**Programmer Guide**





## LAMMPS LIBRARY INTERFACES

As described on the *library interface to LAMMPS* page, LAMMPS can be built as a library (static or shared), so that it can be called by another code, used in a *coupled manner* with other codes, or driven through a *Python script*. Even the LAMMPS standalone executable is essentially a thin wrapper on top of the LAMMPS library, creating a LAMMPS instance, processing input and then existing.

Most of the APIs described below are based on C language wrapper functions in the files `src/library.h` and `src/library.cpp`, but it is also possible to use C++ directly. The basic procedure is always the same: you create one or more instances of *LAMMPS*, pass commands as strings or from files to that LAMMPS instance to execute calculations, and/or call functions that read, manipulate, and update data from the active class instances inside LAMMPS to do analysis or perform operations that are not possible with existing input script commands.

---

### Thread-safety

LAMMPS was initially not conceived as a thread-safe program, but over the years changes have been applied to replace operations that collide with creating multiple LAMMPS instances from multiple-threads of the same process with thread-safe alternatives. This primarily applies to the core LAMMPS code and less so on add-on packages, especially when those packages require additional code in the *lib* folder, interface LAMMPS to Fortran libraries, or the code uses static variables (like the COLVARS package).

Another major issue to deal with is to correctly handle MPI. Creating a LAMMPS instance requires passing an MPI communicator, or it assumes the `MPI_COMM_WORLD` communicator, which spans all MPI processor ranks. When creating multiple LAMMPS object instances from different threads, this communicator has to be different for each thread or else collisions can happen. Or it has to be guaranteed, that only one thread at a time is active. MPI communicators, however, are not a problem, if LAMMPS is compiled with the MPI STUBS library, which implies that there is no MPI communication and only 1 MPI rank.

---

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## 12.1 LAMMPS C Library API

The C library interface is the most commonly used path to manage LAMMPS instances from a compiled code and it is the basis for the *Python* and *Fortran* modules. Almost all functions of the C language API require an argument containing a “handle” in the form of a `void *` type variable, which points to the location of a LAMMPS class instance.

The `library.h` header file by default does not include the `mpi.h` header file and thus hides the `lammps_open()` function which requires the declaration of the `MPI_comm` data type. This is only a problem when the communicator that would be passed is different from `MPI_COMM_WORLD`. Otherwise calling `lammps_open_no_mpi()` will work just as well. To make `lammps_open()` available, you need to compile the code with `-DLAMMPS_LIB_MPI` or add the line `#define LAMMPS_LIB_MPI` before `#include "library.h"`.

Please note the `mpi.h` file must usually be the same (and thus the MPI library in use) for the LAMMPS code and library and the calling code. The exception is when LAMMPS was compiled in serial mode using the STUBS MPI library. In that case the calling code may be compiled with a different MPI library so long as `lammps_open_no_mpi()` is called to create a LAMMPS instance. In that case each MPI rank will run LAMMPS in serial mode.

---

### Errors versus exceptions

If the LAMMPS executable encounters an error condition, it will abort after printing an error message. For a library interface this is usually not desirable. Thus LAMMPS can be compiled to *throw a C++ exception* instead. If enabled, the library functions will catch those exceptions and return. The error status *can be queried* and an *error message retrieved*. We thus recommend enabling C++ exceptions when using the library interface,

---

**Warning:** No checks are made on the arguments of the function calls of the C library interface. *All* function arguments must be non-NULL unless *explicitly* allowed, and must point to consistent and valid data. Buffers for storing returned data must be allocated to a suitable size. Passing invalid or unsuitable information will likely cause crashes or corrupt data.

---

## 12.1.1 Creating or deleting a LAMMPS object

This section documents the following functions:

- `lammps_open()`
  - `lammps_open_no_mpi()`
  - `lammps_open_fortran()`
  - `lammps_close()`
  - `lammps_mpi_init()`
  - `lammps_mpi_finalize()`
  - `lammps_kokkos_finalize()`
  - `lammps_python_finalize()`
- 

The `lammps_open()` and `lammps_open_no_mpi()` functions are used to create and initialize a `LAMMPS()` instance. They return a reference to this instance as a `void *` pointer to be used as the “handle” argument in subsequent function calls until that instance is destroyed by calling `lammps_close()`. Here is a simple example demonstrating its use:

```
#include "library.h"
#include <stdio.h>

int main(int argc, char **argv)
{
    void *handle;
    int version;
    const char *lmpargv[] = { "liblammps", "-log", "none" };
    int lmpargc = sizeof(lmpargv)/sizeof(const char *);
```

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```

/* create LAMMPS instance */
handle = lammps_open_no_mpi(lmpargc, (char **)lmpargv, NULL);
if (handle == NULL) {
    printf("LAMMPS initialization failed");
    lammps_mpi_finalize();
    return 1;
}

/* get and print numerical version code */
version = lammps_version(handle);
printf("LAMMPS Version: %d\n", version);

/* delete LAMMPS instance and shut down MPI */
lammps_close(handle);
lammps_mpi_finalize();
return 0;
}

```

The LAMMPS library uses the MPI library it was compiled with and will either run on all processors in the `MPI_COMM_WORLD` communicator or on the set of processors in the communicator passed as the `comm` argument of `lammps_open()`. This means the calling code can run LAMMPS on all or a subset of processors. For example, a wrapper code might decide to alternate between LAMMPS and another code, allowing them both to run on all the processors. Or it might allocate part of the processors to LAMMPS and the rest to the other code by creating a custom communicator with `MPI_Comm_split()` and running both codes concurrently before syncing them up periodically. Or it might instantiate multiple instances of LAMMPS to perform different calculations and either alternate between them, run them concurrently on split communicators, or run them one after the other. The `lammps_open()` function may be called multiple times for this latter purpose.

The `lammps_close()` function is used to shut down the LAMMPS class pointed to by the handle passed as an argument and free all its memory. This has to be called for every instance created with one of the `lammps_open()` functions. It will, however, **not** call `MPI_Finalize()`, since that may only be called once. See `lammps_mpi_finalize()` for an alternative to invoking `MPI_Finalize()` explicitly from the calling program.

---

```
void *lammps_open(int argc, char **argv, MPI_Comm comm, void **ptr)
```

Create instance of the LAMMPS class and return pointer to it.

The `lammps_open()` function creates a new LAMMPS class instance while passing in a list of strings as if they were *command-line arguments* for the LAMMPS executable, and an MPI communicator for LAMMPS to run under. Since the list of arguments is **exactly** as when called from the command line, the first argument would be the name of the executable and thus is otherwise ignored. However `argc` may be set to 0 and then `argv` may be NULL. If MPI is not yet initialized, `MPI_Init()` will be called during creation of the LAMMPS class instance.

If for some reason the creation or initialization of the LAMMPS instance fails a null pointer is returned.

Changed in version 18Sep2020: This function now has the pointer to the created LAMMPS class instance as return value. For backward compatibility it is still possible to provide the address of a pointer variable as final argument `ptr`.

Deprecated since version 18Sep2020: The `ptr` argument will be removed in a future release of LAMMPS. It should be set to NULL instead.

**See also**

`lammps_open_no_mpi()`, `lammps_open_fortran()`

**Note:** This function is **only** declared when the code using the LAMMPS `library.h` include file is compiled with `-DLAMMPS_LIB_MPI`, or contains a `#define LAMMPS_LIB_MPI 1` statement before `#include "library.h"`. Otherwise you can only use the `lammps_open_no_mpi()` or `lammps_open_fortran()` functions.

---

#### Parameters

- **argc** – number of command line arguments
- **argv** – list of command line argument strings
- **comm** – MPI communicator for this LAMMPS instance
- **ptr** – pointer to a void pointer variable which serves as a handle; may be NULL

#### Returns

pointer to new LAMMPS instance cast to void \*

---

void \***lammps\_open\_no\_mpi**(int argc, char \*\*argv, void \*\*ptr)

Variant of `lammps_open()` that implicitly uses `MPI_COMM_WORLD`.

This function is a version of `lammps_open()`, that is missing the MPI communicator argument. It will use `MPI_COMM_WORLD` instead. The type and purpose of arguments and return value are otherwise the same.

Outside of the convenience, this function is useful, when the LAMMPS library was compiled in serial mode, but the calling code runs in parallel and the `MPI_Comm` data type of the STUBS library would not be compatible with that of the calling code.

If for some reason the creation or initialization of the LAMMPS instance fails a null pointer is returned.

Changed in version 18Sep2020: This function now has the pointer to the created LAMMPS class instance as return value. For backward compatibility it is still possible to provide the address of a pointer variable as final argument *ptr*.

Deprecated since version 18Sep2020: The *ptr* argument will be removed in a future release of LAMMPS. It should be set to NULL instead.

#### See also

`lammps_open()`, `lammps_open_fortran()`

#### Parameters

- **argc** – number of command line arguments
- **argv** – list of command line argument strings
- **ptr** – pointer to a void pointer variable which serves as a handle; may be NULL

#### Returns

pointer to new LAMMPS instance cast to void \*

---

void \***lammmps\_open\_fortran**(int argc, char \*\*argv, int f\_comm)

Variant of `lammmps_open()` using a Fortran MPI communicator.

This function is a version of `lammmps_open()`, that uses an integer for the MPI communicator as the MPI Fortran interface does. It is used in the `lammmps()` constructor of the LAMMPS Fortran module. Internally it converts the `f_comm` argument into a C-style MPI communicator with `MPI_Comm_f2c()` and then calls `lammmps_open()`.

If for some reason the creation or initialization of the LAMMPS instance fails a null pointer is returned.

New in version 18Sep2020.

*See also*

`lammmps_open_fortran()`, `lammmps_open_no_mpi()`

#### Parameters

- **argc** – number of command line arguments
- **argv** – list of command line argument strings
- **f\_comm** – Fortran style MPI communicator for this LAMMPS instance

#### Returns

pointer to new LAMMPS instance cast to void \*

---

void **lammmps\_close**(void \*handle)

Delete a LAMMPS instance created by `lammmps_open()` or its variants.

This function deletes the LAMMPS class instance pointed to by `handle` that was created by one of the `lammmps_open()` variants. It does **not** call `MPI_Finalize()` to allow creating and deleting multiple LAMMPS instances concurrently or sequentially. See `lammmps_mpi_finalize()` for a function performing this operation.

#### Parameters

**handle** – pointer to a previously created LAMMPS instance

---

void **lammmps\_mpi\_init**()

Ensure the MPI environment is initialized.

The MPI standard requires that any MPI application must call `MPI_Init()` exactly once before performing any other MPI function calls. This function checks, whether MPI is already initialized and calls `MPI_Init()` in case it is not.

New in version 18Sep2020.

---

void **lammmps\_mpi\_finalize**()

Shut down the MPI infrastructure.

The MPI standard requires that any MPI application calls `MPI_Finalize()` before exiting. Even if a calling program does not do any MPI calls, MPI is still initialized internally to avoid errors accessing any MPI functions. This function should then be called right before exiting the program to wait until all (parallel) tasks are completed and then MPI is cleanly shut down. After calling this function no more MPI calls may be made.

New in version 18Sep2020.

*See also*

`lammps_kokkos_finalize()`, `lammps_python_finalize()`

---

void `lammps_kokkos_finalize()`

Shut down the Kokkos library environment.

The Kokkos library may only be initialized once during the execution of a process. This is done automatically the first time Kokkos functionality is used. This requires that the Kokkos environment must be explicitly shut down after any LAMMPS instance using it is closed (to release associated resources). After calling this function no Kokkos functionality may be used.

New in version 2Jul2021.

*See also*

`lammps_mpi_finalize()`, `lammps_python_finalize()`

---

void `lammps_python_finalize()`

Clear the embedded Python environment

This function resets and clears an embedded Python environment by calling the `Py_Finalize()` function of the embedded Python library, if enabled. This call would free up all allocated resources and release loaded shared objects.

However, this is **not** done when a LAMMPS instance is deleted because a) LAMMPS may have been used through the Python module and thus the Python interpreter is external and not embedded into LAMMPS and therefore may not be reset by LAMMPS b) some Python modules and extensions, most notably NumPy, are not compatible with being initialized multiple times, which would happen if additional LAMMPS instances using Python would be created *after* after calling `Py_Finalize()`.

This function can be called to explicitly clear the Python environment in case it is safe to do so.

New in version 20Sep2021.

*See also*

`lammps_mpi_finalize()`, `lammps_kokkos_finalize()`

---

## 12.1.2 Executing commands

This section documents the following functions:

- `lammps_file()`
- `lammps_command()`
- `lammps_commands_list()`
- `lammps_commands_string()`

Once a LAMMPS instance is created, there are multiple ways to “drive” a simulation. In most cases it is easiest to process single or multiple LAMMPS commands like in an input file. This can be done through reading a file or passing single commands or lists of commands or blocks of commands with the following functions.

Via these functions, the calling code can have LAMMPS act on a series of *input file commands* that are either read from a file or passed as strings. For example, this allows setup of a problem from an input script, and then running it in stages while performing other operations in between or concurrently. The caller can interleave the LAMMPS function calls with operations it performs, such as calls to extract information from or set information within LAMMPS, or calls to another code’s library.

Just as with *input script parsing* comments can be included in the file or strings, and expansion of variables with `${name}` or `$(expression)` syntax is performed. Below is a short example using some of these functions.

```
/* define to make the otherwise hidden prototype for "lammps_open()" visible */
#define LAMMPS_LIB_MPI
#include "library.h"

#include <mpi.h>
#include <stdio.h>

int main(int argc, char **argv)
{
    void *handle;
    int i;

    MPI_Init(&argc, &argv);
    handle = lammps_open(0, NULL, MPI_COMM_WORLD, NULL);
    lammps_file(handle, "in.sysinit");
    lammps_command(handle, "run 1000 post no");

    for (i=0; i < 100; ++i) {
        lammps_commands_string(handle, "run 100 pre no post no\n"
                                   "print 'PE = $(pe)'\n"
                                   "print 'KE = $(ke)'\n");
    }
    lammps_close(handle);
    MPI_Finalize();
    return 0;
}
```

void **lammers\_file**(void \*handle, const char \*file)

Process LAMMPS input from a file.

This function processes commands in the file pointed to by *filename* line by line and thus functions very similar to the *include* command. The function returns when the end of the file is reached and the commands have completed.

The actual work is done by the functions *Input::file(const char \*)* and *Input::file()*.

### Parameters

- **handle** – pointer to a previously created LAMMPS instance
  - **filename** – name of a file with LAMMPS input
- 

char \***lammers\_command**(void \*handle, const char \*cmd)

Process a single LAMMPS input command from a string.

This function tells LAMMPS to execute the single command in the string *cmd*. The entire string is considered as command and need not have a (final) newline character. Newline characters in the body of the string, however, will be treated as part of the command and will **not** start a second command. The function *lammers\_commands\_string()* processes a string with multiple command lines.

The function returns the name of the command on success or NULL when passing a string without a command.

### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **cmd** – string with a single LAMMPS command

### Returns

string with parsed command name or NULL

---

void **lammers\_commands\_list**(void \*handle, int ncmd, const char \*\*cmds)

Process multiple LAMMPS input commands from list of strings.

This function processes multiple commands from a list of strings by first concatenating the individual strings in *cmds* into a single string, inserting newline characters as needed. The combined string is passed to *lammers\_commands\_string()* for processing.

### Parameters

- **handle** – pointer to a previously created LAMMPS instance
  - **ncmd** – number of lines in *cmds*
  - **cmds** – list of strings with LAMMPS commands
-



void **lammops\_commands\_string**(void \*handle, const char \*str)

Process a block of LAMMPS input commands from a single string.

This function processes a multi-line string similar to a block of commands from a file. The string may have multiple lines (separated by newline characters) and also single commands may be distributed over multiple lines with continuation characters ('&'). Those lines are combined by removing the '&' and the following newline character. After this processing the string is handed to LAMMPS for parsing and executing.

---

**Note:** Multi-line commands enabled by triple quotes will NOT work with this function.

---

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **str** – string with block of LAMMPS input commands

### 12.1.3 System properties

This section documents the following functions:

- *lammops\_get\_natoms()*
- *lammops\_get\_thermo()*
- *lammops\_extract\_box()*
- *lammops\_reset\_box()*
- *lammops\_memory\_usage()*
- *lammops\_get\_mpi\_comm()*
- *lammops\_extract\_setting()*
- *lammops\_extract\_global\_datatype()*
- *lammops\_extract\_global()*

The library interface allows extraction of different kinds of information about the active simulation instance and also modifications to it. This enables combining of a LAMMPS simulation with other processing and simulation methods computed by the calling code, or by another code that is coupled to LAMMPS via the library interface. In some cases the data returned is direct reference to the original data inside LAMMPS, cast to a void pointer. In that case the data needs to be cast to a suitable pointer for the calling program to access it, and you may need to know the correct dimensions and lengths. This also means you can directly change those value(s) from the calling program, e.g. to modify atom positions. Of course, this should be done with care. When accessing per-atom data, please note that this data is the per-processor **local** data and is indexed accordingly. Per-atom data can change sizes and ordering at every neighbor list rebuild or atom sort event as atoms migrate between sub-domains and processors.

```
#include "library.h"
#include <stdio.h>

int main(int argc, char **argv)
{
    void *handle;
    int i;
```

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```

handle = lammps_open_no_mpi(0, NULL, NULL);
lammps_file(handle, "in.sysinit");
printf("Running a simulation with %g atoms.\n",
       lammps_get_natoms(handle));

printf(" %d local and %d ghost atoms. %d atom types\n",
       lammps_extract_setting(handle, "nlocal"),
       lammps_extract_setting(handle, "nghost"),
       lammps_extract_setting(handle, "ntypes"));

double *dt = (double *)lammps_extract_global(handle, "dt");
printf("Changing timestep from %g to 0.5\n", *dt);
*dt = 0.5;

lammps_command(handle, "run 1000 post no");

for (i=0; i < 10; ++i) {
  lammps_command(handle, "run 100 pre no post no");
  printf("PE = %g\nKE = %g\n",
        lammps_get_thermo(handle, "pe"),
        lammps_get_thermo(handle, "ke"));
}
lammps_close(handle);
return 0;
}

```

double **lammps\_get\_natoms**(void \*handle)

Return the total number of atoms in the system.

This number may be very large when running large simulations across multiple processors. Depending on compile time choices, LAMMPS may be using either 32-bit or a 64-bit integer to store this number. For portability this function returns thus a double precision floating point number, which can represent up to a 53-bit signed integer exactly ( $\approx 10^{16}$ ).

As an alternative, you can use `lammps_extract_global()` and cast the resulting pointer to an integer pointer of the correct size and dereference it. The size of that integer (in bytes) can be queried by calling `lammps_extract_setting()` to return the size of a `bigint` integer.

Changed in version 18Sep2020: The type of the return value was changed from `int` to `double` to accommodate reporting atom counts for larger systems that would overflow a 32-bit `int` without having to depend on a 64-bit integer type definition.

#### Parameters

**handle** – pointer to a previously created LAMMPS instance

#### Returns

total number of atoms or 0 if value is too large

double **lammops\_get\_thermo**(void \*handle, const char \*keyword)

Get current value of a thermo keyword.

This function returns the current value of a *thermo keyword*. Unlike *lammops\_extract\_global()* it does not give access to the storage of the desired data but returns its value as a `double`, so it can also return information that is computed on-the-fly.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **keyword** – string with the name of the thermo keyword

#### Returns

value of the requested thermo property or 0.0

---

void **lammops\_extract\_box**(void \*handle, double \*boxlo, double \*boxhi, double \*xy, double \*yz, double \*xz, int \*pflags, int \*boxflag)

Extract simulation box parameters.

This function (re-)initializes the simulation box and boundary information and then assign the designated data to the locations in the pointers passed as arguments. Any argument (except the first) may be a NULL pointer and then will not be assigned.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance
  - **boxlo** – pointer to 3 doubles where the lower box boundary is stored
  - **boxhi** – pointer to 3 doubles where the upper box boundary is stored
  - **xy** – pointer to a double where the xy tilt factor is stored
  - **yz** – pointer to a double where the yz tilt factor is stored
  - **xz** – pointer to a double where the xz tilt factor is stored
  - **pflags** – pointer to 3 ints, set to 1 for periodic boundaries and 0 for non-periodic
  - **boxflag** – pointer to an int, which is set to 1 if the box will be changed during a simulation by a fix and 0 if not.
- 

void **lammops\_reset\_box**(void \*handle, double \*boxlo, double \*boxhi, double xy, double yz, double xz)

Reset simulation box parameters.

This function sets the simulation box dimensions (upper and lower bounds and tilt factors) from the provided data and then re-initializes the box information and all derived settings. It may only be called before atoms are created.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **boxlo** – pointer to 3 doubles containing the lower box boundary
- **boxhi** – pointer to 3 doubles containing the upper box boundary
- **xy** – xy tilt factor
- **yz** – yz tilt factor

- **xz** – xz tilt factor
- 

void **lammops\_memory\_usage**(void \*handle, double \*meminfo)

Get memory usage information

This function will retrieve memory usage information for the current LAMMPS instance or process. The *meminfo* buffer will be filled with 3 different numbers (if supported by the operating system). The first is the tally (in MBytes) of all large memory allocations made by LAMMPS. This is a lower boundary of how much memory is requested and does not account for memory allocated on the stack or allocations via *new*. The second number is the current memory allocation of the current process as returned by a memory allocation reporting in the system library. The third number is the maximum amount of RAM (not swap) used by the process so far. If any of the two latter parameters is not supported by the operating system it will be set to zero.

New in version 18Sep2020.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance
  - **meminfo** – buffer with space for at least 3 double to store data in.
- 

int **lammops\_get\_mpi\_comm**(void \*handle)

Return current LAMMPS world communicator as integer

This will take the LAMMPS “world” communicator and convert it to an integer using `MPI_Comm_c2f()`, so it is equivalent to the corresponding MPI communicator in Fortran. This way it can be safely passed around between different programming languages. To convert it to the C language representation use `MPI_Comm_f2c()`.

If LAMMPS was compiled with `MPI_STUBS`, this function returns -1.

New in version 18Sep2020.

#### See also

[\*lammops\\_open\\_fortran\(\)\*](#)

#### Parameters

**handle** – pointer to a previously created LAMMPS instance

#### Returns

Fortran representation of the LAMMPS world communicator

---

int **lammops\_extract\_setting**(void \*handle, const char \*keyword)

Query LAMMPS about global settings.

This function will retrieve or compute global properties. In contrast to [\*lammops\\_get\\_thermo\(\)\*](#) this function returns an `int`. The following tables list the currently supported keyword. If a keyword is not recognized, the function returns -1.

- *Integer sizes*
-

- *System status*
- *System sizes*
- *Atom style flags*

### Integer sizes

Keyword	Description / Return value
bigint	size of the bigint integer type, 4 or 8 bytes. Set at <i>compile time</i> .
tagint	size of the tagint integer type, 4 or 8 bytes. Set at <i>compile time</i> .
imageint	size of the imageint integer type, 4 or 8 bytes. Set at <i>compile time</i> .

### System status

Keyword	Description / Return value
dimension	Number of dimensions: 2 or 3. See <i>dimension command</i> .
box_exist	1 if the simulation box is defined, 0 if not. See <i>create_box command</i> .
nthreads	Number of requested OpenMP threads for LAMMPS' execution
newton_bond	1 if Newton's 3rd law is applied to bonded interactions, 0 if not.
newton_pair	1 if Newton's 3rd law is applied to non-bonded interactions, 0 if not.
triclinic	1 if the the simulation box is triclinic, 0 if orthogonal. See <i>change_box command</i> .
universe_rank	MPI rank on LAMMPS' universe communicator (0 <= universe_rank < universe_size)
universe_size	Number of ranks on LAMMPS' universe communicator (world_size <= universe_size)
world_rank	MPI rank on LAMMPS' world communicator (0 <= world_rank < world_size)
world_size	Number of ranks on LAMMPS' world communicator

### System sizes

Keyword	Description / Return value
nlocal	number of "owned" atoms of the current MPI rank.
nghost	number of "ghost" atoms of the current MPI rank.
nall	number of all "owned" and "ghost" atoms of the current MPI rank.
nmax	maximum of nlocal+nghost across all MPI ranks (for per-atom data array size).
ntypes	number of atom types
nbondtypes	number of bond types
nangletypes	number of angle types
ndihedraltypes	number of dihedral types
nimproptypes	number of improper types
nellipsoids	number of atoms that have ellipsoid data
nlines	number of atoms that have line data (see <i>pair style line/lj</i> )
ntris	number of atoms that have triangle data (see <i>pair style tri/lj</i> )
nbodies	number of atoms that have body data (see <i>the Body particle HowTo</i> )

### Atom style flags

Keyword	Description / Return value
molecule_flag	1 if the atom style includes molecular topology data. See <i>atom_style command</i> .
q_flag	1 if the atom style includes point charges. See <i>atom_style command</i> .
mu_flag	1 if the atom style includes point dipoles. See <i>atom_style command</i> .
rmass_flag	1 if the atom style includes per-atom masses, 0 if there are per-type masses. See <i>atom_style command</i> .
radius_flag	1 if the atom style includes a per-atom radius. See <i>atom_style command</i> .
sphere_flag	1 if the atom style describes extended particles that can rotate. See <i>atom_style command</i> .
ellipsoid_flag	1 if the atom style describes extended particles that may be ellipsoidal. See <i>atom_style command</i> .
omega_flag	1 if the atom style can store per-atom rotational velocities. See <i>atom_style command</i> .
torque_flag	1 if the atom style can store per-atom torques. See <i>atom_style command</i> .
angmom_flag	1 if the atom style can store per-atom angular momentum. See <i>atom_style command</i> .

*See also*

*lammps\_extract\_global()*

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance
- **keyword** – string with the name of the thermo keyword

**Returns**

value of the queried setting or -1 if unknown

---

int **lammps\_extract\_global\_datatype**(void \*handle, const char \*name)

Get data type of internal global LAMMPS variables or arrays.

This function returns an integer that encodes the data type of the global property with the specified name. See *\_LMP\_DATATYPE\_CONST* for valid values. Callers of *lammps\_extract\_global()* can use this information to then decide how to cast the (void\*) pointer and access the data.

New in version 18Sep2020.

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance (unused)
- **name** – string with the name of the extracted property

**Returns**

integer constant encoding the data type of the property or -1 if not found.

---

void \***lammps\_extract\_global**(void \*handle, const char \*name)

Get pointer to internal global LAMMPS variables or arrays.

This function returns a pointer to the location of some global property stored in one of the constituent classes of a LAMMPS instance. The returned pointer is cast to `void *` and needs to be cast to a pointer of the type that the entity represents. The pointers returned by this function are generally persistent; therefore it is not necessary to

---

call the function again, unless a *clear command* command is issued which wipes out and recreates the contents of the *LAMMPS* class.

Please also see *lammps\_extract\_setting()*, *lammps\_get\_thermo()*, and *lammps\_extract\_box()*.

The following tables list the supported names, their data types, length of the data area, and a short description. The data type can also be queried through calling *lammps\_extract\_global\_datatype()*. The *bigint* type may be defined to be either an *int* or an *int64\_t*. This is set at *compile time* of the LAMMPS library and can be queried through calling *lammps\_extract\_setting()*. The function *lammps\_extract\_global\_datatype()* will directly report the “native” data type. The following tables are provided:

- *Timestep settings*
- *Simulation box settings*
- *System property settings*
- *Unit settings*

### Timestep settings

Name	Type	Length	Description
dt	double	1	length of the time step. See <i>timestep command</i> .
ntimestep	bigint	1	current time step number. See <i>reset_timestep command</i> .
atime	double	1	accumulated simulation time in time units.
atimestep	bigint	1	the number of the timestep when “atime” was last updated.
respa_levels	int	1	number of r-RESPA levels. See <i>run_style command</i> .
respa_dt	double	number of r-RESPA levels	length of the time steps with r-RESPA. See <i>run_style command</i> .

### Simulation box settings

Name	Type	Length	Description
boxlo	double	3	lower box boundaries. See <i>create_box command</i> .
boxhi	double	3	upper box boundaries. See <i>create_box command</i> .
boxxlo	double	1	lower box boundary in x-direction. See <i>create_box command</i> .
boxxhi	double	1	upper box boundary in x-direction. See <i>create_box command</i> .
boxylo	double	1	lower box boundary in y-direction. See <i>create_box command</i> .
boxyhi	double	1	upper box boundary in y-direction. See <i>create_box command</i> .
boxzlo	double	1	lower box boundary in z-direction. See <i>create_box command</i> .
boxzhi	double	1	upper box boundary in z-direction. See <i>create_box command</i> .
sublo	double	3	subbox lower boundaries
subhi	double	3	subbox upper boundaries
sublo_lambda	double	3	subbox lower boundaries in fractional coordinates (for triclinic cells)
subhi_lambda	double	3	subbox upper boundaries in fractional coordinates (for triclinic cells)
periodicity	int	3	0 if non-periodic, 1 if periodic for x, y, and z; See <i>boundary command</i> .
triclinic	int	1	1 if the the simulation box is triclinic, 0 if orthogonal; See <i>change_box command</i> .
xy	double	1	triclinic tilt factor. See <i>Triclinic (non-orthogonal) simulation boxes</i> .
yz	double	1	triclinic tilt factor. See <i>Triclinic (non-orthogonal) simulation boxes</i> .
xz	double	1	triclinic tilt factor. See <i>Triclinic (non-orthogonal) simulation boxes</i> .

### System property settings



Name	Type	Length	Description
ntypes	int	1	number of atom types
nbonds	big-int	1	total number of bonds in the simulation.
nangles	big-int	1	total number of angles in the simulation.
ndihedrals	big-int	1	total number of dihedrals in the simulation.
nimpropers	big-int	1	total number of impropers in the simulation.
natoms	big-int	1	total number of atoms in the simulation.
nlocal	int	1	number of “owned” atoms of the current MPI rank.
nghost	int	1	number of “ghost” atoms of the current MPI rank.
nmax	int	1	maximum of nlocal+nghost across all MPI ranks (for per-atom data array size).
q_flag	int	1	<b>deprecated.</b> Use <code>lammops_extract_setting()</code> instead.

### Unit settings

Name	Type	Length	Description
units	char *	1	string with the current unit style. See <i>units command</i> .
boltz	double	1	value of the “boltz” constant. See <i>units command</i> .
hplanck	double	1	value of the “hplanck” constant. See <i>units command</i> .
mvv2e	double	1	factor to convert $\frac{1}{2}mv^2$ for a particle to the current energy unit; See <i>units command</i> .
ftm2v	double	1	(description missing) See <i>units command</i> .
mv2d	double	1	(description missing) See <i>units command</i> .
nktv2p	double	1	(description missing) See <i>units command</i> .
qqr2e	double	1	factor to convert $\frac{q_i q_j}{r}$ to energy units; See <i>units command</i> .
qe2f	double	1	(description missing) See <i>units command</i> .
vxmu2f	double	1	(description missing) See <i>units command</i> .
xxt2kmu	double	1	(description missing) See <i>units command</i> .
dielectric	double	1	value of the dielectric constant. See <i>dielectric command</i> .
qqr2e	double	1	(description missing) See <i>units command</i> .
e_mass	double	1	(description missing) See <i>units command</i> .
hhmrr2e	double	1	(description missing) See <i>units command</i> .
mvh2r	double	1	(description missing) See <i>units command</i> .
angstrom	double	1	constant to convert current length unit to angstroms; 1.0 for reduced (aka “lj”) units. See <i>units command</i> .
fem- tosecond	double	1	constant to convert current time unit to femtoseconds; 1.0 for reduced (aka “lj”) units
qelectron	double	1	(description missing) See <i>units command</i> .

**Warning:** Modifying the data in the location pointed to by the returned pointer may lead to inconsistent internal data and thus may cause failures or crashes or bogus simulations. In general it is thus usually better to use a LAMMPS input command that sets or changes these parameters. Those will takes care of all side effects and necessary updates of settings derived from such settings. Where possible a reference to such a command or a relevant section of the manual is given below.

### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **name** – string with the name of the extracted property

**Returns**

pointer (cast to `void *`) to the location of the requested property. NULL if name is not known.

### 12.1.4 Per-atom properties

This section documents the following functions:

- `lammmps_extract_atom_datatype()`
- `lammmps_extract_atom()`

---

```
int lammmps_extract_atom_datatype(void *handle, const char *name)
```

Get data type of a LAMMPS per-atom property

This function returns an integer that encodes the data type of the per-atom property with the specified name. See `_LMP_DATATYPE_CONST` for valid values. Callers of `lammmps_extract_atom()` can use this information to then decide how to cast the (`void*`) pointer and access the data.

New in version 18Sep2020.

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance
- **name** – string with the name of the extracted property

**Returns**

integer constant encoding the data type of the property or -1 if not found.

---

```
void *lammmps_extract_atom(void *handle, const char *name)
```

Get pointer to a LAMMPS per-atom property.

This function returns a pointer to the location of per-atom properties (and per-atom-type properties in the case of the ‘mass’ keyword). Per-atom data is distributed across sub-domains and thus MPI ranks. The returned pointer is cast to `void *` and needs to be cast to a pointer of data type that the entity represents.

A table with supported keywords is included in the documentation of the `Atom::extract()` function.

**Warning:** The pointers returned by this function are generally not persistent since per-atom data may be re-distributed, re-allocated, and re-ordered at every re-neighborhood operation.

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance
- **name** – string with the name of the extracted property

**Returns**

pointer (cast to `void *`) to the location of the requested data or NULL if not found.

## 12.1.5 Compute, fixes, variables

This section documents accessing or modifying data stored by computes, fixes, or variables in LAMMPS using the following functions:

- `lammops_extract_compute()`
- `lammops_extract_fix()`
- `lammops_extract_variable()`
- `lammops_set_variable()`

---

```
void *lammops_extract_compute(void *handle, const char*, int, int)
```

Get pointer to data from a LAMMPS compute.

This function returns a pointer to the location of data provided by a *compute command* instance identified by the compute-ID. Computes may provide global, per-atom, or local data, and those may be a scalar, a vector, or an array or they may provide the information about the dimensions of the respective data. Since computes may provide multiple kinds of data, it is required to set style and type flags representing what specific data is desired. This also determines to what kind of pointer the returned pointer needs to be cast to access the data correctly. The function returns NULL if the compute ID is not found or the requested data is not available or current. The following table lists the available options.

Style (see <code>_LMP_STYLE_CONST</code> )	Type (see <code>_LMP_TYPE_CONST</code> )	Returned type	Returned data
LMP_STYLE_GLOBAL	LMP_TYPE_SCALAR	double *	Global scalar
LMP_STYLE_GLOBAL	LMP_TYPE_VECTOR	double *	Global vector
LMP_STYLE_GLOBAL	LMP_TYPE_ARRAY	double **	Global array
LMP_STYLE_GLOBAL	LMP_SIZE_VECTOR	int *	Length of global vector
LMP_STYLE_GLOBAL	LMP_SIZE_ROWS	int *	Rows of global array
LMP_STYLE_GLOBAL	LMP_SIZE_COLS	int *	Columns of global array
LMP_STYLE_ATOM	LMP_TYPE_VECTOR	double *	Per-atom value
LMP_STYLE_ATOM	LMP_TYPE_ARRAY	double **	Per-atom vector
LMP_STYLE_ATOM	LMP_SIZE_COLS	int *	Columns in per-atom array, 0 if vector
LMP_STYLE_LOCAL	LMP_TYPE_VECTOR	double *	Local data vector
LMP_STYLE_LOCAL	LMP_TYPE_ARRAY	double **	Local data array
LMP_STYLE_LOCAL	LMP_SIZE_VECTOR	int *	Alias for using LMP_SIZE_ROWS
LMP_STYLE_LOCAL	LMP_SIZE_ROWS	int *	Number of local array rows or length of vector
LMP_STYLE_LOCAL	LMP_SIZE_COLS	int *	Number of local array columns, 0 if vector

---

**Note:** If the compute's data is not computed for the current step, the compute will be invoked. LAMMPS cannot easily check at that time, if it is valid to invoke a compute, so it may fail with an error. The caller has to check to avoid such an error.

---

**Warning:** The pointers returned by this function are generally not persistent since the computed data may be re-distributed, re-allocated, and re-ordered at every invocation. It is advisable to re-invoke this function before the data is accessed, or make a copy if the data shall be used after other LAMMPS commands have been issued.

### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **id** – string with ID of the compute
- **style** – constant indicating the style of data requested (global, per-atom, or local)
- **type** – constant indicating type of data (scalar, vector, or array) or size of rows or columns

### Returns

pointer (cast to `void *`) to the location of the requested data or NULL if not found.

---

```
void *lammops_extract_fix(void *handle, const char*, int, int, int, int)
```

Get pointer to data from a LAMMPS fix.

This function returns a pointer to data provided by a *fix command* instance identified by its fix-ID. Fixes may provide global, per-atom, or local data, and those may be a scalar, a vector, or an array, or they may provide the information about the dimensions of the respective data. Since individual fixes may provide multiple kinds of data, it is required to set style and type flags representing what specific data is desired. This also determines to what kind of pointer the returned pointer needs to be cast to access the data correctly. The function returns NULL if the fix ID is not found or the requested data is not available.

The following table lists the available options.

Style (see <a href="#">_LMP_STYLE_CONST</a> )	Type (see <a href="#">_LMP_TYPE_CONST</a> )	Returned type	Returned data
LMP_STYLE_GLOBAL	LMP_TYPE_SCALAR	double *	Copy of global scalar
LMP_STYLE_GLOBAL	LMP_TYPE_VECTOR	double *	Copy of global vector element at index nrow
LMP_STYLE_GLOBAL	LMP_TYPE_ARRAY	double *	Copy of global array element at nrow, ncol
LMP_STYLE_GLOBAL	LMP_SIZE_VECTOR	int *	Length of global vector
LMP_STYLE_GLOBAL	LMP_SIZE_ROWS	int *	Rows in global array
LMP_STYLE_GLOBAL	LMP_SIZE_COLS	int *	Columns in global array
LMP_STYLE_ATOM	LMP_TYPE_VECTOR	double *	Per-atom value
LMP_STYLE_ATOM	LMP_TYPE_ARRAY	double **	Per-atom vector
LMP_STYLE_ATOM	LMP_SIZE_COLS	int *	Columns of per-atom array, 0 if vector
LMP_STYLE_LOCAL	LMP_TYPE_VECTOR	double *	Local data vector
LMP_STYLE_LOCAL	LMP_TYPE_ARRAY	double **	Local data array
LMP_STYLE_LOCAL	LMP_SIZE_ROWS	int *	Number of local data rows
LMP_STYLE_LOCAL	LMP_SIZE_COLS	int *	Number of local data columns

---

**Note:** When requesting global data, the fix data can only be accessed one item at a time without access to the pointer itself. Thus this function will allocate storage for a single double value, copy the returned value to it,

and returns a pointer to the location of the copy. Therefore the allocated storage needs to be freed after its use to avoid a memory leak. Example:

```
double *dptr = (double *) lammps_extract_fix(handle,name,0,1,0,0);
double value = *dptr;
lammps_free((void *)dptr);
```

---

**Note:** LAMMPS cannot easily check if it is valid to access the data, so it may fail with an error. The caller has to avoid such an error.

---

**Warning:** The pointers returned by this function for per-atom or local data are generally not persistent, since the computed data may be re-distributed, re-allocated, and re-ordered at every invocation of the fix. It is thus advisable to re-invoke this function before the data is accessed, or make a copy, if the data shall be used after other LAMMPS commands have been issued.

### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **id** – string with ID of the fix
- **style** – constant indicating the style of data requested (global, per-atom, or local)
- **type** – constant indicating type of data (scalar, vector, or array) or size of rows or columns
- **nrow** – row index (only used for global vectors and arrays)
- **ncol** – column index (only used for global arrays)

### Returns

pointer (cast to `void *`) to the location of the requested data or NULL if not found.

---

`void *lammps_extract_variable(void *handle, const char*, const char*)`

Get pointer to data from a LAMMPS variable.

This function returns a pointer to data from a LAMMPS *variable command* identified by its name. When the variable is either an *equal*-style compatible or an *atom*-style variable the variable is evaluated and the corresponding value(s) returned. Variables of style *internal* are compatible with *equal*-style variables and so are *python*-style variables, if they return a numeric value. For other variable styles their string value is returned. The function returns NULL when a variable of the provided *name* is not found or of an incompatible style. The *group* argument is only used for *atom*-style variables and ignored otherwise. If set to NULL when extracting data from an *atom*-style variable, the group is assumed to be “all”.

When requesting data from an *equal*-style or compatible variable this function allocates storage for a single double value, copies the returned value to it, and returns a pointer to the location of the copy. Therefore the allocated storage needs to be freed after its use to avoid a memory leak. Example:

```
double *dptr = (double *) lammps_extract_variable(handle,name,NULL);
double value = *dptr;
lammps_free((void *)dptr);
```

For *atom*-style variables the data returned is a pointer to an allocated block of storage of double of the length `atom->nlocal`. Since the data is returned a copy, the location will persist, but its content will not be updated, in case the variable is re-evaluated. To avoid a memory leak this pointer needs to be freed after use in the calling program.

For other variable styles the returned pointer needs to be cast to a char pointer.

```
const char *cptr = (const char *) lammps_extract_variable(handle,name,NULL);
printf("The value of variable %s is %s\n", name, cptr);
```

---

**Note:** LAMMPS cannot easily check if it is valid to access the data referenced by the variables, e.g. computes or fixes or thermodynamic info, so it may fail with an error. The caller has to make certain, that the data is extracted only when it safe to evaluate the variable and thus an error and crash is avoided.

---

### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **name** – name of the variable
- **group** – group-ID for atom style variable or NULL

### Returns

pointer (cast to `void *`) to the location of the requested data or NULL if not found.

---

int **lammps\_set\_variable**(void\*, char\*, char\*)

Set the value of a string-style variable.

This function assigns a new value from the string `str` to the string-style variable `name`. Returns -1 if a variable of that name does not exist or is not a string-style variable, otherwise 0.

### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **name** – name of the variable
- **str** – new value of the variable

### Returns

0 on success or -1 on failure

---

enum **\_LMP\_DATATYPE\_CONST**

Data type constants for extracting data from atoms, computes and fixes

Must be kept in sync with the equivalent constants in `lammps/constants.py`

*Values:*

enumerator **LAMMPS\_INT**

32-bit integer (array)

enumerator **LAMMPS\_INT\_2D**

two-dimensional 32-bit integer array

---

enumerator **LAMMPS\_DOUBLE**

64-bit double (array)

enumerator **LAMMPS\_DOUBLE\_2D**

two-dimensional 64-bit double array

enumerator **LAMMPS\_INT64**

64-bit integer (array)

enumerator **LAMMPS\_INT64\_2D**

two-dimensional 64-bit integer array

enumerator **LAMMPS\_STRING**

C-String

enum **\_LMP\_STYLE\_CONST**

Style constants for extracting data from computes and fixes.

Must be kept in sync with the equivalent constants in lammps/constants.py

*Values:*

enumerator **LMP\_STYLE\_GLOBAL**

return global data

enumerator **LMP\_STYLE\_ATOM**

return per-atom data

enumerator **LMP\_STYLE\_LOCAL**

return local data

enum **\_LMP\_TYPE\_CONST**

Type and size constants for extracting data from computes and fixes.

Must be kept in sync with the equivalent constants in lammps/constants.py

*Values:*

enumerator **LMP\_TYPE\_SCALAR**

return scalar

enumerator **LMP\_TYPE\_VECTOR**

return vector

enumerator **LMP\_TYPE\_ARRAY**

return array



enumerator **LMP\_SIZE\_VECTOR**

return length of vector

enumerator **LMP\_SIZE\_ROWS**

return number of rows

enumerator **LMP\_SIZE\_COLS**

return number of columns

## 12.1.6 Scatter/gather operations

This section has functions which gather per-atom data from one or more processors into a contiguous global list ordered by atom ID. The same list is returned to all calling processors. It also contains functions which scatter per-atom data from a contiguous global list across the processors that own those atom IDs. It also has a `create_atoms()` function which can create new atoms by scattering them appropriately to owning processors in the LAMMPS spatial decomposition.

It documents the following functions:

- `lammgs_gather_atoms()`
- `lammgs_gather_atoms_concat()`
- `lammgs_gather_atoms_subset()`
- `lammgs_scatter_atoms()`
- `lammgs_scatter_atoms_subset()`
- `lammgs_gather_bonds()`
- `lammgs_gather()`
- `lammgs_gather_concat()`
- `lammgs_gather_subset()`
- `lammgs_scatter()`
- `lammgs_scatter_subset()`
- `lammgs_create_atoms()`

---

void **lammgs\_gather\_atoms**(void \*handle, char \*name, int type, int count, void \*data)

---

void **lammgs\_gather\_atoms\_concat**(void \*handle, char \*name, int type, int count, void \*data)

---

void **lammgs\_gather\_atoms\_subset**(void \*handle, char \*name, int type, int count, int ndata, int \*ids, void \*data)

---

void **lammgs\_scatter\_atoms**(void \*handle, char \*name, int type, int count, void \*data)

---

void `lammops_scatter_atoms_subset`(void \*handle, char \*name, int type, int count, int ndata, int \*ids, void \*data)

---

void `lammops_gather_bonds`(void \*handle, void \*data)

Gather type and constituent atom info for all bonds

This function copies the list of all bonds into a buffer provided by the calling code. The buffer will be filled with bond type, bond atom 1, bond atom 2 for each bond. Thus the buffer has to be allocated to the dimension of 3 times the **total** number of bonds times the size of the LAMMPS “tagint” type, which is either 4 or 8 bytes depending on whether they are stored in 32-bit or 64-bit integers, respectively. This size depends on the compile time settings used when compiling the LAMMPS library and can be queried by calling `lammops_extract_setting()` with the keyword “tagint”.

When running in parallel, the data buffer must be allocated on **all** MPI ranks and will be filled with the information for **all** bonds in the system.

New in version 28Jul2021.

Below is a brief C code demonstrating accessing this collected bond information.

```
#include <stdio.h>
#include <stdlib.h>
#include <stdint.h>
#include "library.h"

int main(int argc, char **argv)
{
    int tagintsize;
    int64_t i, nbonds;
    void *handle, *bonds;

    handle = lammops_open_no_mpi(0, NULL, NULL);
    lammops_file(handle, "in.some_input");

    tagintsize = lammops_extract_setting(handle, "tagint");
    if (tagintsize == 4)
        nbonds = *(int32_t *)lammops_extract_global(handle, "nbonds");
    else
        nbonds = *(int64_t *)lammops_extract_global(handle, "nbonds");
    bonds = malloc(nbonds * 3 * tagintsize);

    lammops_gather_bonds(handle, bonds);

    if (lammops_extract_setting(handle, "world_rank") == 0) {
        if (tagintsize == 4) {
            int32_t *bonds_real = (int32_t *)bonds;
            for (i = 0; i < nbonds; ++i) {
                printf("bond % 4ld: type = %d, atoms: % 4d % 4d\n", i,
                    bonds_real[3*i], bonds_real[3*i+1], bonds_real[3*i+2]);
            }
        } else {
            int64_t *bonds_real = (int64_t *)bonds;
            for (i = 0; i < nbonds; ++i) {
```

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```

        printf("bond % 4ld: type = %ld, atoms: % 4ld % 4ld\n",i,
              bonds_real[3*i], bonds_real[3*i+1], bonds_real[3*i+2]);
    }
}

lammps_close(handle);
lammps_mpi_finalize();
free(bonds);
return 0;
}

```

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance
- **data** – pointer to data to copy the result to

---

void **lammps\_gather**(void \*handle, char \*name, int type, int count, void \*data)

---

void **lammps\_gather\_concat**(void \*handle, char \*name, int type, int count, void \*data)

---

void **lammps\_gather\_subset**(void \*handle, char \*name, int type, int count, int ndata, int \*ids, void \*data)

---

void **lammps\_scatter**(void \*handle, char \*name, int type, int count, void \*data)

---

void **lammps\_scatter\_subset**(void \*handle, char \*name, int type, int count, int ndata, int \*ids, void \*data)

---

int **lammps\_create\_atoms**(void \*handle, int n, const int \*id, const int \*type, const double \*x, const double \*v, const int \*image, int bexpand)

Create N atoms from list of coordinates

The prototype for this function when compiling with `-DLAMMPS_BIGBIG` is:

```

int lammps_create_atoms(void *handle, int n, int64_t *id, int *type, double *x,
↳double *v, int64_t *image, int bexpand);

```

This function creates additional atoms from a given list of coordinates and a list of atom types. Additionally the atom-IDs, velocities, and image flags may be provided. If atom-IDs are not provided, they will be automatically created as a sequence following the largest existing atom-ID.

This function is useful to add atoms to a simulation or - in tandem with `lammps_reset_box()` - to restore a previously extracted and saved state of a simulation. Additional properties for the new atoms can then be assigned via the `lammps_scatter_atoms()` `lammps_extract_atom()` functions.

For non-periodic boundaries, atoms will **not** be created that have coordinates outside the box unless it is a shrink-wrap boundary and the shrinkexceed flag has been set to a non-zero value. For periodic boundaries atoms will be wrapped back into the simulation cell and its image flags adjusted accordingly, unless explicit image flags are provided.

The function returns the number of atoms created or -1 on failure, e.g. when called before a box has been created.

Coordinates and velocities have to be given in a 1d-array in the order X(1),Y(1),Z(1),X(2),Y(2),Z(2),...,X(N),Y(N),Z(N).

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance
- **n** – number of atoms, N, to be added to the system
- **id** – pointer to N atom IDs; NULL will generate IDs
- **type** – pointer to N atom types (required)
- **x** – pointer to 3N doubles with x-,y-,z- positions of the new atoms (required)
- **v** – pointer to 3N doubles with x-,y-,z- velocities of the new atoms (set to 0.0 if NULL)
- **image** – pointer to N imageint sets of image flags, or NULL
- **bexpand** – if 1, atoms outside of shrink-wrap boundaries will still be created and not dropped and the box extended

#### Returns

number of atoms created on success; -1 on failure (no box, no atom IDs, etc.)

### 12.1.7 Neighbor list access

The following functions enable access to neighbor lists generated by LAMMPS or querying of their properties:

- `lammps_find_compute_neighlist()`
- `lammps_find_fix_neighlist()`
- `lammps_find_pair_neighlist()`
- `lammps_neighlist_num_elements()`
- `lammps_neighlist_element_neighbors()`

---

int **lammps\_find\_compute\_neighlist**(void \*handle, const char \*id, int request)

Find index of a neighbor list requested by a compute

The neighbor list request from a compute is identified by the compute ID and the request ID. The request ID is typically 0, but will be

0 in case a compute has multiple neighbor list requests.

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **id** – Identifier of compute instance
- **reqid** – request id to identify neighbor list in case there are multiple requests from the same compute

**Returns**

return neighbor list index if found, otherwise -1

---

int **lammops\_find\_fix\_neighlist**(void \*handle, const char \*id, int request)

Find index of a neighbor list requested by a fix

The neighbor list request from a fix is identified by the fix ID and the request ID. The request ID is typically 0, but will be > 0 in case a fix has multiple neighbor list requests.

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **id** – Identifier of fix instance
- **reqid** – request id to identify neighbor list in case there are multiple requests from the same fix

**Returns**

return neighbor list index if found, otherwise -1

---

int **lammops\_find\_pair\_neighlist**(void \*handle, const char \*style, int exact, int nsub, int request)

Find index of a neighbor list requested by a pair style

This function determines which of the available neighbor lists for pair styles matches the given conditions. It first matches the style name. If exact is 1 the name must match exactly, if exact is 0, a regular expression or sub-string match is done. If the pair style is hybrid or hybrid/overlay the style is matched against the sub styles instead. If a the same pair style is used multiple times as a sub-style, the nsub argument must be > 0 and represents the nth instance of the sub-style (same as for the pair\_coeff command, for example). In that case nsub=0 will not produce a match and this function will return -1.

The final condition to be checked is the request ID (reqid). This will normally be 0, but some pair styles request multiple neighbor lists and set the request ID to a value > 0.

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **style** – String used to search for pair style instance
- **exact** – Flag to control whether style should match exactly or only a regular expression / sub-string match is applied.
- **nsub** – match nsub-th hybrid sub-style instance of the same style
- **reqid** – request id to identify neighbor list in case there are multiple requests from the same pair style instance

**Returns**

return neighbor list index if found, otherwise -1

---

int **lammops\_neighlist\_num\_elements**(void \*handle, int idx)

Return the number of entries in the neighbor list with given index

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance cast to void \*.
- **idx** – neighbor list index

**Returns**

return number of entries in neighbor list, -1 if idx is not a valid index

---

void **lammops\_neighlist\_element\_neighbors**(void \*handle, int idx, int element, int \*iatom, int \*numneigh, int \*\*neighbors)

Return atom local index, number of neighbors, and array of neighbor local atom indices of neighbor list entry

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance cast to void \*.
- **idx** – index of this neighbor list in the list of all neighbor lists
- **element** – index of this neighbor list entry
- **iatom** – [out] local atom index (i.e. in the range [0, nlocal + nghost), -1 if invalid idx or element value
- **numneigh** – [out] number of neighbors of atom iatom or 0
- **neighbors** – [out] pointer to array of neighbor atom local indices or NULL

## 12.1.8 Configuration information

This section documents the following functions:

- *lammops\_version()*
- *lammops\_get\_os\_info()*
- *lammops\_config\_has\_mpi\_support()*
- *lammops\_config\_has\_gzip\_support()*
- *lammops\_config\_has\_png\_support()*
- *lammops\_config\_has\_jpeg\_support()*
- *lammops\_config\_has\_ffmpeg\_support()*
- *lammops\_config\_has\_exceptions()*
- *lammops\_config\_has\_package()*
- *lammops\_config\_package\_count()*
- *lammops\_config\_package\_name()*
- *lammops\_config\_accelerator()*
- *lammops\_has\_gpu\_device()*
- *lammops\_gpu\_device\_info()*
- *lammops\_has\_style()*

- `lammmps_style_count()`
- `lammmps_style_name()`
- `lammmps_has_id()`
- `lammmps_id_count()`
- `lammmps_id_name()`

These library functions can be used to query the LAMMPS library for compile time settings and included packages and styles. This enables programs that use the library interface to determine whether the linked LAMMPS library is compatible with the requirements of the application without crashing during the LAMMPS functions (e.g. due to missing pair styles from packages) or to choose between different options (e.g. whether to use `lj/cut`, `lj/cut/opt`, `lj/cut/omp` or `lj/cut/intel`). Most of the functions can be called directly without first creating a LAMMPS instance. While crashes within LAMMPS may be recovered from by enabling *exceptions*, avoiding them proactively is a safer approach.

Listing 1: Example for using configuration settings functions

```
#include "library.h"
#include <stdio.h>

int main(int argc, char **argv)
{
    void *handle;

    handle = lammmps_open_no_mpi(0, NULL, NULL);
    lammmps_file(handle, "in.missing");
    if (lammmps_has_error(handle)) {
        char errmsg[256];
        int errtype;
        errtype = lammmps_get_last_error_message(handle, errmsg, 256);
        fprintf(stderr, "LAMMPS failed with error: %s\n", errmsg);
        return 1;
    }
    /* write compressed dump file depending on available of options */
    if (lammmps_has_style(handle, "dump", "atom/zstd")) {
        lammmps_command(handle, "dump d1 all atom/zstd 100 dump.zst");
    } else if (lammmps_has_style(handle, "dump", "atom/gz")) {
        lammmps_command(handle, "dump d1 all atom/gz 100 dump.gz");
    } else if (lammmps_config_has_gzip_support()) {
        lammmps_command(handle, "dump d1 all atom 100 dump.gz");
    } else {
        lammmps_command(handle, "dump d1 all atom 100 dump");
    }
    lammmps_close(handle);
    return 0;
}
```

```
int lammmps_version(void *handle)
```

Get numerical representation of the LAMMPS version date.

The `lammops_version()` function returns an integer representing the version of the LAMMPS code in the format YYYYMMDD. This can be used to implement backward compatibility in software using the LAMMPS library interface. The specific format guarantees, that this version number is growing with every new LAMMPS release.

**Parameters**

**handle** – pointer to a previously created LAMMPS instance

**Returns**

an integer representing the version data in the format YYYYMMDD

---

void `lammops_get_os_info`(char \*buffer, int buf\_size)

Get operating system and architecture information

The `lammops_get_os_info()` function can be used to retrieve detailed information about the hosting operating system and compiler/runtime.

A suitable buffer for a C-style string has to be provided and its length. The assembled text will be truncated to not overflow this buffer. The string is typically a few hundred bytes long.

New in version 9Oct2020.

**Parameters**

- **buffer** – string buffer to copy the information to
  - **buf\_size** – size of the provided string buffer
- 

int `lammops_config_has_mpi_support`()

This function is used to query whether LAMMPS was compiled with a real MPI library or in serial. For the real MPI library it reports the size of the MPI communicator in bytes (4 or 8), which allows to check for compatibility with a hosting code.

**Returns**

0 when compiled with MPI STUBS, otherwise the MPI\_Comm size in bytes

---

int `lammops_config_has_gzip_support`()

Check if the LAMMPS library supports reading or writing compressed files via a pipe to gzip or similar compression programs

Several LAMMPS commands (e.g. *read\_data command*, *write\_data command*, *dump styles atom, custom, and xyz*) support reading and writing compressed files via creating a pipe to the `gzip` program. This function checks whether this feature was *enabled at compile time*. It does **not** check whether `gzip` or any other supported compression programs themselves are installed and usable.

**Returns**

1 if yes, otherwise 0

---



int `lammops_config_has_png_support()`

Check if the LAMMPS library supports writing PNG format images

The LAMMPS *dump style image* supports writing multiple image file formats. Most of them need, however, support from an external library and using that has to be *enabled at compile time*. This function checks whether support for the *PNG image file format* is available in the current LAMMPS library.

**Returns**

1 if yes, otherwise 0

---

int `lammops_config_has_jpeg_support()`

Check if the LAMMPS library supports writing JPEG format images

The LAMMPS *dump style image* supports writing multiple image file formats. Most of them need, however, support from an external library and using that has to be *enabled at compile time*. This function checks whether support for the *JPEG image file format* is available in the current LAMMPS library.

**Returns**

1 if yes, otherwise 0

---

int `lammops_config_has_ffmpeg_support()`

Check if the LAMMPS library supports creating movie files via a pipe to ffmpeg

The LAMMPS *dump style movie* supports generating movies from images on-the-fly via creating a pipe to the *ffmpeg* program. This function checks whether this feature was *enabled at compile time*. It does **not** check whether the *ffmpeg* itself is installed and usable.

**Returns**

1 if yes, otherwise 0

---

int `lammops_config_has_exceptions()`

Check whether LAMMPS errors will throw a C++ exception

In case of errors LAMMPS will either abort or throw a C++ exception. The latter has to be *enabled at compile time*. This function checks if exceptions were enabled.

When using the library interface and C++ exceptions are enabled, the library interface functions will “catch” them and the error status can then be checked by calling *lammops\_has\_error()* and the most recent error message can be retrieved via *lammops\_get\_last\_error\_message()*. This can allow to restart a calculation or delete and recreate the LAMMPS instance when C++ exceptions are enabled. One application of using exceptions this way is the *LAMMPS shell*. If C++ exceptions are disabled and an error happens during a call to LAMMPS, the application will terminate.

**Returns**

1 if yes, otherwise 0

---

int **lammers\_config\_has\_package**(const char\*)

Check if a specific package has been included in LAMMPS

This function checks if the LAMMPS library in use includes the specific *LAMMPS package* provided as argument.

**Parameters**

**name** – string with the name of the package

**Returns**

1 if included, 0 if not.

---

int **lammers\_config\_package\_count**()

Count the number of installed packages in the LAMMPS library.

This function counts how many *LAMMPS packages* are included in the LAMMPS library in use.

**Returns**

number of packages included

---

int **lammers\_config\_package\_name**(int, char\*, int)

Get the name of a package in the list of installed packages in the LAMMPS library.

This function copies the name of the package with the index *idx* into the provided C-style string buffer. The length of the buffer must be provided as *buf\_size* argument. If the name of the package exceeds the length of the buffer, it will be truncated accordingly. If the index is out of range, the function returns 0 and *buffer* is set to an empty string, otherwise 1;

**Parameters**

- **idx** – index of the package in the list of included packages ( $0 \leq \text{idx} < \text{package count}$ )
- **buffer** – string buffer to copy the name of the package to
- **buf\_size** – size of the provided string buffer

**Returns**

1 if successful, otherwise 0

---

int **lammers\_config\_accelerator**(const char\*, const char\*, const char\*)

Check for compile time settings in accelerator packages included in LAMMPS.

This function checks availability of compile time settings of included *accelerator packages* in LAMMPS. Supported packages names are “GPU”, “KOKKOS”, “INTEL”, and “OPENMP”. Supported categories are “api” with possible settings “cuda”, “hip”, “phi”, “pthreads”, “opencl”, “openmp”, and “serial”, and “precision” with possible settings “double”, “mixed”, and “single”. If the combination of package, category, and setting is available, the function returns 1, otherwise 0.

**Parameters**

- **package** – string with the name of the accelerator package
- **category** – string with the category name of the setting
- **setting** – string with the name of the specific setting

**Returns**

1 if available, 0 if not.

---

int **lammops\_has\_gpu\_device()**

Check for presence of a viable GPU package device

The *lammops\_has\_gpu\_device()* function checks at runtime if an accelerator device is present that can be used with the *GPU package*. If at least one suitable device is present the function will return 1, otherwise 0.

More detailed information about the available device or devices can be obtained by calling the *lammops\_get\_gpu\_device\_info()* function.

New in version 14May2021.

#### Returns

1 if viable device is available, 0 if not.

---

void **lammops\_get\_gpu\_device\_info**(char \*buffer, int buf\_size)

Get GPU package device information

The *lammops\_get\_gpu\_device\_info()* function can be used to retrieve detailed information about any accelerator devices that are viable for use with the *GPU package*. It will produce a string that is equivalent to the output of the *nvc\_get\_device* or *ocl\_get\_device* or *hip\_get\_device* tools that are compiled alongside LAMMPS if the GPU package is enabled.

A suitable buffer for a C-style string has to be provided and its length. The assembled text will be truncated to not overflow this buffer. This string can be several kilobytes long, if multiple devices are present.

New in version 14May2021.

#### Parameters

- **buffer** – string buffer to copy the information to
  - **buf\_size** – size of the provided string buffer
- 

int **lammops\_has\_style**(void\*, const char\*, const char\*)

Check if a specific style has been included in LAMMPS

This function checks if the LAMMPS library in use includes the specific *style* of a specific *category* provided as an argument. Valid categories are: *atom*, *integrate*, *minimize*, *pair*, *bond*, *angle*, *dihedral*, *improper*, *kpace*, *compute*, *fix*, *region*, *dump*, and *command*.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **category** – category of the style
- **name** – name of the style

#### Returns

1 if included, 0 if not.

---

int **lammops\_style\_count**(void\*, const char\*)

Count the number of styles of category in the LAMMPS library.

This function counts how many styles in the provided *category* are included in the LAMMPS library in use. Please see *lammops\_has\_style()* for a list of valid categories.

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **category** – category of styles

**Returns**

number of styles in category

---

int **lammops\_style\_name**(void\*, const char\*, int, char\*, int)

Look up the name of a style by index in the list of style of a given category in the LAMMPS library.

This function copies the name of the *category* style with the index *idx* into the provided C-style string buffer. The length of the buffer must be provided as *buf\_size* argument. If the name of the style exceeds the length of the buffer, it will be truncated accordingly. If the index is out of range, the function returns 0 and *buffer* is set to an empty string, otherwise 1.

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **category** – category of styles
- **idx** – index of the style in the list of *category* styles ( $0 \leq \text{idx} < \text{style count}$ )
- **buffer** – string buffer to copy the name of the style to
- **buf\_size** – size of the provided string buffer

**Returns**

1 if successful, otherwise 0

---

int **lammops\_has\_id**(void\*, const char\*, const char\*)

Check if a specific ID exists in the current LAMMPS instance

This function checks if the current LAMMPS instance a *category* ID of the given *name* exists. Valid categories are: *compute*, *dump*, *fix*, *group*, *molecule*, *region*, and *variable*.

New in version 9Oct2020.

**Parameters**

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **category** – category of the id
- **name** – name of the id

**Returns**

1 if included, 0 if not.

---

int **lammops\_id\_count**(void\*, const char\*)

Count the number of IDs of a category.

This function counts how many IDs in the provided *category* are defined in the current LAMMPS instance. Please see *lammops\_has\_id()* for a list of valid categories.

New in version 9Oct2020.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **category** – category of IDs

#### Returns

number of IDs in category

---

int **lammops\_id\_name**(void\*, const char\*, int, char\*, int)

Look up the name of an ID by index in the list of IDs of a given category.

This function copies the name of the *category* ID with the index *idx* into the provided C-style string buffer. The length of the buffer must be provided as *buf\_size* argument. If the name of the style exceeds the length of the buffer, it will be truncated accordingly. If the index is out of range, the function returns 0 and *buffer* is set to an empty string, otherwise 1.

New in version 9Oct2020.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **category** – category of IDs
- **idx** – index of the ID in the list of *category* styles (0 <= idx < count)
- **buffer** – string buffer to copy the name of the style to
- **buf\_size** – size of the provided string buffer

#### Returns

1 if successful, otherwise 0

## 12.1.9 Utility functions

To simplify some tasks, the library interface contains these utility functions. They do not directly call the LAMMPS library.

- *lammops\_encode\_image\_flags()*
- *lammops\_decode\_image\_flags()*
- *lammops\_set\_fix\_external\_callback()*
- *lammops\_fix\_external\_set\_energy\_global()*
- *lammops\_fix\_external\_set\_energy\_peratom()*

- `lammps_fix_external_set_virial_global()`
- `lammps_fix_external_set_virial_peratom()`
- `lammps_fix_external_set_vector_length()`
- `lammps_fix_external_set_vector()`
- `lammps_flush_buffers()`
- `lammps_free()`
- `lammps_is_running()`
- `lammps_force_timeout()`
- `lammps_has_error()`
- `lammps_get_last_error_message()`

The `lammps_free()` function is a clean-up function to free memory that the library had allocated previously via other function calls. Look for notes in the descriptions of the individual commands where such memory buffers were allocated that require the use of `lammps_free()`.

---

int **lammps\_encode\_image\_flags**(int ix, int iy, int iz)

Encode three integer image flags into a single imageint.

The prototype for this function when compiling with `-DLAMMPS_BIGBIG` is:

```
int64_t lammps_encode_image_flags(int ix, int iy, int iz);
```

This function performs the bit-shift, addition, and bit-wise OR operations necessary to combine the values of three integers representing the image flags in x-, y-, and z-direction. Unless LAMMPS is compiled with `-DLAMMPS_BIGBIG`, those integers are limited 10-bit signed integers [-512, 511]. Otherwise the return type changes from `int` to `int64_t` and the valid range for the individual image flags becomes [-1048576,1048575], i.e. that of a 21-bit signed integer. There is no check on whether the arguments conform to these requirements.

#### Parameters

- **ix** – image flag value in x
- **iy** – image flag value in y
- **iz** – image flag value in z

#### Returns

encoded image flag integer

---

void **lammps\_decode\_image\_flags**(int image, int \*flags)

Decode a single image flag integer into three regular integers

The prototype for this function when compiling with `-DLAMMPS_BIGBIG` is:

```
void lammps_decode_image_flags(int64_t image, int *flags);
```

This function does the reverse operation of `lammps_encode_image_flags()` and takes an image flag integer does the bit-shift and bit-masking operations to decode it and stores the resulting three regular integers into the buffer pointed to by `flags`.

#### Parameters

- **image** – encoded image flag integer
- **flags** – pointer to storage where the decoded image flags are stored.

---

```
void lammps_set_fix_external_callback(void *handle, const char *id, FixExternalFnPtr funcptr, void *ptr)
```

Set up the callback function for a fix external instance with the given ID.

Fix *external* allows programs that are running LAMMPS through its library interface to modify certain LAMMPS properties on specific timesteps, similar to the way other fixes do.

This function sets the callback function for use with the “pf/callback” mode. The function has to have C language bindings with the prototype:

```
void func(void *ptr, bigint timestep, int nlocal, tagint *ids, double **x, double_
↪ **fexternal);
```

The argument *ptr* to this function will be stored in fix external and the passed as the first argument calling the callback function *func()*. This would usually be a pointer to the active LAMMPS instance, i.e. the same pointer as the *handle* argument. This would be needed to call functions that set the global or per-atom energy or virial contributions from within the callback function.

The callback mechanism is one of the two modes of how forces and can be applied to a simulation with the help of fix external. The alternative is the array mode where you call `lammps_fix_external_get_force()`.

Please see the documentation for *fix external* for more information about how to use the fix and how to couple it with an external code.

Changed in version 28Jul2021.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **id** – fix ID of fix external instance
- **funcptr** – pointer to callback function
- **ptr** – pointer to object in calling code, passed to callback function as first argument

---

```
void lammps_fix_external_set_energy_global(void *handle, const char *id, double eng)
```

Set the global energy contribution for a fix external instance with the given ID.

This is a companion function to `lammps_set_fix_external_callback()` and `lammps_fix_external_get_force()` to also set the contribution to the global energy from the external code. The value of the *eng* argument will be stored in the fix and applied on the current and all following timesteps until changed by another call to this function. The energy is in energy units as determined by the current *units* settings and is the **total** energy of the contribution. Thus when running in parallel all MPI

processes have to call this function with the **same** value and this will be returned as scalar property of the fix external instance when accessed in LAMMPS input commands or from variables.

Please see the documentation for *fix external* for more information about how to use the fix and how to couple it with an external code.

New in version 28Jul2021.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*.
- **id** – fix ID of fix external instance
- **eng** – total energy to be added to the global energy

---

void **lammops\_fix\_external\_set\_energy\_peratom**(void \*handle, const char \*id, double \*eng)

Set the per-atom energy contribution for a fix external instance with the given ID.

This is a companion function to *lammops\_set\_fix\_external\_callback()* to set the per-atom energy contribution due to the fix from the external code as part of the callback function. For this to work, the handle to the LAMMPS object must be passed as the *ptr* argument when registering the callback function.

Please see the documentation for *fix external* for more information about how to use the fix and how to couple it with an external code.

New in version 28Jul2021.

---

**Note:** This function is fully independent from *lammops\_fix\_external\_set\_energy\_global()* and will **NOT** add any contributions to the global energy tally and **NOT** check whether the sum of the contributions added here are consistent with the global added energy.

---

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*.
- **id** – fix ID of fix external instance
- **eng** – pointer to array of length nlocal with the energy to be added to the per-atom energy

---

void **lammops\_fix\_external\_set\_virial\_global**(void \*handle, const char \*id, double \*virial)

Set the global virial contribution for a fix external instance with the given ID.

This is a companion function to *lammops\_set\_fix\_external\_callback()* and *lammops\_fix\_external\_get\_force()* to set the contribution to the global virial from the external code.

The 6 values of the *virial* array will be stored in the fix and applied on the current and all following timesteps until changed by another call to this function. The components of the virial need to be stored in the order: *xx*, *yy*, *zz*, *xy*, *xz*, *yz*. In LAMMPS the virial is stored internally as *stress\*volume* in units of *pressure\*volume* as determined by the current *units* settings and is the **total** contribution. Thus when running in parallel all MPI processes have to call this function with the **same** value and this will then be added by fix external.



Please see the documentation for *fix external* for more information about how to use the fix and how to couple it with an external code.

New in version 28Jul2021.

### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to `void *`.
- **id** – fix ID of fix external instance
- **virial** – the 6 global stress tensor components to be added to the global virial

---

```
void lammps_fix_external_set_virial_peratom(void *handle, const char *id, double **virial)
```

Set the per-atom virial contribution for a fix external instance with the given ID.

This is a companion function to `lammps_set_fix_external_callback()` to set the per-atom virial contribution due to the fix from the external code as part of the callback function. For this to work, the handle to the LAMMPS object must be passed as the *ptr* argument when registering the callback function.

The order and units of the per-atom stress tensor elements are the same as for the global virial. The code in fix external assumes the dimensions of the per-atom virial array is `double virial[nlocal][6]`.

Please see the documentation for *fix external* for more information about how to use the fix and how to couple it with an external code.

New in version 28Jul2021.

---

**Note:** This function is fully independent from `lammps_fix_external_set_virial_global()` and will **NOT** add any contributions to the global virial tally and **NOT** check whether the sum of the contributions added here are consistent with the global added virial.

---

### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to `void *`.
- **id** – fix ID of fix external instance
- **virial** – a list of `nlocal` entries with the 6 per-atom stress tensor components to be added to the per-atom virial

---

```
void lammps_fix_external_set_vector_length(void *handle, const char *id, int len)
```

Set the vector length for a global vector stored with fix external for analysis

This is a companion function to `lammps_set_fix_external_callback()` and `lammps_fix_external_get_force()` to set the length of a global vector of properties that will be stored with the fix via `lammps_fix_external_set_vector()`.

This function needs to be called **before** a call to `lammps_fix_external_set_vector()` and **before** a run or minimize command. When running in parallel it must be called from **all** MPI processes and with the same length parameter.

Please see the documentation for *fix external* for more information about how to use the fix and how to couple it with an external code.

New in version 28Jul2021.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*.
- **id** – fix ID of fix external instance
- **len** – length of the global vector to be stored with the fix

---

void **lammops\_fix\_external\_set\_vector**(void \*handle, const char \*id, int idx, double val)

Store a global vector value for a fix external instance with the given ID.

This is a companion function to *lammops\_set\_fix\_external\_callback()* and *lammops\_fix\_external\_get\_force()* to set the values of a global vector of properties that will be stored with the fix. And can be accessed from within LAMMPS input commands (e.g. fix ave/time or variables) when used in a vector context.

This function needs to be called **after** a call to *lammops\_fix\_external\_set\_vector\_length()* and the and **before** a run or minimize command. When running in parallel it must be called from **all** MPI processes and with the **same** index and value parameters. The value is assumed to be extensive.

Please see the documentation for *fix external* for more information about how to use the fix and how to couple it with an external code.

New in version 28Jul2021.

---

**Note:** The index in the *idx* parameter is 1-based, i.e. the first element is set with *idx* = 1 and the last element of the vector with *idx* = N, where N is the value of the *len* parameter of the call to *lammops\_fix\_external\_set\_vector\_length()*.

---

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*.
- **id** – fix ID of fix external instance
- **idx** – 1-based index of in global vector
- **val** – value to be stored in global vector

---

void **lammops\_flush\_buffers**(void \*ptr)

Flush output buffers

This function can be used to flush buffered output to be written to screen and logfile pointers to simplify capturing output from LAMMPS library calls.

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*.

void **lammgs\_free**(void \*ptr)

Free memory buffer allocated by LAMMPS.

Some of the LAMMPS C library interface functions return data as pointer to a buffer that has been allocated by LAMMPS or the library interface. This function can be used to delete those in order to avoid memory leaks.

**Parameters**

**ptr** – pointer to data allocated by LAMMPS

---

int **lammgs\_is\_running**(void \*handle)

Check if LAMMPS is currently inside a run or minimization

This function can be used from signal handlers or multi-threaded applications to determine if the LAMMPS instance is currently active.

**Parameters**

**handle** – pointer to a previously created LAMMPS instance cast to void \*.

**Returns**

0 if idle or >0 if active

---

void **lammgs\_force\_timeout**(void \*handle)

Force a timeout to cleanly stop an ongoing run

This function can be used from signal handlers or multi-threaded applications to cleanly terminate an ongoing run.

**Parameters**

**handle** – pointer to a previously created LAMMPS instance cast to void \*

---

int **lammgs\_has\_error**(void \*handle)

Check if there is a (new) error message available

This function can be used to query if an error inside of LAMMPS has thrown a *C++ exception*.

---

**Note:** This function will always report “no error” when the LAMMPS library has been compiled without `-DLAMMPS_EXCEPTIONS` which turns fatal errors aborting LAMMPS into a C++ exceptions. You can use the library function `lammgs_config_has_exceptions()` to check if this is the case.

---

**Parameters**

**handle** – pointer to a previously created LAMMPS instance cast to void \*.

**Returns**

0 on no error, 1 on error.

---

```
int lammops_get_last_error_message(void *handle, char *buffer, int buf_size)
```

Copy the last error message into the provided buffer

This function can be used to retrieve the error message that was set in the event of an error inside of LAMMPS which resulted in a C++ *exception*. A suitable buffer for a C-style string has to be provided and its length. If the internally stored error message is longer, it will be truncated accordingly. The return value of the function corresponds to the kind of error: a “1” indicates an error that occurred on all MPI ranks and is often recoverable, while a “2” indicates an abort that would happen only in a single MPI rank and thus may not be recoverable as other MPI ranks may be waiting on the failing MPI ranks to send messages.

---

**Note:** This function will do nothing when the LAMMPS library has been compiled without `-DLAMMPS_EXCEPTIONS` which turns errors aborting LAMMPS into a C++ exceptions. You can use the library function `lammops_config_has_exceptions()` to check if this is the case.

---

#### Parameters

- **handle** – pointer to a previously created LAMMPS instance cast to void \*
- **buffer** – string buffer to copy the error message to
- **buf\_size** – size of the provided string buffer

#### Returns

1 when all ranks had the error, 2 on a single rank error.

### 12.1.10 Extending the C API

The functionality of the LAMMPS library interface has historically been motivated by the needs of its users. Functions have been added or expanded as they were needed and used. Contributions to the interface are always welcome. However with a refactoring of the library interface and its documentation that started in Spring 2020, there are now a few requirements for including new changes or extensions.

- New functions should be orthogonal to existing ones and not implement functionality that can already be achieved with the existing APIs.
- All changes and additions should be documented with [Doxygen](#) style comments and references to those functions added to the corresponding files in the `doc/src` folder.
- If possible, new unit tests to test those new features should be added.
- New features should also be implemented and documented not just for the C interface, but also the Python and Fortran interfaces.
- All additions should work and be compatible with `-DLAMMPS_BIGBIG`, `-DLAMMPS_SMALLBIG`, `-DLAMMPS_SMALLSMALL` as well as when compiling with and without MPI support.
- The `library.h` file should be kept compatible to C code at a level similar to C89. Its interfaces may not reference any custom data types (e.g. `bigint`, `tagint`, and so on) that are only known inside of LAMMPS; instead `int` and `int64_t` should be used.
- only use C style comments, not C++ style

Please note that these are not **strict** requirements, but the LAMMPS developers very much appreciate, if they are followed and can assist with implementing what is missing. It helps maintaining the code base and keeping it consistent.

---

## 12.2 LAMMPS Python APIs

The LAMMPS Python module enables calling the LAMMPS C library API from Python by dynamically loading functions in the LAMMPS shared library through the Python `ctypes` module. Because of the dynamic loading, it is **required** that LAMMPS is compiled in “*shared*” mode. The Python interface is object oriented, but otherwise tries to be very similar to the C library API. Three different Python classes to run LAMMPS are available and they build on each other. More information on this is in the *Use Python with LAMMPS* section of the manual. Use of the LAMMPS Python module is described in *The lammmps Python module*.

---

## 12.3 LAMMPS Fortran API

The LAMMPS Fortran module is a wrapper around calling functions from the LAMMPS C library API. This is done using the `ISO_C_BINDING` feature in Fortran 2003. The interface is object oriented but otherwise tries to be very similar to the C library API and the basic Python module.

### 12.3.1 The LIBLAMMPS Fortran Module

The LIBLAMMPS module provides an interface to call LAMMPS from a Fortran code. It is based on the LAMMPS C-library interface and requires a Fortran 2003 compatible compiler to be compiled.

While C libraries have a defined binary interface (ABI) and can thus be used from multiple compiler versions from different vendors for as long as they are compatible with the hosting operating system, the same is not true for Fortran codes. Thus the LAMMPS Fortran module needs to be compiled alongside the code using it from the source code in `fortran/lammmps.f90`. When linking, you also need to *link to the LAMMPS library*. A typical command line for a simple program using the Fortran interface would be:

```
mpifort -o testlib.x lammmps.f90 testlib.f90 -L. -llammmps
```

Please note, that the MPI compiler wrapper is only required when the calling the library from an MPI parallel code. Please also note the order of the source files: the `lammmps.f90` file needs to be compiled first, since it provides the LIBLAMMPS module that is imported by the Fortran code using the interface. A working example code can be found together with equivalent examples in C and C++ in the `examples/COUPLE/simple` folder of the LAMMPS distribution.

New in version 9Oct2020.

---

### Work in Progress

This Fortran module is work in progress and only the documented functionality is currently available. The final implementation should cover the entire range of functionality available in the C and Python library interfaces.

---

**Note:** A contributed (and more complete!) Fortran interface that more closely resembles the C-library interface is available in the `examples/COUPLE/fortran2` folder. Please see the README file in that folder for more information about it and how to contact its author and maintainer.

---

---

### 12.3.2 Creating or deleting a LAMMPS object

With the Fortran interface the creation of a *LAMMPS* instance is included in the constructor for creating the `lammps()` derived type. To import the definition of that type and its type bound procedures you need to add a `USE LIBLAMMPS` statement. Internally it will call either `lammps_open_fortran()` or `lammps_open_no_mpi()` from the C library API to create the class instance. All arguments are optional and `lammps_mpi_init()` will be called automatically, if it is needed. Similarly, a possible call to `lammps_finalize()` is integrated into the `close()` function and triggered with the optional logical argument set to `.true.`. Here is a simple example:

```
PROGRAM testlib
  USE LIBLAMMPS           ! include the LAMMPS library interface
  IMPLICIT NONE
  TYPE(lammps)           :: lmp      ! derived type to hold LAMMPS instance
  CHARACTER(len=*) , PARAMETER :: args(3) = &
    [ CHARACTER(len=12) :: 'liblammps', '-log', 'none' ]

  ! create a LAMMPS instance (and initialize MPI)
  lmp = lammps(args)
  ! get and print numerical version code
  PRINT*, 'LAMMPS Version: ', lmp%version()
  ! delete LAMMPS instance (and shuts down MPI)
  CALL lmp%close(.true.)

END PROGRAM testlib
```

It is also possible to pass command line flags from Fortran to C/C++ and thus make the resulting executable behave similar to the standalone executable (it will ignore the `-in/-i` flag, though). This allows to use the command line to configure accelerator and suffix settings, configure screen and logfile output, or to set index style variables from the command line and more. Here is a correspondingly adapted version of the previous example:

```
PROGRAM testlib2
  USE LIBLAMMPS           ! include the LAMMPS library interface
  IMPLICIT NONE
  TYPE(lammps)           :: lmp      ! derived type to hold LAMMPS instance
  CHARACTER(len=128) , ALLOCATABLE :: command_args(:)
  INTEGER :: i, argc

  ! copy command line flags to `command_args()`
  argc = COMMAND_ARGUMENT_COUNT()
  ALLOCATE(command_args(0:argc))
  DO i=0, argc
    CALL GET_COMMAND_ARGUMENT(i, command_args(i))
  END DO

  ! create a LAMMPS instance (and initialize MPI)
  lmp = lammps(command_args)
  ! get and print numerical version code
  PRINT*, 'Program name: ', command_args(0)
  PRINT*, 'LAMMPS Version: ', lmp%version()
  ! delete LAMMPS instance (and shuts down MPI)
  CALL lmp%close(.TRUE.)
  DEALLOCATE(command_args)

END PROGRAM testlib2
```

## Executing LAMMPS commands

Once a LAMMPS instance is created, it is possible to “drive” the LAMMPS simulation by telling LAMMPS to read commands from a file, or pass individual or multiple commands from strings or lists of strings. This is done similar to how it is implemented in the *C-library* `<pg_lib_execute>` interface. Before handing off the calls to the C-library interface, the corresponding Fortran versions of the calls (`file()`, `command()`, `commands_list()`, and `commands_string()`) have to make a copy of the strings passed as arguments so that they can be modified to be compatible with the requirements of strings in C without affecting the original strings. Those copies are automatically deleted after the functions return. Below is a small demonstration of the uses of the different functions:

```
PROGRAM testcmd
  USE LIBLAMMPS
  TYPE(lammps)      :: lmp
  CHARACTER(len=512) :: cmds
  CHARACTER(len=40), ALLOCATABLE :: cmdlist(:)
  CHARACTER(len=10)  :: trimmed
  INTEGER           :: i

  lmp = lammps()
  CALL lmp%file('in.melt')
  CALL lmp%command('variable zpos index 1.0')
  ! define 10 groups of 10 atoms each
  ALLOCATE(cmdlist(10))
  DO i=1, 10
    WRITE(trimmed,'(I10)') 10*i
    WRITE(cmdlist(i),'(A,I1,A,I10,A,A)')      &
      'group g', i-1, ' id ', 10*(i-1)+1, ':', ADJUSTL(trimmed)
  END DO
  CALL lmp%commands_list(cmdlist)
  ! run multiple commands from multi-line string
  cmds = 'clear' // NEW_LINE('A') // &
        'region box block 0 2 0 2 0 2' // NEW_LINE('A') // &
        'create_box 1 box' // NEW_LINE('A') // &
        'create_atoms 1 single 1.0 1.0 ${zpos}'
  CALL lmp%commands_string(cmds)
  CALL lmp%close(.TRUE.)

END PROGRAM testcmd
```

### 12.3.3 The LIBLAMMPS module API

Below are the detailed descriptions of definitions and interfaces of the contents of the LIBLAMMPS Fortran interface to LAMMPS.

#### type `lammps`

Derived type that is the general class of the Fortran interface. It holds a reference to the `LAMMPS` class instance that any of the included calls are forwarded to.

##### Type fields

- `% handle [c_ptr]` :: reference to the LAMMPS class
- `% close` :: `close()`
- `% version` :: `version()`
- `% file` :: `file()`
- `% command` :: `command()`
- `% commands_list` :: `commands_list()`
- `% commands_string` :: `commands_string()`

#### function `lammps`(*args*[, *comm*])

This is the constructor for the Fortran class and will forward the arguments to a call to either `lammps_open_fortran()` or `lammps_open_no_mpi()`. If the LAMMPS library has been compiled with MPI support, it will also initialize MPI, if it has not already been initialized before.

The *args* argument with the list of command line parameters is optional and so is the *comm* argument with the MPI communicator. If *comm* is not provided, `MPI_COMM_WORLD` is assumed. For more details please see the documentation of `lammps_open()`.

##### Parameters

`args (*) [character(len=*),optional]` :: arguments as list of strings

##### Options

`comm [integer,optional]` :: MPI communicator

##### Return

`lammps` :: an instance of the `lammps` derived type

#### subroutine `close`([*finalize*])

This method will close down the LAMMPS instance through calling `lammps_close()`. If the *finalize* argument is present and has a value of `.true.`, then this subroutine also calls `lammps_mpi_finalize()`.

##### Options

`finalize [logical,optional]` :: shut down the MPI environment of the LAMMPS library if true.

#### function `version`()

This method returns the numeric LAMMPS version like `lammps_version()`

##### Return

`integer` :: LAMMPS version

---

#### subroutine `file`(*filename*)

This method will call `lammps_file()` to have LAMMPS read and process commands from a file.

##### Parameters

`filename [character(len=*)]` :: name of file with LAMMPS commands



**subroutine `command(cmd)`**

This method will call `lammers_command()` to have LAMMPS execute a single command.

**Parameters**

`cmd` [*character(len=\*)*] :: single LAMMPS command

**subroutine `commands_list(cmds)`**

This method will call `lammers_commands_list()` to have LAMMPS execute a list of input lines.

**Parameters**

`cmd` (\*) [*character(len=\*)*] :: list of LAMMPS input lines

**subroutine `commands_string(str)`**

This method will call `lammers_commands_string()` to have LAMMPS execute a block of commands from a string.

**Parameters**

`str` [*character(len=\*)*] :: LAMMPS input in string

## 12.4 LAMMPS C++ API

It is also possible to invoke the LAMMPS C++ API directly in your code. It lacks some of the convenience of the C library API, but it allows more direct access to simulation data and thus more low-level manipulations. The following links provide some examples and references to the C++ API.

### 12.4.1 Using the C++ API directly

Using the C++ classes of the LAMMPS library is lacking some of the convenience of the C library API, but it allows a more direct access to simulation data and thus more low-level manipulations and tighter integration of LAMMPS into another code. While for the complete C library API is provided in the `library.h` header file, for using the C++ API it is required to include the individual header files defining the individual classes in use. Typically the name of the class and the name of the header follow some simple rule. Examples are given below.

### 12.4.2 Creating or deleting a LAMMPS object

When using the LAMMPS library interfaces, the core task is to create an instance of the `LAMMPS_NS::LAMMPS` class. In C++ this can be done directly through the `new` operator. All further operations are then initiated through calling member functions of some of the components of the LAMMPS class or accessing their data members. The destruction of the LAMMPS instance is correspondingly initiated by using the `delete` operator. Here is a simple example:

```
#include "lammps.h"

#include <mpi.h>
#include <iostream>

int main(int argc, char **argv)
{
    LAMMPS_NS::LAMMPS *lmp;
    // custom argument vector for LAMMPS library
    const char *lmpargv[] {"liblammps", "-log", "none"};
```

(continues on next page)

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```

int lmpargc = sizeof(lmpargv)/sizeof(const char *);

// explicitly initialize MPI
MPI_Init(&argc, &argv);

// create LAMMPS instance
lmp = new LAMMPS_NS::LAMMPS(lmpargc, (char **)lmpargv, MPI_COMM_WORLD);
// output numerical version string
std::cout << "LAMMPS version ID: " << lmp->num_ver << std::endl;
// delete LAMMPS instance
delete lmp;

// stop MPI environment
MPI_Finalize();
return 0;
}

```

This minimal example only requires to include the `lammps.h` header file since it only accesses a non-pointer member of the LAMMPS class.

### 12.4.3 Executing LAMMPS commands

Once a LAMMPS instance is created by your C++ code, you need to set up a simulation and that is most conveniently done by “driving” it through issuing commands like you would do when running a LAMMPS simulation from an input script. Processing of input in LAMMPS is handled by the `Input` class an instance of which is a member of the `LAMMPS` class. You have two options: reading commands from a file, or executing a single command from a string. See below for a small example:

```

#include "lammps.h"
#include "input.h"
#include <mpi.h>

using namespace LAMMPS_NS;

int main(int argc, char **argv)
{
    const char *lmpargv[] {"liblammps", "-log", "none"};
    int lmpargc = sizeof(lmpargv)/sizeof(const char *);

    MPI_Init(&argc, &argv);
    LAMMPS *lmp = new LAMMPS(lmpargc, (char **)lmpargv, MPI_COMM_WORLD);
    lmp->input->file("in.melt");
    lmp->input->one("run 100 post no");
    delete lmp;
    return 0;
}

```

## USE PYTHON WITH LAMMPS

These pages describe various ways that LAMMPS and Python can be used together.

### 13.1 Overview

The LAMMPS distribution includes a `python` directory with the Python code needed to run LAMMPS from Python. The `python/lammps` package contains *the “lammps” Python module* that wraps the LAMMPS C-library interface. This module makes it possible to do the following either from a Python script, or interactively from a Python prompt:

- create one or more instances of LAMMPS
- invoke LAMMPS commands or read them from an input script
- run LAMMPS incrementally
- extract LAMMPS results
- and modify internal LAMMPS data structures.

From a Python script you can do this in serial or in parallel. Running Python interactively in parallel does not generally work, unless you have a version of Python that extends Python to enable multiple instances of Python to read what you type.

To do all of this, you must build LAMMPS in *“shared” mode* and make certain that your Python interpreter can find the `lammps` Python package and the LAMMPS shared library file.

The Python wrapper for LAMMPS uses the `ctypes` package in Python, which auto-generates the interface code needed between Python and a set of C-style library functions. `Ctypes` has been part of the standard Python distribution since version 2.5. You can check which version of Python you have by simply typing “python” at a shell prompt. Below is an example output for Python version 3.8.5.

```
$ python
Python 3.8.5 (default, Aug 12 2020, 00:00:00)
[GCC 10.2.1 20200723 (Red Hat 10.2.1-1)] on linux
Type "help", "copyright", "credits" or "license" for more information.
>>>
```

**Warning:** The options described in this section of the manual for using Python with LAMMPS currently support either Python 2 or 3. Specifically version 2.7 or later and 3.6 or later. Since the Python community no longer maintains Python 2 (see [this notice](#)), we recommend use of Python 3 with LAMMPS. While Python 2 code should continue to work, that is not something we can guarantee long-term.

LAMMPS can work together with Python in three ways. First, Python can wrap LAMMPS through the its *library interface*, so that a Python script can create one or more instances of LAMMPS and launch one or more simulations. In Python terms, this is referred to as “extending” Python with a LAMMPS module.

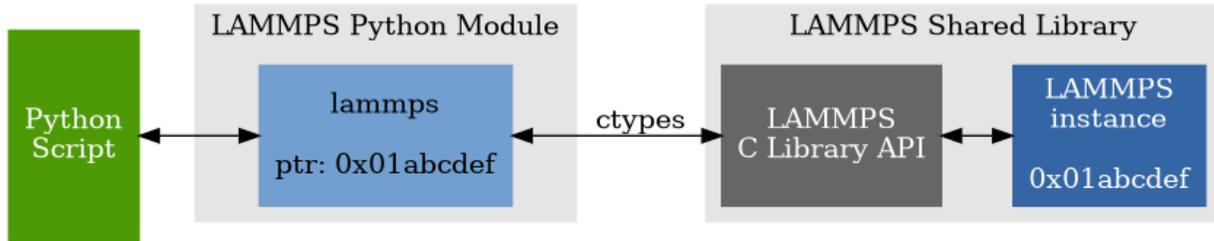


Fig. 1: Launching LAMMPS via Python

Second, the lower-level Python interface in the *lammmps Python class* can be used indirectly through the provided *PyLammmps* and *IPyLammmps* wrapper classes, also written in Python. These wrappers try to simplify the usage of LAMMPS in Python by providing a more object-based interface to common LAMMPS functionality. They also reduce the amount of code necessary to parameterize LAMMPS scripts through Python and make variables and computes directly accessible.

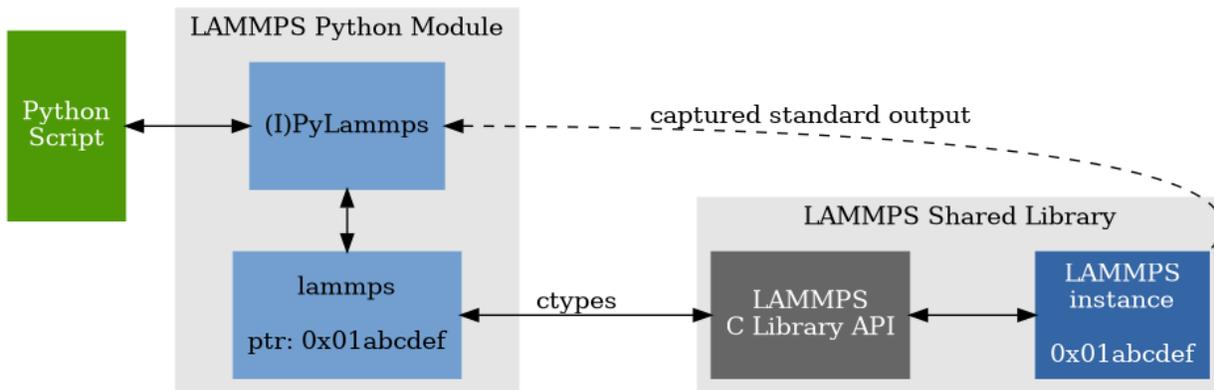


Fig. 2: Using the PyLammmps / IPyLammmps wrappers

Third, LAMMPS can use the Python interpreter, so that a LAMMPS input script or styles can invoke Python code directly, and pass information back-and-forth between the input script and Python functions you write. This Python code can also call back to LAMMPS to query or change its attributes through the LAMMPS Python module mentioned above. In Python terms, this is called “embedding” Python into LAMMPS. When used in this mode, Python can perform script operations that the simple LAMMPS input script syntax can not.

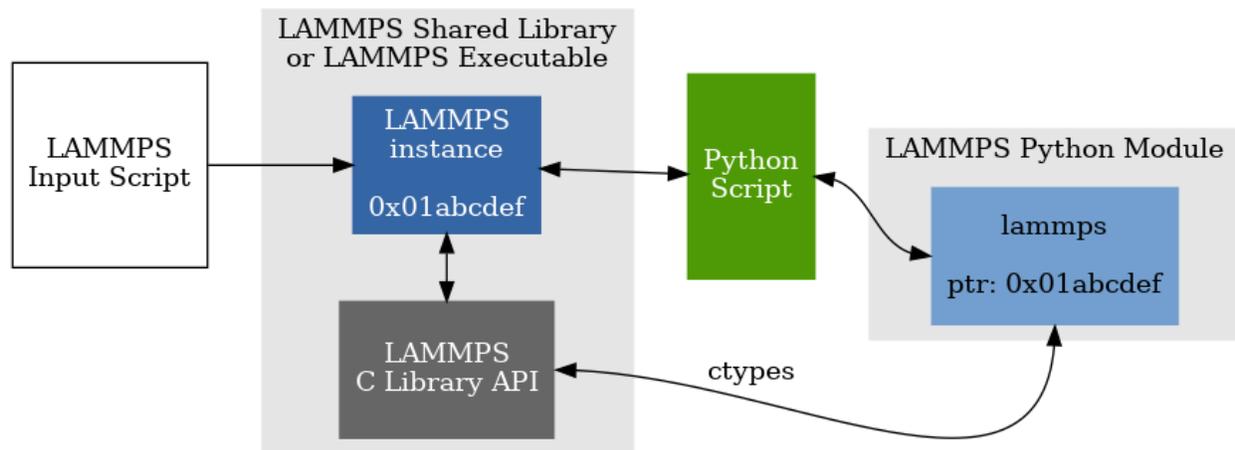


Fig. 3: Calling Python code from LAMMPS

## 13.2 Installation

The LAMMPS Python module enables calling the *LAMMPS C library API* from Python by dynamically loading functions in the LAMMPS shared library through the Python `ctypes` module. Because of the dynamic loading, it is required that LAMMPS is compiled in “*shared*” mode. It is also recommended to compile LAMMPS with *C++ exceptions* enabled.

Two components are necessary for Python to be able to invoke LAMMPS code:

- The LAMMPS Python Package (`lammmps`) from the `python` folder
- The LAMMPS Shared Library (`liblammmps.so`, `liblammmps.dylib` or `liblammmps.dll`) from the folder where you compiled LAMMPS.

### 13.2.1 Installing the LAMMPS Python Module and Shared Library

Making LAMMPS usable within Python and vice versa requires putting the LAMMPS Python package (`lammmps`) into a location where the Python interpreter can find it and installing the LAMMPS shared library into a folder that the dynamic loader searches or inside of the installed `lammmps` package folder. There are multiple ways to achieve this.

1. Do a full LAMMPS installation of libraries, executables, selected headers, documentation (if enabled), and supporting files (only available via CMake), which can also be either system-wide or into user specific folders.
2. Install both components into a Python `site-packages` folder, either system-wide or in the corresponding user-specific folder. This way no additional environment variables need to be set, but the shared library is otherwise not accessible.
3. Do an installation into a virtual environment. This can either be an installation of the Python package only or a full installation of LAMMPS.
4. Leave the files where they are in the source/development tree and adjust some environment variables.

#### Full install (CMake-only)

*Build the LAMMPS executable and library* with `-DBUILD_SHARED_LIBS=on`, `-DLAMMPS_EXCEPTIONS=on` and `-DPKG_PYTHON=on` (The first option is required, the other two are optional by recommended). The exact file name of the shared library depends on the platform (Unix/Linux, MacOS, Windows) and the build configuration

being used. The installation base folder is already set by default to the `$HOME/.local` directory, but it can be changed to a custom location defined by the `CMAKE_INSTALL_PREFIX` CMake variable. This uses a folder called `build` to store files generated during compilation.

```
# create build folder
mkdir build
cd build

# configure LAMMPS compilation
cmake -C ../cmake/presets/basic.cmake -D BUILD_SHARED_LIBS=on \
      -D LAMMPS_EXCEPTIONS=on -D PKG_PYTHON=on ../cmake

# compile LAMMPS
cmake --build .

# install LAMMPS into $HOME/.local
cmake --install .
```

This leads to an installation to the following locations:

File	Location	Notes
LAMMPS Python package	<ul style="list-style-type: none"> <li><code>\$HOME/.local/lib/pythonX.Y/site-packages/lammps</code> (32bit)</li> <li><code>\$HOME/.local/lib64/pythonX.Y/site-packages/lammps</code> (64bit)</li> </ul>	X.Y depends on the installed Python version
LAMMPS shared library	<ul style="list-style-type: none"> <li><code>\$HOME/.local/lib/</code> (32bit)</li> <li><code>\$HOME/.local/lib64/</code> (64bit)</li> </ul>	Set shared loader environment variable to this path (see below for more info on this)
LAMMPS executable	<ul style="list-style-type: none"> <li><code>\$HOME/.local/bin/</code></li> </ul>	
LAMMPS potential files	<ul style="list-style-type: none"> <li><code>\$HOME/.local/share/lammps/potentials/</code></li> </ul>	Set <code>LAMMPS_POTENTIALS</code> environment variable to this path

For a system-wide installation you need to set `CMAKE_INSTALL_PREFIX` to a system folder like `/usr` (or `/usr/local`); the default is `${HOME}/.local`. The installation step for a system folder installation (**not** the configuration/compilation) needs to be done with superuser privilege, e.g. by using `sudo cmake --install ..`. The installation folders will then be changed to (assuming `/usr` as prefix):

File	Location	Notes
LAMMPS Python package	<ul style="list-style-type: none"> <li>• /usr/lib/pythonX.Y/site-packages/lammps (32bit)</li> <li>• /usr/lib64/pythonX.Y/site-packages/lammps (64bit)</li> </ul>	X.Y depends on the installed Python version
LAMMPS shared library	<ul style="list-style-type: none"> <li>• /usr/lib/ (32bit)</li> <li>• /usr/lib64/ (64bit)</li> </ul>	
LAMMPS executable	<ul style="list-style-type: none"> <li>• /usr/bin/</li> </ul>	
LAMMPS potential files	<ul style="list-style-type: none"> <li>• /usr/share/lammps/potentials/</li> </ul>	

To be able to use the “user” installation you have to ensure that the folder containing the LAMMPS shared library is either included in a path searched by the shared linker (e.g. like /usr/lib64/) or part of the LD\_LIBRARY\_PATH environment variable (or DYLD\_LIBRARY\_PATH on MacOS). Otherwise you will get an error when trying to create a LAMMPS object through the Python module.

```
# Unix/Linux
export LD_LIBRARY_PATH=$HOME/.local/lib:$LD_LIBRARY_PATH

# MacOS
export DYLD_LIBRARY_PATH=$HOME/.local/lib:$DYLD_LIBRARY_PATH
```

If you plan to use the LAMMPS executable (e.g., lmp), you may also need to adjust the PATH environment variable (but many newer Linux distributions already have \$HOME/.local/bin included). Example:

```
export PATH=$HOME/.local/bin:$PATH
```

To make those changes permanent, you can add the commands to your \$HOME/.bashrc file. For a system-wide installation it is not necessary due to files installed in system folders that are loaded automatically when a login shell is started.

### Python package only

Compile LAMMPS with either *CMake* or the *traditional make* procedure in *shared mode*. After compilation has finished type (in the compilation folder):

```
make install-python
```

This will try to build a so-called (binary) ‘wheel’, a compressed binary python package and then install it with the python package manager ‘pip’. Installation will be attempted into a system-wide site-packages folder and if that fails into the corresponding folder in the user’s home directory. For a system-wide installation you would have to gain superuser privilege, e.g. through `sudo`

File	Location	Notes
LAMMPS Python package	<ul style="list-style-type: none"> <li>• \$HOME/.local/lib/pythonX.Y/site-packages/lammps</li> </ul>	X.Y depends on the installed Python version
LAMMPS shared library	<ul style="list-style-type: none"> <li>• \$HOME/.local/lib/pythonX.Y/site-packages/lammps</li> </ul>	X.Y depends on the installed Python version

For a system-wide installation those folders would then become.

File	Location	Notes
LAMMPS Python package	<ul style="list-style-type: none"> <li>• /usr/lib/pythonX.Y/site-packages/lammps</li> </ul>	X.Y depends on the installed Python version
LAMMPS shared library	<ul style="list-style-type: none"> <li>• /usr/lib/pythonX.Y/site-packages/lammps</li> </ul>	X.Y depends on the installed Python version

No environment variables need to be set for those, as those folders are searched by default by Python or the LAMMPS Python package.

Changed in version 24Mar2022.

---

**Note:** If there is an existing installation of the LAMMPS python module, `make install-python` will try to update it. However, that will fail if the older version of the module was installed by LAMMPS versions until 17Feb2022. Those were using the `distutils` package, which does not create a “manifest” that allows a clean uninstall. The `make install-python` command will always produce a `lammps-<version>-<python>-<abi>-<os>-<arch>.whl` file (the ‘wheel’). And this file can be later installed directly with `python -m pip install <wheel file>.whl` without having to type `make install-python` again and repeating the build step, too.

---

For the traditional `make` process you can override the python version to version `x.y` when calling `make` with `PYTHON=pythonX.Y`. For a CMake based compilation this choice has to be made during the CMake configuration step.

If the default settings of `make install-python` are not what you want, you can invoke `install.py` from the python directory manually as

```
$ python install.py -p <python package> -l <shared library> [-n]
```

- The `-p` flag points to the `lammps` Python package folder to be installed,
  - the `-l` flag points to the LAMMPS shared library file to be installed,
  - and the optional `-n` instructs the script to only build a wheel file but not attempt to install it.
- 

## Virtual environment



A virtual environment is a minimal Python installation inside of a folder. It allows isolating and customizing a Python environment that is mostly independent from a user or system installation. For the core Python environment, it uses symbolic links to the system installation and thus it can be set up quickly and will not take up much disk space. This gives you the flexibility to install (newer/different) versions of Python packages that would potentially conflict with already installed system packages. It also does not require any superuser privileges. See [PEP 405: Python Virtual Environments](#) for more information.

To create a virtual environment in the folder `$HOME/myenv`, use the `venv` module as follows.

```
# create virtual environment in folder $HOME/myenv
python3 -m venv $HOME/myenv
```

For Python versions prior 3.3 you can use `virtualenv` command instead of “`python3 -m venv`”. This step has to be done only once.

To activate the virtual environment type:

```
source $HOME/myenv/bin/activate
```

This has to be done every time you log in or open a new terminal window and after you turn off the virtual environment with the `deactivate` command.

When using CMake to build LAMMPS, you need to set `CMAKE_INSTALL_PREFIX` to the value of the `$VIRTUAL_ENV` environment variable during the configuration step. For the traditional make procedure, no additional steps are needed. After compiling LAMMPS you can do a “Python package only” installation with `make install-python` and the LAMMPS Python package and the shared library file are installed into the following locations:

File	Location	Notes
LAMMPS Python Module	<ul style="list-style-type: none"> <li><code>\$VIRTUAL_ENV/lib/pythonX.Y/site-packages/lammps</code></li> </ul>	X.Y depends on the installed Python version
LAMMPS shared library	<ul style="list-style-type: none"> <li><code>\$VIRTUAL_ENV/lib/pythonX.Y/site-packages/lammps</code></li> </ul>	X.Y depends on the installed Python version

If you do a full installation (CMake only) with “`install`”, this leads to the following installation locations:

File	Location	Notes
LAMMPS Python Module	<ul style="list-style-type: none"> <li>• \$VIRTUAL_ENV/lib/pythonX.Y/site-packages/lammps</li> </ul>	X.Y depends on the installed Python version
LAMMPS shared library	<ul style="list-style-type: none"> <li>• \$VIRTUAL_ENV/lib/ (32bit)</li> <li>• \$VIRTUAL_ENV/lib64/ (64bit)</li> </ul>	Set shared loader environment variable to this path (see below for more info on this)
LAMMPS executable	<ul style="list-style-type: none"> <li>• \$VIRTUAL_ENV/bin/</li> </ul>	
LAMMPS potential files	<ul style="list-style-type: none"> <li>• \$VIRTUAL_ENV/share/lammps/potentials/</li> </ul>	Set LAMMPS_POTENTIALS environment variable to this path

In that case you need to modify the \$HOME/myenv/bin/activate script in a similar fashion you need to update your \$HOME/.bashrc file to include the shared library and executable locations in LD\_LIBRARY\_PATH (or DYLD\_LIBRARY\_PATH on MacOS) and PATH, respectively.

For example with:

```
# Unix/Linux
echo 'export LD_LIBRARY_PATH=$VIRTUAL_ENV/lib:$LD_LIBRARY_PATH' >> $HOME/myenv/bin/
↪activate

# MacOS
echo 'export DYLD_LIBRARY_PATH=$VIRTUAL_ENV/lib:$DYLD_LIBRARY_PATH' >> $HOME/myenv/
↪bin/activate
```

### In place usage

You can also *compile LAMMPS* as usual in “*shared*” mode leave the shared library and Python package inside the source/compilation folders. Instead of copying the files where they can be found, you need to set the environment variables PYTHONPATH (for the Python package) and LD\_LIBRARY\_PATH (or DYLD\_LIBRARY\_PATH on MacOS

For Bourne shells (bash, ksh and similar) the commands are:

```
export PYTHONPATH=${PYTHONPATH}:${HOME}/lammps/python
export LD_LIBRARY_PATH=${LD_LIBRARY_PATH}:${HOME}/lammps/src
```

For the C-shells like csh or tcsh the commands are:

```
setenv PYTHONPATH ${PYTHONPATH}:${HOME}/lammps/python
setenv LD_LIBRARY_PATH ${LD_LIBRARY_PATH}:${HOME}/lammps/src
```

On MacOS you may also need to set DYLD\_LIBRARY\_PATH accordingly. You can make those changes permanent by editing your \$HOME/.bashrc or \$HOME/.login files, respectively.

---

**Note:** The PYTHONPATH needs to point to the parent folder that contains the lammps package!

---

To verify if LAMMPS can be successfully started from Python, start the Python interpreter, load the lammps Python module and create a LAMMPS instance. This should not generate an error message and produce output similar to the following:

```
$ python
Python 3.8.5 (default, Sep  5 2020, 10:50:12)
[GCC 10.2.0] on linux
Type "help", "copyright", "credits" or "license" for more information.
>>> import lammps
>>> lmp = lammps.lammps()
LAMMPS (18 Sep 2020)
using 1 OpenMP thread(s) per MPI task
>>>
```

---

**Note:** Unless you opted for “In place use”, you will have to rerun the installation any time you recompile LAMMPS to ensure the latest Python package and shared library are installed and used.

---



---

**Note:** If you want Python to be able to load different versions of the LAMMPS shared library with different settings, you will need to manually copy the files under different names (e.g. liblammps\_mpi.so or liblammps\_gpu.so) into the appropriate folder as indicated above. You can then select the desired library through the *name* argument of the LAMMPS object constructor (see *Creating or deleting a LAMMPS object*).

---

### 13.2.2 Extending Python to run in parallel

If you wish to run LAMMPS in parallel from Python, you need to extend your Python with an interface to MPI. This also allows you to make MPI calls directly from Python in your script, if you desire.

We have tested this with [MPI for Python](#) (aka mpi4py) and you will find installation instruction for it below.

Installation of mpi4py (version 3.0.3 as of Sep 2020) can be done as follows:

- Via pip into a local user folder with:

```
pip install --user mpi4py
```

- Via dnf into a system folder for RedHat/Fedora systems:

```
# for use with OpenMPI
sudo dnf install python3-mpi4py-openmpi
# for use with MPICH
sudo dnf install python3-mpi4py-openmpi
```

- Via pip into a virtual environment (see above):

```
$ source $HOME/myenv/activate
(myenv)$ pip install mpi4py
```

- Via pip into a system folder (not recommended):

```
sudo pip install mpi4py
```

For more detailed installation instructions and additional options, please see the [mpi4py installation](#) page.

To use mpi4py and LAMMPS in parallel from Python, you **must** make certain that **both** are using the **same** implementation and version of MPI library. If you only have one MPI library installed on your system this is not an issue, but it can be if you have multiple MPI installations (e.g. on an HPC cluster to be selected through environment modules). Your LAMMPS build is explicit about which MPI it is using, since it is either detected during CMake configuration or in the traditional make build system you specify the details in your low-level `src/MAKE/Makefile.foo` file. The installation process of mpi4py uses the `mpicc` command to find information about the MPI it uses to build against. And it tries to load “libmpi.so” from the `LD_LIBRARY_PATH`. This may or may not find the MPI library that LAMMPS is using. If you have problems running both mpi4py and LAMMPS together, this is an issue you may need to address, e.g. by loading the module for different MPI installation so that mpi4py finds the right one.

If you have successfully installed mpi4py, you should be able to run Python and type

```
from mpi4py import MPI
```

without error. You should also be able to run Python in parallel on a simple test script

```
$ mpirun -np 4 python3 test.py
```

where `test.py` contains the lines

```
from mpi4py import MPI
comm = MPI.COMM_WORLD
print("Proc %d out of %d procs" % (comm.Get_rank(), comm.Get_size()))
```

and see one line of output for each processor you run on.

```
# NOTE: the line order is not deterministic
$ mpirun -np 4 python3 test.py
Proc 0 out of 4 procs
Proc 1 out of 4 procs
Proc 2 out of 4 procs
Proc 3 out of 4 procs
```

## 13.3 Run LAMMPS from Python

After compiling the LAMMPS shared library and making it ready to use, you can now write and run Python scripts that import the LAMMPS Python module and launch LAMMPS simulations through Python scripts. The following pages take you through the various steps necessary, what functionality is available and give some examples how to use it.

---

### 13.3.1 Running LAMMPS and Python in serial

To run a LAMMPS in serial, type these lines into Python interactively from the bench directory:

```
>>> from lammps import lammps
>>> lmp = lammps()
>>> lmp.file("in.lj")
```

Or put the same lines in the file `test.py` and run it as

```
$ python3 test.py
```

Either way, you should see the results of running the `in.lj` benchmark on a single processor appear on the screen, the same as if you had typed something like:

```
lmp_serial -in in.lj
```

### 13.3.2 Running LAMMPS and Python in parallel with MPI

To run LAMMPS in parallel, assuming you have installed the `mpi4py` package as discussed *Extending Python to run in parallel*, create a `test.py` file containing these lines:

```
from mpi4py import MPI
from lammps import lammps
lmp = lammps()
lmp.file("in.lj")
me = MPI.COMM_WORLD.Get_rank()
nprocs = MPI.COMM_WORLD.Get_size()
print("Proc %d out of %d procs has" % (me,nprocs),lmp)
MPI.Finalize()
```

You can run the script in parallel as:

```
$ mpirun -np 4 python3 test.py
```

and you should see the same output as if you had typed

```
$ mpirun -np 4 lmp_mpi -in in.lj
```

Note that without the `mpi4py` specific lines from `test.py`

```
from lammps import lammps
lmp = lammps()
lmp.file("in.lj")
```

running the script with `mpirun` on  $P$  processors would lead to  $P$  independent simulations to run parallel, each with a single processor. Therefore, if you use the `mpi4py` lines and you see multiple LAMMPS single processor outputs, `mpi4py` is not working correctly.

Also note that once you import the `mpi4py` module, `mpi4py` initializes MPI for you, and you can use MPI calls directly in your Python script, as described in the `mpi4py` documentation. The last line of your Python script should be `MPI.finalize()`, to insure MPI is shut down correctly.

### 13.3.3 Running Python scripts

Note that any Python script (not just for LAMMPS) can be invoked in one of several ways:

```
$ python script.py
$ python -i script.py
$ ./script.py
```

The last command requires that the first line of the script be something like this:

```
#!/usr/bin/python
#!/usr/bin/python -i
```

where the path points to where you have Python installed, and that you have made the script file executable:

```
$ chmod +x script.py
```

Without the `-i` flag, Python will exit when the script finishes. With the `-i` flag, you will be left in the Python interpreter when the script finishes, so you can type subsequent commands. As mentioned above, you can only run Python interactively when running Python on a single processor, not in parallel.

---

### 13.3.4 Creating or deleting a LAMMPS object

With the Python interface the creation of a *LAMMPS* instance is included in the constructors for the *lammps*, *PyLammps*, and *IPyLammps* classes. Internally it will call either *lammps\_open()* or *lammps\_open\_no\_mpi()* from the C library API to create the class instance.

All arguments are optional. The *name* argument allows loading a LAMMPS shared library that is named *liblammps\_machine.so* instead of the default name of *liblammps.so*. In most cases the latter will be installed or used. The *ptr* argument is for use of the *lammps* module from inside a LAMMPS instance, e.g. with the *python* command, where a pointer to the already existing *LAMMPS* class instance can be passed to the Python class and used instead of creating a new instance. The *comm* argument may be used in combination with the *mpi4py* module to pass an MPI communicator to LAMMPS and thus it is possible to run the Python module like the library interface on a subset of the MPI ranks after splitting the communicator.

Here are simple examples using all three Python interfaces:

---

#### lammps API

```
from lammps import lammps

# NOTE: argv[0] is set by the lammps class constructor
args = ["-log", "none"]

# create LAMMPS instance
lmp = lammps(cmdargs=args)

# get and print numerical version code
print("LAMMPS Version: ", lmp.version())

# explicitly close and delete LAMMPS instance (optional)
lmp.close()
```

---

## PyLammps API

The `PyLammps` class is a wrapper around the `lammps` class and all of its lower level functions. By default, it will create a new instance of `lammps` passing along all arguments to the constructor of `lammps`.

```
from lammps import PyLammps

# NOTE: argv[0] is set by the lammps class constructor
args = ["-log", "none"]

# create LAMMPS instance
L = PyLammps(cmdargs=args)

# get and print numerical version code
print("LAMMPS Version: ", L.version())

# explicitly close and delete LAMMPS instance (optional)
L.close()
```

`PyLammps` objects can also be created on top of an existing `lammps` object:

```
from lammps import lammps, PyLammps
...
# create LAMMPS instance
lmp = lammps(cmdargs=args)

# create PyLammps instance using previously created LAMMPS instance
L = PyLammps(ptr=lmp)
```

This is useful if you have to create the `lammps` instance in a specific way, but want to take advantage of the `PyLammps` interface.

---

## IPyLammps API

The `IPyLammps` class is an extension of the `PyLammps` class. It has the same construction behavior. By default, it will create a new instance of `lammps` passing along all arguments to the constructor of `lammps`.

```
from lammps import IPyLammps

# NOTE: argv[0] is set by the lammps class constructor
args = ["-log", "none"]

# create LAMMPS instance
L = IPyLammps(cmdargs=args)

# get and print numerical version code
print("LAMMPS Version: ", L.version())

# explicitly close and delete LAMMPS instance (optional)
L.close()
```

You can also initialize `IPyLammps` on top of an existing `lammps` or `PyLammps` object:

```
from lammps import lammps, IPyLammps
...
# create LAMMPS instance
lmp = lammps(cmdargs=args)

# create PyLammps instance using previously created LAMMPS instance
L = PyLammps(ptr=lmp)
```

This is useful if you have to create the `lammps` instance in a specific way, but want to take advantage of the `IPyLammps` interface.

---

In all of the above cases, same as with the *C library API*, this will use the `MPI_COMM_WORLD` communicator for the MPI library that LAMMPS was compiled with.

The `lmp.close()` call is optional since the LAMMPS class instance will also be deleted automatically during the `lammps` class destructor. Instead of `lmp.close()` it is also possible to call `lmp.finalize()`; this will destruct the LAMMPS instance, but also finalized and release the MPI and/or Kokkos environment if enabled and active.

Note that you can create multiple LAMMPS objects in your Python script, and coordinate and run multiple simulations, e.g.

```
from lammps import lammps
lmp1 = lammps()
lmp2 = lammps()
lmp1.file("in.file1")
lmp2.file("in.file2")
```

### 13.3.5 Executing commands

Once an instance of the `lammps`, `PyLammps`, or `IPyLammps` class is created, there are multiple ways to “feed” it commands. In a way that is not very different from running a LAMMPS input script, except that Python has many more facilities for structured programming than the LAMMPS input script syntax. Furthermore it is possible to “compute” what the next LAMMPS command should be.

---

#### lammps API

Same as in the equivalent *C library functions*, commands can be read from a file, a single string, a list of strings and a block of commands in a single multi-line string. They are processed under the same boundary conditions as the C library counterparts. The example below demonstrates the use of `lammps.file()`, `lammps.command()`, `lammps.commands_list()`, and `lammps.commands_string()`:

```
from lammps import lammps
lmp = lammps()

# read commands from file 'in.melt'
lmp.file('in.melt')

# issue a single command
lmp.command('variable zpos index 1.0')

# create 10 groups with 10 atoms each
cmds = ["group g{} id {}:{}".format(i,10*i+1,10*(i+1)) for i in range(10)]
```

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```

Imp.commands_list(cmds)

# run commands from a multi-line string
block = """
clear
region box block 0 2 0 2 0 2
create_box 1 box
create_atoms 1 single 1.0 1.0 ${zpos}
"""
Imp.commands_string(block)

```

### PyLammps/IPyLammps API

Unlike the lammps API, the PyLammps/IPyLammps APIs allow running LAMMPS commands by calling equivalent member functions of *PyLammps* and *IPyLammps* instances.

For instance, the following LAMMPS command

```
region box block 0 10 0 5 -0.5 0.5
```

can be executed using with the lammps API with the following Python code if `Imp` is an instance of *lammps*:

```

from lammps import lammps

Imp = lammps()
Imp.command("region box block 0 10 0 5 -0.5 0.5")

```

With the PyLammps interface, any LAMMPS command can be split up into arbitrary parts. These parts are then passed to a member function with the name of the *command*. For the *region* command that means the `region()` method can be called. The arguments of the command can be passed as one string, or individually.

```

from lammps import PyLammps

L = PyLammps()

# pass command parameters as one string
L.region("box block 0 10 0 5 -0.5 0.5")

# OR pass them individually
L.region("box block", 0, 10, 0, 5, -0.5, 0.5)

```

In the latter example, all parameters except the first are Python floating-point literals. The member function takes the entire parameter list and transparently merges it to a single command string.

The benefit of this approach is avoiding redundant command calls and easier parameterization. In the lammps API parameterization needed to be done manually by creating formatted command strings.

```

Imp.command("region box block %f %f %f %f %f %f" % (xlo, xhi, ylo, yhi, zlo,
↪zhi))

```

In contrast, methods of PyLammps accept parameters directly and will convert them automatically to a final command string.

```
L.region("box block", xlo, xhi, ylo, yhi, zlo, zhi)
```

Using these facilities, the example shown for the lammps API can be rewritten as follows:

```
from lammps import PyLammps
L = PyLammps()

# read commands from file 'in.melt'
L.file('in.melt')

# issue a single command
L.variable('zpos', 'index', 1.0)

# create 10 groups with 10 atoms each
for i in range(10):
    L.group(f"g{i}", "id", f"{10*i+1}:{10*(i+1)}")

L.clear()
L.region("box block", 0, 2, 0, 2, 0, 2)
L.create_box(1, "box")
L.create_atoms(1, "single", 1.0, 1.0, "${zpos}")
```

### 13.3.6 System properties

Similar to what is described in *System properties*, the instances of *lammps*, *PyLammps*, or *IPyLammps* can be used to extract different kinds of information about the active LAMMPS instance and also to modify some of it. The main difference between the interfaces is how the information is exposed.

While the *lammps* is just a thin layer that wraps C API calls, *PyLammps* and *IPyLammps* expose information as objects and properties.

In some cases the data returned is a direct reference to the original data inside LAMMPS cast to ctypes pointers. Where possible, the wrappers will determine the ctypes data type and cast pointers accordingly. If numpy is installed arrays can also be extracted as numpy arrays, which will access the C arrays directly and have the correct dimensions to protect against invalid accesses.

**Warning:** When accessing per-atom data, please note that this data is the per-processor local data and indexed accordingly. These arrays can change sizes and order at every neighbor list rebuild and atom sort event as atoms are migrating between sub-domains.

---

#### lammps API

```
from lammps import lammps

lmp = lammps()
lmp.file("in.sysinit")

natoms = lmp.get_natoms()
print(f"running simulation with {natoms} atoms")

lmp.command("run 1000 post no");
```

```

for i in range(10):
    lmp.command("run 100 pre no post no")
    pe = lmp.get_thermo("pe")
    ke = lmp.get_thermo("ke")
    print(f"PE = {pe}\nKE = {ke}")

```

```
lmp.close()
```

#### Methods:

- `version()`: return the numerical version id, e.g. LAMMPS 2 Sep 2015 -> 20150902
- `get_thermo()`: return current value of a thermo keyword
- `get_natoms()`: total # of atoms as int
- `reset_box()`: reset the simulation box size
- `extract_setting()`: return a global setting
- `extract_global()`: extract a global quantity
- `extract_box()`: extract box info
- `create_atoms()`: create N atoms with IDs, types, x, v, and image flags

---

#### PyLammps/IPyLammps API

In addition to the functions provided by `lammps`, `PyLammps` objects have several properties which allow you to query the system state:

##### **L.system**

Is a dictionary describing the system such as the bounding box or number of atoms

##### **L.system.xlo, L.system.xhi**

bounding box limits along x-axis

##### **L.system.ylo, L.system.yhi**

bounding box limits along y-axis

##### **L.system.zlo, L.system.zhi**

bounding box limits along z-axis

##### **L.communication**

configuration of communication subsystem, such as the number of threads or processors

##### **L.communication.nthreads**

number of threads used by each LAMMPS process

##### **L.communication.nprocs**

number of MPI processes used by LAMMPS

##### **L.fixes**

List of fixes in the current system

##### **L.computes**

List of active computes in the current system

##### **L.dump**

List of active dumps in the current system

**L.groups**

List of groups present in the current system

**Retrieving the value of an arbitrary LAMMPS expressions**

LAMMPS expressions can be immediately evaluated by using the `eval` method. The passed string parameter can be any expression containing global *thermo command* values, variables, compute or fix data (see *Output from LAMMPS (thermo, dumps, computes, fixes, variables)*):

```
result = L.eval("ke") # kinetic energy
result = L.eval("pe") # potential energy

result = L.eval("v_t/2.0")
```

**Example**

```
from lammps import PyLammps

L = PyLammps()
L.file("in.sysinit")

print(f"running simulation with {L.system.natoms} atoms")

L.run(1000, "post no");

for i in range(10):
    L.run(100, "pre no post no")
    pe = L.eval("pe")
    ke = L.eval("ke")
    print(f"PE = {pe}\nKE = {ke}")
```

### 13.3.7 Per-atom properties

Similar to what is described in *Per-atom properties*, the instances of *lammps*, *PyLammps*, or *IPyLammps* can be used to extract atom quantities and modify some of them. The main difference between the interfaces is how the information is exposed.

While the *lammps* is just a thin layer that wraps C API calls, *PyLammps* and *IPyLammps* expose information as objects and properties.

In some cases the data returned is a direct reference to the original data inside LAMMPS cast to ctypes pointers. Where possible, the wrappers will determine the ctypes data type and cast pointers accordingly. If numpy is installed arrays can also be extracted as numpy arrays, which will access the C arrays directly and have the correct dimensions to protect against invalid accesses.

**Warning:** When accessing per-atom data, please note that this data is the per-processor local data and indexed accordingly. These arrays can change sizes and order at every neighbor list rebuild and atom sort event as atoms are migrating between sub-domains.

**lammps API**

```
from lammps import lammps
```

```

lmp = lammps()
lmp.file("in.sysinit")

nlocal = lmp.extract_global("nlocal")
x = lmp.extract_atom("x")

for i in range(nlocal):
    print("(x,y,z) = (" + x[i][0], x[i][1], x[i][2], ")")

lmp.close()

```

**Methods:**

- `extract_atom()`: extract a per-atom quantity

**Numpy Methods:**

- `numpy.extract_atom()`: extract a per-atom quantity as numpy array

**PyLammps/IPyLammps API**

All atoms in the current simulation can be accessed by using the `PyLammps.atoms` property. Each element of this list is a `Atom` or `Atom2D` object. The attributes of these objects provide access to their data (id, type, position, velocity, force, etc.):

```

# access first atom
L.atoms[0].id
L.atoms[0].type

# access second atom
L.atoms[1].position
L.atoms[1].velocity
L.atoms[1].force

```

Some attributes can be changed:

```

# set position in 2D simulation
L.atoms[0].position = (1.0, 0.0)

# set position in 3D simulation
L.atoms[0].position = (1.0, 0.0, 1.0)

```

### 13.3.8 Compute, fixes, variables

This section documents accessing or modifying data from objects like computes, fixes, or variables in LAMMPS using the `lammps` module.

**lammps API**

For `lammps.extract_compute()` and `lammps.extract_fix()`, the global, per-atom, or local data calculated by the compute or fix can be accessed. What is returned depends on whether the compute or fix calculates a scalar or vector or array. For a scalar, a single double value is returned. If the compute or fix calculates a vector or array, a pointer to the internal LAMMPS data is returned, which you can use via

normal Python subscripting.

The one exception is that for a fix that calculates a global vector or array, a single double value from the vector or array is returned, indexed by I (vector) or I and J (array). I,J are zero-based indices. See the [Howto output](#) page for a discussion of global, per-atom, and local data, and of scalar, vector, and array data types. See the doc pages for individual *computes* and *fixes* for a description of what they calculate and store.

For `lammops.extract_variable()`, an *equal-style or atom-style variable* is evaluated and its result returned.

For equal-style variables a single `c_double` value is returned and the group argument is ignored. For atom-style variables, a vector of `c_double` is returned, one value per atom, which you can use via normal Python subscripting. The values will be zero for atoms not in the specified group.

`lammops.numpy.extract_compute()`, `lammops.numpy.extract_fix()`, and `lammops.numpy.extract_variable()` are equivalent NumPy implementations that return NumPy arrays instead of ctypes pointers.

The `lammops.set_variable()` method sets an existing string-style variable to a new string value, so that subsequent LAMMPS commands can access the variable.

#### Methods:

- `lammops.extract_compute()`: extract value(s) from a compute
- `lammops.extract_fix()`: extract value(s) from a fix
- `lammops.extract_variable()`: extract value(s) from a variable
- `lammops.set_variable()`: set existing named string-style variable to value

#### NumPy Methods:

- `lammops.numpy.extract_compute()`: extract value(s) from a compute, return arrays as numpy arrays
  - `lammops.numpy.extract_fix()`: extract value(s) from a fix, return arrays as numpy arrays
  - `lammops.numpy.extract_variable()`: extract value(s) from a variable, return arrays as numpy arrays
- 

### PyLammps/IPyLammps API

PyLammps and IPyLammps classes currently do not add any additional ways of retrieving information out of computes and fixes. This information can still be accessed by using the lammps API:

```
L.lmp.extract_compute(...)  
L.lmp.extract_fix(...)  
# OR  
L.lmp.numpy.extract_compute(...)  
L.lmp.numpy.extract_fix(...)
```

LAMMPS variables can be both defined and accessed via the `PyLammps` interface.

To define a variable you can use the `variable` command:

```
L.variable("a index 2")
```

A dictionary of all variables is returned by the `PyLammps.variables` property:

you can access an individual variable by retrieving a variable object from the `L.variables` dictionary by name

```
a = L.variables['a']
```

The variable value can then be easily read and written by accessing the `value` property of this object.

```
print(a.value)
a.value = 4
```

### 13.3.9 Scatter/gather operations

```
data = lmp.gather_atoms(name,type,count) # return per-atom property of all atoms.
↳gathered into data, ordered by atom ID
                                     # name = "x", "charge", "type", etc
data = lmp.gather_atoms_concat(name,type,count) # ditto, but concatenated atom values.
↳from each proc (unordered)
data = lmp.gather_atoms_subset(name,type,count,ndata,ids) # ditto, but for subset of.
↳Ndata atoms with IDs

lmp.scatter_atoms(name,type,count,data) # scatter per-atom property to all atoms from.
↳data, ordered by atom ID
                                     # name = "x", "charge", "type", etc
                                     # count = # of per-atom values, 1 or 3, etc

lmp.scatter_atoms_subset(name,type,count,ndata,ids,data) # ditto, but for subset of.
↳Ndata atoms with IDs
```

The gather methods collect peratom info of the requested type (atom coords, atom types, forces, etc) from all processors, and returns the same vector of values to each calling processor. The scatter functions do the inverse. They distribute a vector of peratom values, passed by all calling processors, to individual atoms, which may be owned by different processors.

Note that the data returned by the gather methods, e.g. `gather_atoms("x")`, is different from the data structure returned by `extract_atom("x")` in four ways. (1) `Gather_atoms()` returns a vector which you index as `x[i]`; `extract_atom()` returns an array which you index as `x[i][j]`. (2) `Gather_atoms()` orders the atoms by atom ID while `extract_atom()` does not. (3) `Gather_atoms()` returns a list of all atoms in the simulation; `extract_atoms()` returns just the atoms local to each processor. (4) Finally, the `gather_atoms()` data structure is a copy of the atom coords stored internally in LAMMPS, whereas `extract_atom()` returns an array that effectively points directly to the internal data. This means you can change values inside LAMMPS from Python by assigning a new values to the `extract_atom()` array. To do this with the `gather_atoms()` vector, you need to change values in the vector, then invoke the `scatter_atoms()` method.

For the scatter methods, the array of coordinates passed to must be a ctypes vector of ints or doubles, allocated and initialized something like this:

```
from ctypes import c_double
natoms = lmp.get_natoms()
n3 = 3*natoms
x = (n3*c_double)()
x[0] = x coord of atom with ID 1
x[1] = y coord of atom with ID 1
x[2] = z coord of atom with ID 1
```

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```
x[3] = x coord of atom with ID 2
...
x[n3-1] = z coord of atom with ID natoms
lmp.scatter_atoms("x",1,3,x)
```

Alternatively, you can just change values in the vector returned by the gather methods, since they are also ctypes vectors.

### 13.3.10 Neighbor list access

Access to neighbor lists is handled through a couple of wrapper classes that allows to treat it like either a python list or a NumPy array. The access procedure is similar to that of the C-library interface: use one of the “find” functions to look up the index of the neighbor list in the global table of neighbor lists and then get access to the neighbor list data. The code sample below demonstrates reading the neighbor list data using the NumPy access method.

```
from lammps import lammps
import numpy as np

lmp = lammps()
lmp.commands_string("""
region box block -2 2 -2 2 -2 2
lattice fcc 1.0
create_box 1 box
create_atoms 1 box
mass 1 1.0
pair_style lj/cut 2.5
pair_coeff 1 1 1.0 1.0
run 0 post no""")

# look up the neighbor list
nlidx = lmp.find_pair_neighlist('lj/cut')
nl = lmp.numpy.get_neighlist(nlidx)
tags = lmp.extract_atom('id')
print("half neighbor list with {} entries".format(nl.size))
# print neighbor list contents
for i in range(0,nl.size):
    idx, nlist = nl.get(i)
    print("\natom {} with ID {} has {} neighbors:".format(idx,tags[idx],nlist.size))
    if nlist.size > 0:
        for n in np.nditer(nlist):
            print(" atom {} with ID {}".format(n,tags[n]))
```

#### Methods:

- `lammps.get_neighlist()`: Get neighbor list for given index
- `lammps.get_neighlist_size()`: Get number of elements in neighbor list
- `lammps.get_neighlist_element_neighbors()`: Get element in neighbor list and its neighbors
- `lammps.find_pair_neighlist()`: Find neighbor list of pair style
- `lammps.find_fix_neighlist()`: Find neighbor list of pair style
- `lammps.find_compute_neighlist()`: Find neighbor list of pair style

#### NumPy Methods:



- `lammps.numpy.get_neighlist()`: Get neighbor list for given index, which uses NumPy arrays for its element neighbor arrays
- `lammps.numpy.get_neighlist_element_neighbors()`: Get element in neighbor list and its neighbors (as numpy array)

### 13.3.11 Configuration information

The following methods can be used to query the LAMMPS library about compile time settings and included packages and styles.

Listing 1: Example for using configuration settings functions

```

from lammps import lammps

lmp = lammps()

try:
    lmp.file("in.missing")
except Exception as e:
    print("LAMMPS failed with error:", e)

# write compressed dump file depending on available of options

if lmp.has_style("dump", "atom/zstd"):
    lmp.command("dump d1 all atom/zstd 100 dump.zst")
elif lmp.has_style("dump", "atom/gz"):
    lmp.command("dump d1 all atom/gz 100 dump.gz")
elif lmp.has_gzip_support():
    lmp.command("dump d1 all atom 100 dump.gz")
else:
    lmp.command("dump d1 all atom 100 dump")

```

#### Methods:

- `lammps.has_mpi_support`
- `lammps.has_exceptions`
- `lammps.has_gzip_support`
- `lammps.has_png_support`
- `lammps.has_jpeg_support`
- `lammps.has_ffmpeg_support`
- `lammps.installed_packages`
- `lammps.get_accelerator_config`
- `lammps.has_style()`
- `lammps.available_styles()`

## 13.4 The `lammps` Python module

The LAMMPS Python interface is implemented as a module called `lammps` which is defined in the `lammps` package in the `python` folder of the LAMMPS source code distribution. After compilation of LAMMPS, the module can be installed into a Python system folder or a user folder with `make install-python`. Components of the module can then be loaded into a Python session with the `import` command.

There are multiple Python interface classes in the `lammps` module:

- the `lammps` class. This is a wrapper around the C-library interface and its member functions try to replicate the *C-library API* closely. This is the most feature-complete Python API.
- the `PyLammps` class. This is a more high-level and more Python style class implemented on top of the `lammps` class.
- the `IPyLammps` class is derived from `PyLammps` and adds embedded graphics features to conveniently include LAMMPS into Jupyter notebooks.

---

### Version check

The `lammps` module stores the version number of the LAMMPS version it is installed from. When initializing the `lammps` class, this version is checked to be the same as the result from `lammps.version()`, the version of the LAMMPS shared library that the module interfaces to. If they are not the same an `AttributeError` exception is raised since a mismatch of versions (e.g. due to incorrect use of the `LD_LIBRARY_PATH` or `PYTHONPATH` environment variables) can lead to crashes or data corruption and otherwise incorrect behavior.

---

LAMMPS module global members:

`lammps.__version__`

Numerical representation of the LAMMPS version this module was taken from. Has the same format as the result of `lammps.version()`.

---

### 13.4.1 The `lammps` class API

The `lammps` class is the core of the LAMMPS Python interfaces. It is a wrapper around the *LAMMPS C library API* using the `Python ctypes` module and a shared library compiled from the LAMMPS source code. The individual methods in this class try to closely follow the corresponding C functions. The handle argument that needs to be passed to the C functions is stored internally in the class and automatically added when calling the C library functions. Below is a detailed documentation of the API.

**class** `lammps.lammps`(*name=""*, *cmdargs=None*, *ptr=None*, *comm=None*)

Create an instance of the LAMMPS Python class.

This is a Python wrapper class that exposes the LAMMPS C-library interface to Python. It either requires that LAMMPS has been compiled as shared library which is then dynamically loaded via the `ctypes` Python module or that this module is called from a Python function that is called from a Python interpreter embedded into a LAMMPS executable, for example through the `python invoke` command. When the class is instantiated it calls the `lammps_open()` function of the LAMMPS C-library interface, which in turn will create an instance of the *LAMMPS C++* class. The handle to this C++ class is stored internally and automatically passed to the calls to the C library interface.

#### Parameters

- **name** (*string*) – “machine” name of the shared LAMMPS library (“mpi” loads `liblammps_mpi.so`, “” loads `liblammps.so`)

- **cmdargs** (*list*) – list of command line arguments to be passed to the `lammps_open()` function. The executable name is automatically added.
- **ptr** (*pointer*) – pointer to a LAMMPS C++ class instance when called from an embedded Python interpreter. None means load symbols from shared library.
- **comm** (*MPI\_Comm*) – MPI communicator (as provided by `mpi4py`). None means use `MPI_COMM_WORLD` implicitly.

#### property `numpy`

Return object to access numpy versions of API

It provides alternative implementations of API functions that return numpy arrays instead of ctypes pointers. If numpy is not installed, accessing this property will lead to an ImportError.

##### Returns

instance of numpy wrapper object

##### Return type

`numpy_wrapper`

#### `close()`

Explicitly delete a LAMMPS instance through the C-library interface.

This is a wrapper around the `lammps_close()` function of the C-library interface.

#### `finalize()`

Shut down the MPI communication and Kokkos environment (if active) through the library interface by calling `lammps_mpi_finalize()` and `lammps_kokkos_finalize()`.

You cannot create or use any LAMMPS instances after this function is called unless LAMMPS was compiled without MPI and without Kokkos support.

#### `version()`

Return a numerical representation of the LAMMPS version in use.

This is a wrapper around the `lammps_version()` function of the C-library interface.

##### Returns

version number

##### Return type

int

#### `get_os_info()`

Return a string with information about the OS and compiler runtime

This is a wrapper around the `lammps_get_os_info()` function of the C-library interface.

##### Returns

OS info string

##### Return type

string

#### `get_mpi_comm()`

Get the MPI communicator in use by the current LAMMPS instance

This is a wrapper around the `lammps_get_mpi_comm()` function of the C-library interface. It will return None if either the LAMMPS library was compiled without MPI support or the `mpi4py` Python module is not available.

**Returns**

MPI communicator

**Return type**

MPI\_Comm

**file(*path*)**

Read LAMMPS commands from a file.

This is a wrapper around the `lammps_file()` function of the C-library interface. It will open the file with the name/path *file* and process the LAMMPS commands line by line until the end. The function will return when the end of the file is reached.

**Parameters****path** (*string*) – Name of the file/path with LAMMPS commands**command(*cmd*)**

Process a single LAMMPS input command from a string.

This is a wrapper around the `lammps_command()` function of the C-library interface.

**Parameters****cmd** (*string*) – a single lammps command**commands\_list(*cmdlist*)**

Process multiple LAMMPS input commands from a list of strings.

This is a wrapper around the `lammps_commands_list()` function of the C-library interface.

**Parameters****cmdlist** (*list of strings*) – a single lammps command**commands\_string(*multicmd*)**

Process a block of LAMMPS input commands from a string.

This is a wrapper around the `lammps_commands_string()` function of the C-library interface.

**Parameters****multicmd** (*string*) – text block of lammps commands**get\_natoms()**

Get the total number of atoms in the LAMMPS instance.

Will be precise up to 53-bit signed integer due to the underlying `lammps_get_natoms()` function returning a double.

**Returns**

number of atoms

**Return type**

int

**extract\_box()**

Extract simulation box parameters

This is a wrapper around the `lammps_extract_box()` function of the C-library interface. Unlike in the C function, the result is returned as a list.

**Returns**

list of the extracted data: boxlo, boxhi, xy, yz, xz, periodicity, box\_change

**Return type**

[ 3\*double, 3\*double, double, double, 3\*int, int]

**reset\_box**(*boxlo*, *boxhi*, *xy*, *yz*, *xz*)

Reset simulation box parameters

This is a wrapper around the `lammps_reset_box()` function of the C-library interface.

**Parameters**

- **boxlo** (*list of 3 floating point numbers*) – new lower box boundaries
- **boxhi** (*list of 3 floating point numbers*) – new upper box boundaries
- **xy** (*float*) – xy tilt factor
- **yz** (*float*) – yz tilt factor
- **xz** (*float*) – xz tilt factor

**get\_thermo**(*name*)

Get current value of a thermo keyword

This is a wrapper around the `lammps_get_thermo()` function of the C-library interface.

**Parameters**

**name** (*string*) – name of thermo keyword

**Returns**

value of thermo keyword

**Return type**

double or None

**extract\_setting**(*name*)

Query LAMMPS about global settings that can be expressed as an integer.

This is a wrapper around the `lammps_extract_setting()` function of the C-library interface. Its documentation includes a list of the supported keywords.

**Parameters**

**name** (*string*) – name of the setting

**Returns**

value of the setting

**Return type**

int

**extract\_global\_datatype**(*name*)

Retrieve global property datatype from LAMMPS

This is a wrapper around the `lammps_extract_global_datatype()` function of the C-library interface. Its documentation includes a list of the supported keywords. This function returns None if the keyword is not recognized. Otherwise it will return a positive integer value that corresponds to one of the *data type* constants define in the `lammps` module.

**Parameters**

**name** (*string*) – name of the property

**Returns**

data type of global property, see *Data Types*

**Return type**

int

**extract\_global**(*name*, *dtype=None*)

Query LAMMPS about global settings of different types.

This is a wrapper around the `lammps_extract_global()` function of the C-library interface. Since there are no pointers in Python, this method will - unlike the C function - return the value or a list of values. The `lammps_extract_global()` documentation includes a list of the supported keywords and their data types. Since Python needs to know the data type to be able to interpret the result, by default, this function will try to auto-detect the data type by asking the library. You can also force a specific data type. For that purpose the `lammps` module contains *data type* constants. This function returns `None` if either the keyword is not recognized, or an invalid data type constant is used.

**Parameters**

- **name** (*string*) – name of the property
- **dtype** (*int*, *optional*) – data type of the returned data (see *Data Types*)

**Returns**

value of the property or list of values or `None`

**Return type**

`int`, `float`, `list`, or `NoneType`

**extract\_atom\_datatype**(*name*)

Retrieve per-atom property datatype from LAMMPS

This is a wrapper around the `lammps_extract_atom_datatype()` function of the C-library interface. Its documentation includes a list of the supported keywords. This function returns `None` if the keyword is not recognized. Otherwise it will return an integer value that corresponds to one of the *data type* constants defined in the `lammps` module.

**Parameters**

**name** (*string*) – name of the property

**Returns**

data type of per-atom property (see *Data Types*)

**Return type**

`int`

**extract\_atom**(*name*, *dtype=None*)

Retrieve per-atom properties from LAMMPS

This is a wrapper around the `lammps_extract_atom()` function of the C-library interface. Its documentation includes a list of the supported keywords and their data types. Since Python needs to know the data type to be able to interpret the result, by default, this function will try to auto-detect the data type by asking the library. You can also force a specific data type by setting `dtype` to one of the *data type* constants defined in the `lammps` module. This function returns `None` if either the keyword is not recognized, or an invalid data type constant is used.

---

**Note:** While the returned arrays of per-atom data are dimensioned for the range `[0:nmax]` - as is the underlying storage - the data is usually only valid for the range of `[0:nlocal]`, unless the property of interest is also updated for ghost atoms. In some cases, this depends on a LAMMPS setting, see for example `comm_modify vel yes`.

---

**Parameters**

- **name** (*string*) – name of the property

- **dtype** (*int*, *optional*) – data type of the returned data (see *Data Types*)

**Returns**

requested data or None

**Return type**

ctypes.POINTER(ctypes.c\_int32), ctypes.POINTER(ctypes.POINTER(ctypes.c\_int32)),  
 ctypes.POINTER(ctypes.c\_int64), ctypes.POINTER(ctypes.POINTER(ctypes.c\_int64)),  
 ctypes.POINTER(ctypes.c\_double), ctypes.POINTER(ctypes.POINTER(ctypes.c\_double)),  
 or NoneType

**extract\_compute**(*cid*, *cstyle*, *ctype*)

Retrieve data from a LAMMPS compute

This is a wrapper around the `lammers_extract_compute()` function of the C-library interface. This function returns None if either the compute id is not recognized, or an invalid combination of *cstyle* and *ctype* constants is used. The names and functionality of the constants are the same as for the corresponding C-library function. For requests to return a scalar or a size, the value is returned, otherwise a pointer.

**Parameters**

- **cid** (*string*) – compute ID
- **cstyle** (*int*) – style of the data retrieve (global, atom, or local), see *Style Constants*
- **ctype** (*int*) – type or size of the returned data (scalar, vector, or array), see *Type Constants*

**Returns**

requested data as scalar, pointer to 1d or 2d double array, or None

**Return type**

c\_double, ctypes.POINTER(c\_double), ctypes.POINTER(ctypes.POINTER(c\_double)), or  
 NoneType

**extract\_fix**(*fid*, *fstyle*, *ftype*, *nrow=0*, *ncol=0*)

Retrieve data from a LAMMPS fix

This is a wrapper around the `lammers_extract_fix()` function of the C-library interface. This function returns None if either the fix id is not recognized, or an invalid combination of *fstyle* and *ftype* constants is used. The names and functionality of the constants are the same as for the corresponding C-library function. For requests to return a scalar or a size, the value is returned, also when accessing global vectors or arrays, otherwise a pointer.

**Parameters**

- **fid** (*string*) – fix ID
- **fstyle** (*int*) – style of the data retrieve (global, atom, or local), see *Style Constants*
- **ftype** (*int*) – type or size of the returned data (scalar, vector, or array), see *Type Constants*
- **nrow** (*int*) – index of global vector element or row index of global array element
- **ncol** (*int*) – column index of global array element

**Returns**

requested data or None

**Return type**

c\_double, ctypes.POINTER(c\_double), ctypes.POINTER(ctypes.POINTER(c\_double)), or  
 NoneType

**extract\_variable**(*name*, *group=None*, *vartype=0*)

Evaluate a LAMMPS variable and return its data

This function is a wrapper around the function `lammps_extract_variable()` of the C-library interface, evaluates variable name and returns a copy of the computed data. The memory temporarily allocated by the C-interface is deleted after the data is copied to a Python variable or list. The variable must be either an equal-style (or equivalent) variable or an atom-style variable. The variable type has to be provided as `vartype` parameter which may be one of two constants: `LMP_VAR_EQUAL` or `LMP_VAR_ATOM`; it defaults to equal-style variables. The `group` parameter is only used for atom-style variables and defaults to the group “all” if set to `None`, which is the default.

**Parameters**

- **name** (*string*) – name of the variable to execute
- **group** (*string*, *only for atom-style variables*) – name of group for atom-style variable
- **vartype** (*int*) – type of variable, see *Variable Type Constants*

**Returns**

the requested data

**Return type**

`c_double`, (`c_double`), or `NoneType`

**flush\_buffers**()

Flush output buffers

This is a wrapper around the `lammps_flush_buffers()` function of the C-library interface.

**set\_variable**(*name*, *value*)

Set a new value for a LAMMPS string style variable

This is a wrapper around the `lammps_set_variable()` function of the C-library interface.

**Parameters**

- **name** (*string*) – name of the variable
- **value** (*any. will be converted to a string*) – new variable value

**Returns**

either 0 on success or -1 on failure

**Return type**

`int`

**gather\_bonds**()

Retrieve global list of bonds

This is a wrapper around the `lammps_gather_bonds()` function of the C-library interface.

This function returns a tuple with the number of bonds and a flat list of ctypes integer values with the bond type, bond atom1, bond atom2 for each bond.

New in version 28Jul2021.

**Returns**

a tuple with the number of bonds and a list of `c_int` or `c_long`

**Return type**

(`int`, `3*nbonds*c_tagint`)



**encode\_image\_flags**(*ix, iy, iz*)

convert 3 integers with image flags for x-, y-, and z-direction into a single integer like it is used internally in LAMMPS

This method is a wrapper around the `lammeps_encode_image_flags()` function of library interface.

**Parameters**

- **ix** (*int*) – x-direction image flag
- **iy** (*int*) – y-direction image flag
- **iz** (*int*) – z-direction image flag

**Returns**

encoded image flags

**Return type**

`lammeps.c_imageint`

**decode\_image\_flags**(*image*)

Convert encoded image flag integer into list of three regular integers.

This method is a wrapper around the `lammeps_decode_image_flags()` function of library interface.

**Parameters**

**image** (`lammeps.c_imageint`) – encoded image flags

**Returns**

list of three image flags in x-, y-, and z- direction

**Return type**

list of 3 int

**create\_atoms**(*n, id, type, x, v=None, image=None, shrinkexceed=False*)

Create N atoms from list of coordinates and properties

This function is a wrapper around the `lammeps_create_atoms()` function of the C-library interface, and the behavior is similar except that the *v*, *image*, and *shrinkexceed* arguments are optional and default to *None*, *None*, and *False*, respectively. With *None* being equivalent to a NULL pointer in C.

The lists of coordinates, types, atom IDs, velocities, image flags can be provided in any format that may be converted into the required internal data types. Also the list may contain more than *N* entries, but not fewer. In the latter case, the function will return without attempting to create atoms. You may use the `encode_image_flags` method to properly combine three integers with image flags into a single integer.

**Parameters**

- **n** (*int*) – number of atoms for which data is provided
- **id** (*list of lammeps.tagint*) – list of atom IDs with at least n elements or None
- **type** (*list of int*) – list of atom types
- **x** (*list of float*) – list of coordinates for x-, y-, and z (flat list of 3n entries)
- **v** (*list of float*) – list of velocities for x-, y-, and z (flat list of 3n entries) or None (optional)
- **image** (*list of lammeps.imageint*) – list of encoded image flags (optional)
- **shrinkexceed** (*bool*) – whether to expand shrink-wrap boundaries if atoms are outside the box (optional)

**Returns**

number of atoms created. 0 if insufficient or invalid data

**Return type**

int

**property has\_mpi\_support**

Report whether the LAMMPS shared library was compiled with a real MPI library or in serial.

This is a wrapper around the `lammops_config_has_mpi_support()` function of the library interface.

**Returns**

False when compiled with MPI STUBS, otherwise True

**Return type**

bool

**property is\_running**

Report whether being called from a function during a run or a minimization

Various LAMMPS commands must not be called during an ongoing run or minimization. This property allows to check for that. This is a wrapper around the `lammops_is_running()` function of the library interface.

New in version 9Oct2020.

**Returns**

True when called during a run otherwise false

**Return type**

bool

**force\_timeout()**

Trigger an immediate timeout, i.e. a “soft stop” of a run.

This function allows to cleanly stop an ongoing run or minimization at the next loop iteration. This is a wrapper around the `lammops_force_timeout()` function of the library interface.

New in version 9Oct2020.

**property has\_exceptions**

Report whether the LAMMPS shared library was compiled with C++ exceptions handling enabled

This is a wrapper around the `lammops_config_has_exceptions()` function of the library interface.

**Returns**

state of C++ exception support

**Return type**

bool

**property has\_gzip\_support**

Report whether the LAMMPS shared library was compiled with support for reading and writing compressed files through gzip.

This is a wrapper around the `lammops_config_has_gzip_support()` function of the library interface.

**Returns**

state of gzip support

**Return type**

bool

**property has\_png\_support**

Report whether the LAMMPS shared library was compiled with support for writing images in PNG format.

This is a wrapper around the `lammops_config_has_png_support()` function of the library interface.

**Returns**

state of PNG support

**Return type**

bool

**property has\_jpeg\_support**

Report whether the LAMMPS shared library was compiled with support for writing images in JPEG format.

This is a wrapper around the `lammops_config_has_jpeg_support()` function of the library interface.

**Returns**

state of JPEG support

**Return type**

bool

**property has\_ffmpeg\_support**

State of support for writing movies with ffmpeg in the LAMMPS shared library

This is a wrapper around the `lammops_config_has_ffmpeg_support()` function of the library interface.

**Returns**

state of ffmpeg support

**Return type**

bool

**property accelerator\_config**

Return table with available accelerator configuration settings.

This is a wrapper around the `lammops_config_accelerator()` function of the library interface which loops over all known packages and categories and returns enabled features as a nested dictionary with all enabled settings as list of strings.

**Returns**

nested dictionary with all known enabled settings as list of strings

**Return type**

dictionary

**property has\_gpu\_device**

Availability of GPU package compatible device

This is a wrapper around the `lammops_has_gpu_device()` function of the C library interface.

**Returns**

True if a GPU package compatible device is present, otherwise False

**Return type**

bool

**get\_gpu\_device\_info()**

Return a string with detailed information about any devices that are usable by the GPU package.

This is a wrapper around the `lammops_get_gpu_device_info()` function of the C-library interface.

**Returns**

GPU device info string

**Return type**

string

**property installed\_packages**

List of the names of enabled packages in the LAMMPS shared library

This is a wrapper around the functions `lammps_config_package_count()` and `:cpp:func`lammps_config_package_name`` of the library interface.

:return

**has\_style(category, name)**

Returns whether a given style name is available in a given category

This is a wrapper around the function `lammps_has_style()` of the library interface.

**Parameters**

- **category** (*string*) – name of category
- **name** (*string*) – name of the style

**Returns**

true if style is available in given category

**Return type**

bool

**available\_styles(category)**

Returns a list of styles available for a given category

This is a wrapper around the functions `lammps_style_count()` and `lammps_style_name()` of the library interface.

**Parameters**

**category** (*string*) – name of category

**Returns**

list of style names in given category

**Return type**

list

**has\_id(category, name)**

Returns whether a given ID name is available in a given category

This is a wrapper around the function `lammps_has_id()` of the library interface.

New in version 9Oct2020.

**Parameters**

- **category** (*string*) – name of category
- **name** (*string*) – name of the ID

**Returns**

true if ID is available in given category

**Return type**

bool

**available\_ids**(*category*)

Returns a list of IDs available for a given category

This is a wrapper around the functions `lammops_id_count()` and `lammops_id_name()` of the library interface.

New in version 9Oct2020.

**Parameters**

**category** (*string*) – name of category

**Returns**

list of id names in given category

**Return type**

list

**available\_plugins**(*category*)

Returns a list of plugins available for a given category

This is a wrapper around the functions `lammops_plugin_count()` and `lammops_plugin_name()` of the library interface.

New in version 10Mar2021.

**Returns**

list of style/name pairs of loaded plugins

**Return type**

list

**set\_fix\_external\_callback**(*fix\_id*, *callback*, *caller=None*)

Set the callback function for a fix external instance with a given fix ID.

Optionally also set a reference to the calling object.

This is a wrapper around the `lammops_set_fix_external_callback()` function of the C-library interface. However this is set up to call a Python function with the following arguments.

- object is the value of the “caller” argument
- n timestep is the current timestep
- nlocal is the number of local atoms on the current MPI process
- tag is a 1d NumPy array of integers representing the atom IDs of the local atoms
- x is a 2d NumPy array of doubles of the coordinates of the local atoms
- f is a 2d NumPy array of doubles of the forces on the local atoms that will be added

Changed in version 28Jul2021.

**Parameters**

- **fix\_id** – Fix-ID of a fix external instance
- **callback** – Python function that will be called from fix external
- **caller** – reference to some object passed to the callback function

**Type**

string

**Type**

function

**Type**

object, optional

**fix\_external\_get\_force**(*fix\_id*)

Get access to the array with per-atom forces of a fix external instance with a given fix ID.

This is a wrapper around the `lammops_fix_external_get_force()` function of the C-library interface.

New in version 28Jul2021.

**Parameters**

**fix\_id** – Fix-ID of a fix external instance

**Type**

string

**Returns**

requested data

**Return type**`ctypes.POINTER(ctypes.POINTER(ctypes.double))`**fix\_external\_set\_energy\_global**(*fix\_id, eng*)

Set the global energy contribution for a fix external instance with the given ID.

This is a wrapper around the `lammops_fix_external_set_energy_global()` function of the C-library interface.

New in version 28Jul2021.

**Parameters**

- **fix\_id** – Fix-ID of a fix external instance
- **eng** – potential energy value to be added by fix external

**Type**

string

**Type**

float

**fix\_external\_set\_virial\_global**(*fix\_id, virial*)

Set the global virial contribution for a fix external instance with the given ID.

This is a wrapper around the `lammops_fix_external_set_virial_global()` function of the C-library interface.

New in version 28Jul2021.

**Parameters**

- **fix\_id** – Fix-ID of a fix external instance
- **eng** – list of 6 floating point numbers with the virial to be added by fix external

**Type**

string

**Type**

float

**fix\_external\_set\_energy\_peratom**(*fix\_id, eatom*)

Set the per-atom energy contribution for a fix external instance with the given ID.

This is a wrapper around the `lammops_fix_external_set_energy_peratom()` function of the C-library interface.

New in version 28Jul2021.

**Parameters**

- **fix\_id** – Fix-ID of a fix external instance
- **eatom** – list of potential energy values for local atoms to be added by fix external

**Type**

string

**Type**

float

**fix\_external\_set\_virial\_peratom**(*fix\_id, vatom*)

Set the per-atom virial contribution for a fix external instance with the given ID.

This is a wrapper around the `lammops_fix_external_set_virial_peratom()` function of the C-library interface.

New in version 28Jul2021.

**Parameters**

- **fix\_id** – Fix-ID of a fix external instance
- **vatom** – list of natoms lists with 6 floating point numbers to be added by fix external

**Type**

string

**Type**

float

**fix\_external\_set\_vector\_length**(*fix\_id, length*)

Set the vector length for a global vector stored with fix external for analysis

This is a wrapper around the `lammops_fix_external_set_vector_length()` function of the C-library interface.

New in version 28Jul2021.

**Parameters**

- **fix\_id** – Fix-ID of a fix external instance
- **length** – length of the global vector

**Type**

string

**Type**

int

**fix\_external\_set\_vector**(*fix\_id, idx, val*)

Store a global vector value for a fix external instance with the given ID.

This is a wrapper around the `lammops_fix_external_set_vector()` function of the C-library interface.

New in version 28Jul2021.

**Parameters**

- **fix\_id** – Fix-ID of a fix external instance
- **idx** – 1-based index of the value in the global vector
- **val** – value to be stored in the global vector

**Type**

string

**Type**

int

**Type**

float

**get\_neighlist(*idx*)**

Returns an instance of *NeighList* which wraps access to the neighbor list with the given index

See *lammps.numpy.get\_neighlist()* if you want to use NumPy arrays instead of *c\_int* pointers.

**Parameters**

**idx** (*int*) – index of neighbor list

**Returns**

an instance of *NeighList* wrapping access to neighbor list data

**Return type**

*NeighList*

**get\_neighlist\_size(*idx*)**

Return the number of elements in neighbor list with the given index

**Parameters**

**idx** (*int*) – neighbor list index

**Returns**

number of elements in neighbor list with index *idx*

**Return type**

int

**get\_neighlist\_element\_neighbors(*idx*, *element*)**

Return data of neighbor list entry

**Parameters**

- **element** (*int*) – neighbor list index
- **element** – neighbor list element index

**Returns**

tuple with atom local index, number of neighbors and array of neighbor local atom indices

**Return type**

(int, int, POINTER(c\_int))

**find\_pair\_neighlist(*style*, *exact=True*, *nsub=0*, *reqid=0*)**

Find neighbor list index of pair style neighbor list

Search for a neighbor list requested by a pair style instance that matches “style”. If *exact* is True, the pair style name must match exactly. If *exact* is False, the pair style name is matched against “style” as regular expression or sub-string. If the pair style is a hybrid pair style, the style is instead matched against the



hybrid sub-styles. If the same pair style is used as sub-style multiple types, you must set `nsub` to a value `n > 0` which indicates the `n`th instance of that sub-style to be used (same as for the `pair_coeff` command). The default value of 0 will fail to match in that case.

Once the pair style instance has been identified, it may have requested multiple neighbor lists. Those are uniquely identified by a request ID `> 0` as set by the pair style. Otherwise the request ID is 0.

#### Parameters

- **style** (*string*) – name of pair style that should be searched for
- **exact** (*bool, optional*) – controls whether style should match exactly or only must be contained in pair style name, defaults to True
- **nsub** (*int, optional*) – match `nsub`-th hybrid sub-style, defaults to 0
- **reqid** (*int, optional*) – list request id, `> 0` in case there are more than one, defaults to 0

#### Returns

neighbor list index if found, otherwise -1

#### Return type

int

#### **find\_fix\_neighlist** (*fixid, reqid=0*)

Find neighbor list index of fix neighbor list

The fix instance requesting the neighbor list is uniquely identified by the fix ID. In case the fix has requested multiple neighbor lists, those are uniquely identified by a request ID `> 0` as set by the fix. Otherwise the request ID is 0 (the default).

#### Parameters

- **fixid** (*string*) – name of fix
- **reqid** (*int, optional*) – id of neighbor list request, in case there are more than one request, defaults to 0

#### Returns

neighbor list index if found, otherwise -1

#### Return type

int

#### **find\_compute\_neighlist** (*computeid, reqid=0*)

Find neighbor list index of compute neighbor list

The compute instance requesting the neighbor list is uniquely identified by the compute ID. In case the compute has requested multiple neighbor lists, those are uniquely identified by a request ID `> 0` as set by the compute. Otherwise the request ID is 0 (the default).

#### Parameters

- **computeid** (*string*) – name of compute
- **reqid** (*int, optional*) – index of neighbor list request, in case there are more than one request, defaults to 0

#### Returns

neighbor list index if found, otherwise -1

#### Return type

int

```
class lammps.numpy_wrapper.numpy_wrapper(lmp)
```

lammps API NumPy Wrapper

This is a wrapper class that provides additional methods on top of an existing *lammps* instance. The methods transform raw ctypes pointers into NumPy arrays, which give direct access to the original data while protecting against out-of-bounds accesses.

There is no need to explicitly instantiate this class. Each instance of *lammps* has a `numpy` property that returns an instance.

#### Parameters

**lmp** (*Lammps*) – instance of the *lammps* class

```
extract_atom(name, dtype=None, nelem=None, dim=None)
```

Retrieve per-atom properties from LAMMPS as NumPy arrays

This is a wrapper around the `lammps.extract_atom()` method. It behaves the same as the original method, but returns NumPy arrays instead of ctypes pointers.

---

**Note:** While the returned arrays of per-atom data are dimensioned for the range `[0:nmax]` - as is the underlying storage - the data is usually only valid for the range of `[0:nlocal]`, unless the property of interest is also updated for ghost atoms. In some cases, this depends on a LAMMPS setting, see for example *comm\_modify vel yes*.

---

#### Parameters

- **name** (*string*) – name of the property
- **dtype** (*int, optional*) – type of the returned data (see *Data Types*)
- **nelem** (*int, optional*) – number of elements in array
- **dim** (*int, optional*) – dimension of each element

#### Returns

requested data as NumPy array with direct access to C data or None

#### Return type

numpy.array or NoneType

```
extract_compute(cid, cstyle, ctype)
```

Retrieve data from a LAMMPS compute

This is a wrapper around the `lammps.extract_compute()` method. It behaves the same as the original method, but returns NumPy arrays instead of ctypes pointers.

#### Parameters

- **cid** (*string*) – compute ID
- **cstyle** (*int*) – style of the data retrieve (global, atom, or local), see *Style Constants*
- **ctype** (*int*) – type of the returned data (scalar, vector, or array), see *Type Constants*

#### Returns

requested data either as float, as NumPy array with direct access to C data, or None

#### Return type

float, numpy.array, or NoneType

**extract\_fix**(*fid, fstyle, ftype, nrow=0, ncol=0*)

Retrieve data from a LAMMPS fix

This is a wrapper around the `lammops.extract_fix()` method. It behaves the same as the original method, but returns NumPy arrays instead of ctypes pointers.

**Parameters**

- **fid** (*string*) – fix ID
- **fstyle** (*int*) – style of the data retrieve (global, atom, or local), see *Style Constants*
- **ftype** (*int*) – type or size of the returned data (scalar, vector, or array), see *Type Constants*
- **nrow** (*int*) – index of global vector element or row index of global array element
- **ncol** (*int*) – column index of global array element

**Returns**

requested data

**Return type**

integer or double value, pointer to 1d or 2d double array or None

**extract\_variable**(*name, group=None, vartype=0*)

Evaluate a LAMMPS variable and return its data

This function is a wrapper around the function `lammops.extract_variable()` method. It behaves the same as the original method, but returns NumPy arrays instead of ctypes pointers.

**Parameters**

- **name** (*string*) – name of the variable to execute
- **group** (*string*) – name of group for atom-style variable (ignored for equal-style variables)
- **vartype** (*int*) – type of variable, see *Variable Type Constants*

**Returns**

the requested data or None

**Return type**

c\_double, numpy.array, or NoneType

**gather\_bonds**()

Retrieve global list of bonds as NumPy array

This is a wrapper around `lammops.gather_bonds()` It behaves the same as the original method, but returns a NumPy array instead of a ctypes list.

New in version 28Jul2021.

**Returns**

the requested data as a 2d-integer numpy array

**Return type**

numpy.array(nbonds,3)

**fix\_external\_get\_force**(*fix\_id*)

Get access to the array with per-atom forces of a fix external instance with a given fix ID.

This function is a wrapper around the `lammops.fix_external_get_force()` method. It behaves the same as the original method, but returns a NumPy array instead of a ctypes pointer.

Changed in version 28Jul2021.

**Parameters**

**fix\_id** – Fix-ID of a fix external instance

**Type**

string

**Returns**

requested data

**Return type**

numpy.array

**fix\_external\_set\_energy\_peratom**(*fix\_id, eatom*)

Set the per-atom energy contribution for a fix external instance with the given ID.

This function is an alternative to `lammops.fix_external_set_energy_peratom()` method. It behaves the same as the original method, but accepts a NumPy array instead of a list as argument.

New in version 28Jul2021.

**Parameters**

- **fix\_id** – Fix-ID of a fix external instance
- **eatom** – per-atom potential energy

**Type**

string

**Type**

numpy.array

**fix\_external\_set\_virial\_peratom**(*fix\_id, vatom*)

Set the per-atom virial contribution for a fix external instance with the given ID.

This function is an alternative to `lammops.fix_external_set_virial_peratom()` method. It behaves the same as the original method, but accepts a NumPy array instead of a list as argument.

New in version 28Jul2021.

**Parameters**

- **fix\_id** – Fix-ID of a fix external instance
- **eatom** – per-atom potential energy

**Type**

string

**Type**

numpy.array

**get\_neighlist**(*idx*)

Returns an instance of `NumPyNeighList` which wraps access to the neighbor list with the given index

**Parameters**

**idx** (*int*) – index of neighbor list

**Returns**

an instance of `NumPyNeighList` wrapping access to neighbor list data

**Return type**

`NumPyNeighList`

**get\_neighlist\_element\_neighbors**(*idx, element*)

Return data of neighbor list entry

This function is a wrapper around the function `lammops.get_neighlist_element_neighbors()` method. It behaves the same as the original method, but returns a NumPy array containing the neighbors instead of a ctypes pointer.

**Parameters**

- **element** (*int*) – neighbor list index
- **element** – neighbor list element index

**Returns**

tuple with atom local index and numpy array of neighbor local atom indices

**Return type**

(int, numpy.array)

### 13.4.2 The PyLammps class API

The `PyLammps` class is a wrapper that creates a simpler, more “Pythonic” interface to common LAMMPS functionality. LAMMPS data structures are exposed through objects and properties. This makes Python scripts shorter and more concise. See the *PyLammps Tutorial* for an introduction on how to use this interface.

```
class lammops.PyLammps(name="", cmdargs=None, ptr=None, comm=None, verbose=False)
```

This is a Python wrapper class around the lower-level `lammops` class, exposing a more Python-like, object-oriented interface for prototyping system inside of IPython and Jupyter notebooks.

It either creates its own instance of `lammops` or can be initialized with an existing instance. The arguments are the same of the lower-level interface. The original interface can still be accessed via `PyLammps.lmp`.

**Parameters**

- **name** (*string*) – “machine” name of the shared LAMMPS library (“mpi” loads `liblammops_mpi.so`, “” loads `liblammops.so`)
- **cmdargs** (*list*) – list of command line arguments to be passed to the `lammops_open()` function. The executable name is automatically added.
- **ptr** (*pointer*) – pointer to a LAMMPS C++ class instance when called from an embedded Python interpreter. None means load symbols from shared library.
- **comm** (*MPI\_Comm*) – MPI communicator (as provided by `mpi4py`). None means use `MPI_COMM_WORLD` implicitly.
- **verbose** (*bool*) – print all LAMMPS output to stdout

**Variables**

- **lmp** (*lammops*) – instance of original LAMMPS Python interface
- **runs** – list of completed runs, each storing the thermo output

**close()**

Explicitly delete a LAMMPS instance

This is a wrapper around the `lammops.close()` of the Python interface.

**version()**

Return a numerical representation of the LAMMPS version in use.

This is a wrapper around the `lammps.version()` function of the Python interface.

**Returns**

version number

**Return type**

int

**file(*file*)**

Read LAMMPS commands from a file.

This is a wrapper around the `lammps.file()` function of the Python interface.

**Parameters**

**path** (*string*) – Name of the file/path with LAMMPS commands

**property enable\_cmd\_history****Getter**

Return whether command history is saved

**Setter**

Set if command history should be saved

**Type**

bool

**write\_script(*filepath*)**

Write LAMMPS script file containing all commands executed up until now

**Parameters**

**filepath** (*string*) – path to script file that should be written

**clear\_cmd\_history()**

Clear LAMMPS command history up to this point

**command(*cmd*)**

Execute LAMMPS command

If `PyLammps.enable_cmd_history` is set to `True`, commands executed will be recorded. The entire command history can be written to a file using `PyLammps.write_script()`. To clear the command history, use `PyLammps.clear_cmd_history()`.

**Parameters**

**cmd** – command string that should be executed

**Type**

cmd: string

**run(\*args, \*\*kwargs)**

Execute LAMMPS run command with given arguments

All thermo output during the run is captured and saved as new entry in `PyLammps.runs`. The latest run can be retrieved by `PyLammps.last_run`.

**property last\_run**

Return data produced of last completed run command

**Getter**

Returns an object containing information about the last run command

**Type**  
dict

**property atoms**

All atoms of this LAMMPS instance

**Getter**  
Returns a list of atoms currently in the system

**Type**  
*AtomList*

**property system**

The system state of this LAMMPS instance

**Getter**  
Returns an object with properties storing the current system state

**Type**  
namedtuple

**property communication**

The communication state of this LAMMPS instance

**Getter**  
Returns an object with properties storing the current communication state

**Type**  
namedtuple

**property computes**

The list of active computes of this LAMMPS instance

**Getter**  
Returns a list of computes that are currently active in this LAMMPS instance

**Type**  
list

**property dumps**

The list of active dumps of this LAMMPS instance

**Getter**  
Returns a list of dumps that are currently active in this LAMMPS instance

**Type**  
list

**property fixes**

The list of active fixes of this LAMMPS instance

**Getter**  
Returns a list of fixes that are currently active in this LAMMPS instance

**Type**  
list

**property groups**

The list of active atom groups of this LAMMPS instance

**Getter**  
Returns a list of atom groups that are currently active in this LAMMPS instance

**Type**  
list

**property variables**

Returns a dictionary of all variables defined in the current LAMMPS instance

**Getter**  
Returns a dictionary of all variables that are defined in this LAMMPS instance

**Type**  
dict

**eval(*expr*)**

Evaluate expression

**Parameters**

**expr** (*string*) – the expression string that should be evaluated inside of LAMMPS

**Returns**

the value of the evaluated expression

**Return type**

float if numeric, string otherwise

**lmp\_print(*s*)**

needed for Python2 compatibility, since print is a reserved keyword

**class** `lammmps.AtomList(pylammmps_instance)`

A dynamic list of atoms that returns either an `Atom` or `Atom2D` instance for each atom. Instances are only allocated when accessed.

**Variables**

- **natoms** – total number of atoms
- **dimensions** – number of dimensions in system

**class** `lammmps.Atom(pylammmps_instance, index)`

A wrapper class then represents a single atom inside of LAMMPS

It provides access to properties of the atom and allows you to change some of them.

**property id**

Return the atom ID

**Type**  
int

**property type**

Return the atom type

**Type**  
int

**property mol**

Return the atom molecule index

**Type**  
int



**property mass**

Return the atom mass

**Type**

float

**property radius**

Return the particle radius

**Type**

float

**property position****Getter**

Return position of atom

**Setter**

Set position of atom

**Type**

numpy.array (float, float, float)

**property velocity****Getter**

Return velocity of atom

**Setter**

Set velocity of atom

**Type**

numpy.array (float, float, float)

**property force**

Return the total force acting on the atom

**Type**

numpy.array (float, float, float)

**property omega**

Return the rotational velocity of the particle

**Type**

numpy.array (float, float, float)

**property torque**

Return the total torque acting on the particle

**Type**

numpy.array (float, float, float)

**property angular\_momentum**

Return the angular momentum of the particle

**Type**

numpy.array (float, float, float)

**property charge**

Return the atom charge

**Type**

float

**class** `lammops.Atom2D`(*pylammops\_instance, index*)

A wrapper class then represents a single 2D atom inside of LAMMPS

Inherits all properties from the `Atom` class, but returns 2D versions of position, velocity, and force.

It provides access to properties of the atom and allows you to change some of them.

**property position**

Access to coordinates of an atom

**Getter**

Return position of atom

**Setter**

Set position of atom

**Type**

`numpy.array` (float, float)

**property velocity**

Access to velocity of an atom :getter: Return velocity of atom :setter: Set velocity of atom :type: `numpy.array` (float, float)

**property force**

Access to force of an atom :getter: Return force of atom :setter: Set force of atom :type: `numpy.array` (float, float)

---

### 13.4.3 The `IPyLammops` class API

The `IPyLammops` class is an extension of `PyLammops`, adding additional functions to quickly display visualizations such as images and videos inside of IPython. See the [PyLammops Tutorial](#) for examples.

**class** `lammops.IPyLammops`(*name=""*, *cmdargs=None*, *ptr=None*, *comm=None*)

IPython wrapper for LAMMPS which adds embedded graphics capabilities to PyLammops interface

It either creates its own instance of `lammops` or can be initialized with an existing instance. The arguments are the same of the lower-level interface. The original interface can still be accessed via `PyLammops.lmp`.

**Parameters**

- **name** (*string*) – “machine” name of the shared LAMMPS library (“mpi” loads `liblammops_mpi.so`, “” loads `liblammops.so`)
- **cmdargs** (*list*) – list of command line arguments to be passed to the `lammops_open()` function. The executable name is automatically added.
- **ptr** (*pointer*) – pointer to a LAMMPS C++ class instance when called from an embedded Python interpreter. None means load symbols from shared library.
- **comm** (*MPI\_Comm*) – MPI communicator (as provided by `mpi4py`). None means use `MPI_COMM_WORLD` implicitly.

**image**(*filename='snapshot.png'*, *group='all'*, *color='type'*, *diameter='type'*, *size=None*, *view=None*, *center=None*, *up=None*, *zoom=1.0*, *background\_color='white'*)

Generate image using `write_dump` command and display it

See [dump image](#) for more information.

**Parameters**

- **filename** (*string*) – Name of the image file that should be generated. The extension determines whether it is PNG or JPEG
- **group** (*string*) – the group of atoms `write_image` should use
- **color** (*string*) – name of property used to determine color
- **diameter** (*string*) – name of property used to determine atom diameter
- **size** (*tuple (width, height)*) – dimensions of image
- **view** (*tuple (theta, phi)*) – view parameters
- **center** (*tuple (flag, center\_x, center\_y, center\_z)*) – center parameters
- **up** (*tuple (up\_x, up\_y, up\_z)*) – vector pointing to up direction
- **zoom** (*float*) – zoom factor
- **background\_color** (*string*) – background color of scene

**Returns**

Image instance used to display image in notebook

**Return type**

`IPython.core.display.Image`

**video**(*filename*)

Load video from file

Can be used to visualize videos from *dump movie*.

**Parameters**

**filename** (*string*) – Path to video file

**Returns**

HTML Video Tag used by notebook to embed a video

**Return type**

`IPython.display.HTML`

---

### 13.4.4 Additional components of the `lammmps` module

The `lammmps` module additionally contains several constants and the `NeighList` class:

#### Data Types

`LAMMPS_INT`, `LAMMPS_INT_2D`, `LAMMPS_DOUBLE`, `LAMMPS_DOUBLE_2D`, `LAMMPS_INT64`,  
`LAMMPS_INT64_2D`, `LAMMPS_STRING`

Constants in the `lammmps` module to indicate how to cast data when the C library function returns a void pointer. Used in `lammmps.extract_global()` and `lammmps.extract_atom()`. See `_LMP_DATATYPE_CONST` for the equivalent constants in the C library interface.

## Style Constants

### LMP\_STYLE\_GLOBAL, LMP\_STYLE\_ATOM, LMP\_STYLE\_LOCAL

Constants in the *lammps* module to select what style of data to request from computes or fixes. See *\_LMP\_STYLE\_CONST* for the equivalent constants in the C library interface. Used in *lammps.extract\_compute()*, *lammps.extract\_fix()*, and their NumPy variants *lammps.numpy.extract\_compute()* and *lammps.numpy.extract\_fix()*.

## Type Constants

### LMP\_TYPE\_SCALAR, LMP\_TYPE\_VECTOR, LMP\_TYPE\_ARRAY, LMP\_SIZE\_VECTOR, LMP\_SIZE\_ROWS, LMP\_SIZE\_COLS

Constants in the *lammps* module to select what type of data to request from computes or fixes. See *\_LMP\_TYPE\_CONST* for the equivalent constants in the C library interface. Used in *lammps.extract\_compute()*, *lammps.extract\_fix()*, and their NumPy variants *lammps.numpy.extract\_compute()* and *lammps.numpy.extract\_fix()*.

## Variable Type Constants

### LMP\_VAR\_EQUAL, LMP\_VAR\_ATOM

Constants in the *lammps* module to select what type of variable to query when calling *lammps.extract\_variable()*. See also: *variable command*.

## Classes representing internal objects

### `class lammps.NeighList(lmp, idx)`

This is a wrapper class that exposes the contents of a neighbor list.

It can be used like a regular Python list. Each element is a tuple of:

- the atom local index
- its number of neighbors
- and a pointer to an `c_int` array containing local atom indices of its neighbors

Internally it uses the lower-level LAMMPS C-library interface.

#### Parameters

- **`lmp`** (`lammps`) – reference to instance of *lammps*
- **`idx`** (`int`) – neighbor list index

#### property `size`

##### Returns

number of elements in neighbor list

#### `get(element)`

Access a specific neighbor list entry. “element” must be a number from 0 to the size-1 of the list

##### Returns

tuple with atom local index, number of neighbors and ctypes pointer to neighbor’s local atom indices

**Return type**

(int, int, ctypes.POINTER(c\_int))

**find**(*iatom*)

Find the neighbor list for a specific (local) atom *iatom*. If there is no list for *iatom*, (-1, None) is returned.

**Returns**

tuple with number of neighbors and ctypes pointer to neighbor's local atom indices

**Return type**

(int, ctypes.POINTER(c\_int))

**class** `lammops.numpy_wrapper.NumPyNeighList`(*lmp*, *idx*)

This is a wrapper class that exposes the contents of a neighbor list.

It can be used like a regular Python list. Each element is a tuple of:

- the atom local index
- a NumPy array containing the local atom indices of its neighbors

Internally it uses the lower-level LAMMPS C-library interface.

**Parameters**

- **lmp** (`lammops`) – reference to instance of `lammops`
- **idx** (`int`) – neighbor list index

**get**(*element*)

Access a specific neighbor list entry. “element” must be a number from 0 to the size-1 of the list

**Returns**

tuple with atom local index, numpy array of neighbor local atom indices

**Return type**

(int, numpy.array)

**find**(*iatom*)

Find the neighbor list for a specific (local) atom *iatom*. If there is no list for *iatom*, None is returned.

**Returns**

numpy array of neighbor local atom indices

**Return type**

numpy.array or None

## 13.5 Extending the Python interface

As noted previously, most of the `lammops` Python class methods correspond one-to-one with the functions in the LAMMPS library interface in `src/library.cpp` and `library.h`. This means you can extend the Python wrapper by following these steps:

- Add a new interface function to `src/library.cpp` and `src/library.h`.
- Rebuild LAMMPS as a shared library.
- Add a wrapper method to `python/lammops/core.py` for this interface function.
- Define the corresponding `argtypes` list and `restype` in the `lammops.__init__()` function.
- Re-install the shared library and the python module, if needed

- You should now be able to invoke the new interface function from a Python script.

## 13.6 Calling Python from LAMMPS

LAMMPS has several commands which can be used to invoke Python code directly from an input script:

- *python*
- *variable python*
- *fix python/invoke*
- *pair\_style python*

The *python* command which can be used to define and execute a Python function that you write the code for. The Python function can also be assigned to a LAMMPS python-style variable via the *variable* command. Each time the variable is evaluated, either in the LAMMPS input script itself, or by another LAMMPS command that uses the variable, this will trigger the Python function to be invoked.

The Python code for the function can be included directly in the input script or in an auxiliary file. The function can have arguments which are mapped to LAMMPS variables (also defined in the input script) and it can return a value to a LAMMPS variable. This is thus a mechanism for your input script to pass information to a piece of Python code, ask Python to execute the code, and return information to your input script.

Note that a Python function can be arbitrarily complex. It can import other Python modules, instantiate Python classes, call other Python functions, etc. The Python code that you provide can contain more code than the single function. It can contain other functions or Python classes, as well as global variables or other mechanisms for storing state between calls from LAMMPS to the function.

The Python function you provide can consist of “pure” Python code that only performs operations provided by standard Python. However, the Python function can also “call back” to LAMMPS through its Python-wrapped library interface, in the manner described in the *Python run* doc page. This means it can issue LAMMPS input script commands or query and set internal LAMMPS state. As an example, this can be useful in an input script to create a more complex loop with branching logic, than can be created using the simple looping and branching logic enabled by the *next* and *if* commands.

See the *python* page and the *variable* doc page for its python-style variables for more info, including examples of Python code you can write for both pure Python operations and callbacks to LAMMPS.

## 13.7 Output Readers

The Python package contains the *lammops.formats* module, which provides classes to post-process some of the output files generated by LAMMPS.

**class** `lammops.formats.LogFile(filename)`

Reads LAMMPS log files and extracts the thermo information

It supports the line, multi, and yaml thermo output styles.

### Parameters

**filename** (*str*) – path to log file

### Variables

- **runs** – List of LAMMPS runs in log file. Each run is a dictionary with thermo fields as keys, storing the values over time
- **errors** – List of error lines in log file

```
class lammops.formats.AvgChunkFile(filename)
```

Reads files generated by fix ave/chunk

#### Parameters

**filename** (*str*) – path to ave/chunk file

#### Variables

- **timesteps** – List of timesteps stored in file
- **total\_count** – total count over time
- **chunks** – List of chunks. Each chunk is a dictionary containing its ID, the coordinates, and the averaged quantities

## 13.8 Example Python scripts

The `python/examples` directory has Python scripts which show how Python can run LAMMPS, grab data, change it, and put it back into LAMMPS.

These are the Python scripts included as demos in the `python/examples` directory of the LAMMPS distribution, to illustrate the kinds of things that are possible when Python wraps LAMMPS. If you create your own scripts, send them to us and we can include them in the LAMMPS distribution.

<code>trivial.py</code>	read/run a LAMMPS input script through Python
<code>demo.py</code>	invoke various LAMMPS library interface routines
<code>simple.py</code>	run in parallel, similar to <code>examples/COUPLE/simple/simple.cpp</code>
<code>split.py</code>	same as <code>simple.py</code> but running in parallel on a subset of procs
<code>gui.py</code>	GUI go/stop/temperature-slider to control LAMMPS
<code>plot.py</code>	real-time temperature plot with GnuPlot via Pizza.py
<code>viz_TOOL.py</code>	real-time viz via some viz package
<code>vizplotgui_TOOL.py</code>	combination of <code>viz_TOOL.py</code> and <code>plot.py</code> and <code>gui.py</code>

For the `viz_TOOL.py` and `vizplotgui_TOOL.py` commands, replace `TOOL` with `gl` or `atomeye` or `pymol` or `vmd`, depending on what visualization package you have installed.

Note that for GL, you need to be able to run the Pizza.py GL tool, which is included in the `pizza` sub-directory. See the Pizza.py doc pages for more info:

- <https://lammops.github.io/pizza>

Note that for AtomEye, you need version 3, and there is a line in the scripts that specifies the path and name of the executable. See the AtomEye web pages for more details:

- <http://li.mit.edu/Archive/Graphics/A/>
- <http://li.mit.edu/Archive/Graphics/A3/A3.html>

The latter link is to AtomEye 3 which has the scripting capability needed by these Python scripts.

Note that for PyMol, you need to have built and installed the open-source version of PyMol in your Python, so that you can import it from a Python script. See the PyMol web pages for more details:

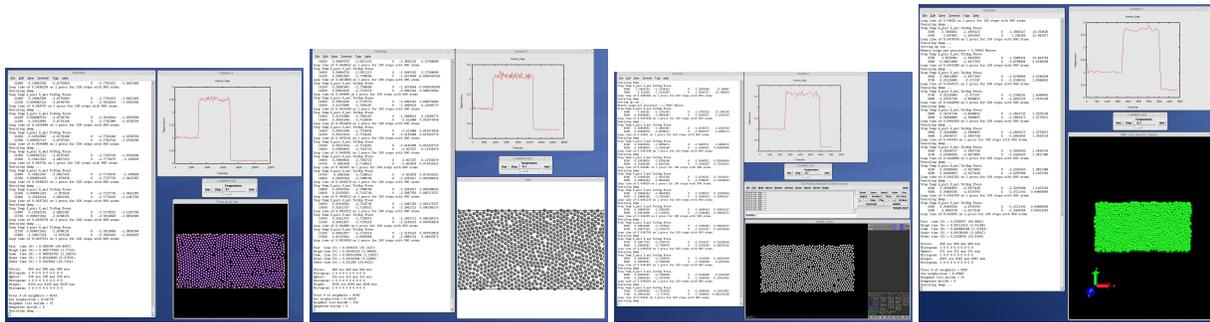
- <https://www.pymol.org>
- <https://github.com/schrodinger/pymol-open-source>

The latter link is to the open-source version.

Note that for VMD, you need a fairly current version (1.8.7 works for me) and there are some lines in the `pizza/vmd.py` script for 4 PIZZA variables that have to match the VMD installation on your system.

See the `python/README` file for instructions on how to run them and the source code for individual scripts for comments about what they do.

Here are screenshots of the `vizplotgui_tool.py` script in action for different visualization package options:



## 13.9 Handling LAMMPS errors

Compiling the shared library with *C++ exception support* provides a better error handling experience. Without exceptions the LAMMPS code will terminate the current Python process with an error message. C++ exceptions allow capturing them on the C++ side and rethrowing them on the Python side. This way LAMMPS errors can be handled through the Python exception handling mechanism.

```
from lammps import lammps, MPIAbortException

lmp = lammps()

try:
    # LAMMPS will normally terminate itself and the running process if an error
    # occurs. This would kill the Python interpreter. To avoid this, make sure to
    # compile with LAMMPS_EXCEPTIONS enabled. This ensures the library API calls
    # will not terminate the parent process. Instead, the library wrapper will
    # detect that an error has occurred and throw a Python exception

    lmp.command('unknown')
except MPIAbortException as ae:
    # Single MPI process got killed. This would normally be handled by an MPI abort
    pass
except Exception as e:
    # All (MPI) processes have reached this error
    pass
```

**Warning:** Capturing a LAMMPS exception in Python can still mean that the current LAMMPS process is in an illegal state and must be terminated. It is advised to save your data and terminate the Python instance as quickly as possible.



## 13.10 Troubleshooting

### 13.10.1 Testing if Python can launch LAMMPS

To test if LAMMPS is callable from Python, launch Python interactively and type:

```
>>> from lammps import lammps
>>> lmp = lammps()
```

If you get no errors, you're ready to use LAMMPS from Python. If the second command fails, the most common error to see is

```
OSError: Could not load LAMMPS dynamic library
```

which means Python was unable to load the LAMMPS shared library. This typically occurs if the system can't find the LAMMPS shared library or one of the auxiliary shared libraries it depends on, or if something about the library is incompatible with your Python. The error message should give you an indication of what went wrong.

If your shared library uses a suffix, such as `liblammps_mpi.so`, change the constructor call as follows (see [Creating or deleting a LAMMPS object](#) for more details):

```
>>> lmp = lammps(name='mpi')
```

You can also test the load directly in Python as follows, without first importing from the `lammps` module:

```
>>> from ctypes import CDLL
>>> CDLL("liblammps.so")
```

If an error occurs, carefully go through the steps in [Installing the LAMMPS Python Module and Shared Library](#) and on the [Build\\_basics](#) page about building a shared library.

If you are not familiar with [Python](#), it is a powerful scripting and programming language which can do almost everything that compiled languages like C, C++, or Fortran can do in fewer lines of code. It also comes with a large collection of add-on modules for many purposes (either bundled or easily installed from Python code repositories). The major drawback is slower execution speed of the script code compared to compiled programming languages. But when the script code is interfaced to optimized compiled code, performance can be on par with a standalone executable, for as long as the scripting is restricted to high-level operations. Thus Python is also convenient to use as a “glue” language to “drive” a program through its library interface, or to hook multiple pieces of software together, such as a simulation code and a visualization tool, or to run a coupled multi-scale or multi-physics model.

See the [Coupling LAMMPS to other codes](#) page for more ideas about coupling LAMMPS to other codes. See the [LAMMPS Library Interfaces](#) page for a description of the LAMMPS library interfaces. That interface is exposed to Python either when calling LAMMPS from Python or when calling Python from a LAMMPS input script and then calling back to LAMMPS from Python code. The C-library interface is designed to be easy to add functionality to, thus the Python interface to LAMMPS is easy to extend as well.

If you create interesting Python scripts that run LAMMPS or interesting Python functions that can be called from a LAMMPS input script, that you think would be generally useful, please post them as a pull request to our [GitHub site](#), and they can be added to the LAMMPS distribution or web page.



## MODIFYING & EXTENDING LAMMPS

LAMMPS is designed in a modular fashion and to be easy to modify or extend with new functionality. In fact, about 95% of its source code are optional. The following pages give basic instructions on what is required when adding new styles of different kinds to LAMMPS.

If you add a new feature to LAMMPS and think it will be of general interest to other users, we encourage you to submit it for inclusion in LAMMPS as a pull request on our [GitHub site](#), after reading about *how to prepare your code for submission* and *the style requirements and recommendations*.

### 14.1 Overview

The best way to add a new feature to LAMMPS is to find a similar feature and look at the corresponding source and header files to figure out what it does. You will need some knowledge of C++ to be able to understand the high-level structure of LAMMPS and its class organization, but functions (class methods) that do actual computations are mostly written in C-style code and operate on simple C-style data structures (vectors and arrays). A high-level overview of the programming style choices in LAMMPS is *given elsewhere*.

Most of the new features described on the *Modify* doc page require you to write a new C++ derived class (except for exceptions described below, where you can make small edits to existing files). Creating a new class requires 2 files, a source code file (\*.cpp) and a header file (\*.h). The derived class must provide certain methods to work as a new option. Depending on how different your new feature is compared to existing features, you can either derive from the base class itself, or from a derived class that already exists. Enabling LAMMPS to invoke the new class is as simple as putting the two source files in the src directory and re-building LAMMPS.

The advantage of C++ and its object-orientation is that all the code and variables needed to define the new feature are in the 2 files you write, and thus should not make the rest of LAMMPS more complex or cause side-effect bugs.

Here is a concrete example. Suppose you write 2 files `pair_foo.cpp` and `pair_foo.h` that define a new class `PairFoo` that computes pairwise potentials described in the classic 1997 *paper* by Foo, et al. If you wish to invoke those potentials in a LAMMPS input script with a command like

```
pair_style foo 0.1 3.5
```

then your `pair_foo.h` file should be structured as follows:

```
#ifndef PAIR_CLASS
// clang-format off
PairStyle(foo,PairFoo);
#else
// clang-format on
...
(class definition for PairFoo)
```

(continues on next page)

```
...  
#endif
```

where “foo” is the style keyword in the pair\_style command, and PairFoo is the class name defined in your pair\_foo.cpp and pair\_foo.h files.

When you re-build LAMMPS, your new pairwise potential becomes part of the executable and can be invoked with a pair\_style command like the example above. Arguments like 0.1 and 3.5 can be defined and processed by your new class.

---

**Note:** With the traditional make process, simply adding the new files to the src folder and compiling LAMMPS again for the desired configuration with “make machine” is sufficient. When using CMake, you need to re-run CMake with “cmake .” in the build folder to have it recognize the added files and include them into the build system.

---

As illustrated by this example pair style, many kinds of options are referred to in the LAMMPS documentation as the “style” of a particular command.

The *Modify page* lists all the common styles in LAMMPS, and discusses the header file for the base class that these styles are derived from. Public variables in that file are ones used and set by the derived classes which are also used by the base class. Sometimes they are also used by the rest of LAMMPS. Pure functions, which means functions declared as virtual in the base class header file which are also set to 0, are functions you **must** implement in your new derived class to give it the functionality LAMMPS expects. Virtual functions that are not set to 0 are functions you may override or not. Those are usually defined with an empty function body.

Additionally, new output options can be added directly to the thermo.cpp, dump\_custom.cpp, and variable.cpp files. These are also listed on the *Modify page*.

Here are additional guidelines for modifying LAMMPS and adding new functionality:

- Think about whether what you want to do would be better as a pre- or post-processing step. Many computations are more easily and more quickly done that way.
- Do not try to do anything within the timestepping of a run that is not parallel. For example do not accumulate a bunch of data on a single processor and analyze it. You run the risk of seriously degrading the parallel efficiency this way.
- If your new feature reads arguments or writes output, make sure you follow the unit conventions discussed by the *units* command.

---

(Foo) Foo, Morefoo, and Maxfoo, J of Classic Potentials, 75, 345 (1997).

## 14.2 Submitting new features for inclusion in LAMMPS

We encourage LAMMPS users to submit new features they wrote for LAMMPS to be included into the LAMMPS distribution and thus become easily accessible to all LAMMPS users. The LAMMPS source code is managed with git and public development is hosted on [GitHub](#). You can monitor the repository to be notified of releases, follow the ongoing development, and comment on topics of interest to you.

### 14.2.1 Communication with the LAMMPS developers

For any larger modifications or programming project, you are encouraged to contact the LAMMPS developers ahead of time in order to discuss implementation strategies and coding guidelines. That will make it easier to integrate your contribution and results in less work for everybody involved. You are also encouraged to search through the list of [open issues on GitHub](#) and submit a new issue for a planned feature, so you would not duplicate the work of others (and possibly get scooped by them) or have your work duplicated by others.

For informal communication with the LAMMPS developers you may ask to join the [LAMMPS developers on Slack](#). This slack work space is by invitation only. Thus for access, please send an e-mail to [slack@lammips.org](mailto:slack@lammips.org) explaining what part of LAMMPS you are working on. Only discussions related to LAMMPS development are tolerated in that work space, so this is **NOT** for people that look for help with compiling, installing, or using LAMMPS. Please post a message to the [LAMMPS forum](#) for those purposes.

### 14.2.2 Packages versus individual files

The remainder of this chapter describes how to add new “style” files of various kinds to LAMMPS. Packages are simply collections of one or more such new class files which are invoked as a new style within a LAMMPS input script. In some cases also collections of supporting functions or classes are included as separate files in a package, especially when they can be shared between multiple styles. If designed correctly, these additions typically do not require any changes to the core code of LAMMPS; they are simply add-on files that are compiled with the rest of LAMMPS. To make those styles work, you may need some trivial changes to the core code; an example of a trivial change is making a parent-class method “virtual” when you derive a new child class from it.

If you think your new feature or package requires some non-trivial changes in core LAMMPS files, you should communicate with the LAMMPS developers [on Slack](#), [on GitHub](#), or [via email](#), since we may have recommendations about what changes to do where, or may not want to include certain changes for some reason and thus you would need to look for alternatives.

### 14.2.3 Time and effort required

How quickly your contribution will be integrated can vary a lot. It depends largely on how much effort it will cause the LAMMPS developers to integrate and test it, how many and what kind of changes to the core code are required, how quickly you can address them and of how much interest it is to the larger LAMMPS community. Please see the section on [LAMMPS programming style and requirements](#) for instructions, recommendations, and formal requirements. A small, modular, well written contribution may be integrated within hours, but a complex change that will require a redesign of some core functionality in LAMMPS for a clean integration can take many months until it is considered ready for inclusion (though this is rare).

### 14.2.4 Submission procedure

All changes to LAMMPS (including those from LAMMPS developers) are integrated via pull requests on GitHub and cannot be merged without passing the automated testing and an approving review by a LAMMPS core developer. Thus before submitting your contribution, you should first make certain, that your added or modified code compiles and works correctly with the latest patch-level or development version of LAMMPS and contains all bug fixes from it.

Once you have prepared everything, see the [LAMMPS GitHub Tutorial](#) page for instructions on how to submit your changes or new files through a GitHub pull request yourself. If you are unable or unwilling to submit via GitHub yourself, you may also submit patch files or full files to the LAMMPS developers and ask them to submit a pull request on GitHub on your behalf. Then create a gzipped tar file of all changed or added files or a corresponding patch file using ‘diff -u’ or ‘diff -c’ format and compress it with gzip. Please only use gzip compression, as this works well and is available on all platforms.

If the new features/files are broadly useful we may add them as core files to LAMMPS or as part of a *package*. All packages are listed and described on the *Packages details* doc page.

### 14.2.5 Licensing

Note that by providing us files to release, you agree to make them open-source, i.e. we can release them under the terms of the GPL (version 2) with the rest of LAMMPS. And similarly as part of a LGPL (version 2.1) distribution of LAMMPS that we make available to developers on request only and with files that are not authorized for that kind of distribution removed (e.g. interface to FFTW). See the *LAMMPS license* page for details.

### 14.2.6 External contributions

If you prefer to do so, you can also develop and support your add-on feature **without** having it included in the LAMMPS distribution, for example as a download from a website of your own. See the [External LAMMPS packages and tools](#) page of the LAMMPS website for examples of groups that do this. We are happy to advertise your package and website from that page. Simply email the [developers](#) with info about your package and we will post it there. We recommend to name external packages USER-<name> so they can be easily distinguished from bundled packages that do not have the USER- prefix.

## 14.3 LAMMPS programming style and requirements for contributions

The following is a summary of the current requirements and recommendations for including contributed source code or documentation into the LAMMPS software distribution.

### 14.3.1 Motivation

The LAMMPS developers are committed to providing a software package that is versatile, reliable, high-quality, efficient, portable, and easy to maintain and modify. Achieving all of these goals is challenging since a large part of LAMMPS consists of contributed code from many different authors and not many of them are professionally trained programmers and familiar with the idiosyncrasies of maintaining a large software package. In addition, changes that interfere with the parallel efficiency of the core code must be avoided. As LAMMPS continues to grow and more features and functionality are added, it becomes a necessity to be more discriminating with new contributions while also working at the same time to improve the existing code.

The following requirements and recommendations are provided to help maintaining or improving that status. Where possible we utilize available continuous integration tools to search for common programming mistakes, portability limitations, incompatible formatting, and undesired side effects. It is indicated which requirements are strict, and which represent a preference and thus are negotiable or optional.

Please feel free to contact the LAMMPS core developers in case you need additional explanations or clarifications or in case you need assistance in realizing the (strict) requirements for your contributions.

### 14.3.2 Licensing requirements (strict)

Contributing authors agree when submitting a pull request that their contributions can be distributed under the LAMMPS license conditions. This is the GNU public license in version 2 (not 3 or later) for the publicly distributed versions, e.g. on the LAMMPS homepage or on GitHub. On request we also make a version of LAMMPS available under LGPL 2.1 terms; this will usually be the latest available or a previous stable version with a few LGPL 2.1 incompatible files removed.

Your new source files should have the LAMMPS copyright, GPL notice, and your name and email address at the top, like other user-contributed LAMMPS source files.

Contributions may be under a different license for long as that license does not conflict with the aforementioned terms. Contributions that use code with a conflicting license can be split into two parts:

1. the core parts (i.e. parts that must be in the *src* tree) that are licensed under compatible terms and bundled with the LAMMPS sources
2. an external library that must be downloaded and compiled (either separately or as part of the LAMMPS compilation)

Please note, that this split licensed mode may complicate including the contribution in binary packages.

### 14.3.3 Using Pull Requests on GitHub (preferred)

All contributions to LAMMPS are processed as pull requests on GitHub (this also applies to the work of the core LAMMPS developers). A [tutorial for submitting pull requests on GitHub](#) is provided. If this is still problematic, contributors may contact any of the core LAMMPS developers for help or to create a pull request on their behalf. This latter way of submission may delay the integration as it depends on the amount of time required to prepare the pull request and free time available by the LAMMPS developer in question to spend on this task.

### 14.3.4 Integration Testing (strict)

Contributed code, like all pull requests, must pass the automated tests on GitHub before it can be merged with the LAMMPS distribution. These tests compile LAMMPS in a variety of environments and settings and run the bundled unit tests. At the discretion of the LAMMPS developer managing the pull request, additional tests may be activated that test for “side effects” on running a collection of input decks and create consistent results. Also, the translation of the documentation to HTML and PDF is tested for.

More specifically, this means that contributed source code **must** compile with the most current version of LAMMPS with `-DLAMMPS_BIGBIG` in addition to the default setting of `-DLAMMPS_SMALLBIG`. The code needs to work correctly in both cases and also in serial and parallel using MPI.

Some “disruptive” changes may break tests and require updates to the testing tools or scripts or tests themselves. This is rare. If in doubt, contact the LAMMPS developer that is assigned to the pull request for further details and explanations and suggestions of what needs to be done.

### 14.3.5 Documentation (strict)

Contributions that add new styles or commands or augment existing ones must include the corresponding new or modified documentation in `ReStructuredText` format (.rst files in the `doc/src/` folder). The documentation shall be written in American English and the .rst file must use only ASCII characters so it can be cleanly translated to PDF files (via `sphinx` and PDFLaTeX). Special characters may be included via embedded math expression typeset in a LaTeX subset.

When adding new commands, they need to be integrated into the `sphinx` documentation system, and the corresponding command tables and lists updated. When translating the documentation into html files there should be no warnings. When adding a new package also some lists describing packages must be updated as well as a package specific description added and, if necessary, some package specific build instructions included.

As appropriate, the text files with the documentation can include inline mathematical expression or figures (see `doc/JPG` for examples). Additional PDF files with further details (see `doc/PDF` for examples) may also be included. The page should also include literature citations as appropriate; see the bottom of `doc/fix_nh.rst` for examples and the earlier part of the same file for how to format the cite itself. Citation labels must be unique across **all** .rst files. The “Restrictions” section of the page should indicate if your command is only available if LAMMPS is built with the appropriate FOO package. See other package doc files for examples of how to do this.

Please run at least “make html” and “make spelling” and carefully inspect and proofread the resulting HTML format doc page before submitting your code. Upon submission of a pull request, checks for error free completion of the HTML and PDF build will be performed and also a spell check, a check for correct anchors and labels, and a check for completeness of references all styles in their corresponding tables and lists is run. In case the spell check reports false positives they can be added to the file `doc/utils/sphinx-config/false_positives.txt`

Contributions that add or modify the library interface or “public” APIs from the C++ code or the Fortran module must include suitable doxygen comments in the source and corresponding changes to the documentation sources for the “Programmer Guide” guide section of the LAMMPS manual.

### 14.3.6 Examples (preferred)

In most cases, it is preferred that example scripts (simple, small, fast to complete on 1 CPU) are included that demonstrate the use of new or extended functionality. These are typically under the `examples` or `examples/PACKAGES` directory. A few guidelines for such example input decks.

- commands that generate output should be commented out (except when the output is the sole purpose or the feature, e.g. for a new compute).
- commands like `log`, `echo`, `package`, `processors`, `suffix` may **not** be used in the input file (exception: “processors \* \* 1” or similar is acceptable when used to avoid unwanted domain decomposition of empty volumes).
- outside of the log files no generated output should be included
- custom `thermo_style` settings may not include output measuring CPU or other time as that makes comparing the thermo output between different runs more complicated.
- input files should be named `in.name`, data files should be named `data.name` and log files should be named `log.version.name.<compiler>.<ncpu>`
- the total file size of all the inputs and outputs should be small
- where possible potential files from the “potentials” folder or data file from other folders should be re-used through symbolic links



### 14.3.7 Howto document (optional)

If your feature requires some more complex steps and explanations to be used correctly or some external or bundled tools or scripts, we recommend that you also contribute a *Howto document* providing some more background information and some tutorial material. This can also be used to provide more in-depth explanations for bundled examples.

As a general rule-of-thumb, the more clear and self-explanatory you make your documentation, README files and examples, and the easier you make it for people to get started, the more likely it is that users will try out your new feature.

### 14.3.8 Programming Style Requirements (varied)

The LAMMPS developers aim to employ a consistent programming style and naming conventions across the entire code base, as this helps with maintenance, debugging, and understanding the code, both for developers and users.

The files *pair\_lj\_cut.h*, *pair\_lj\_cut.cpp*, *utils.h*, and *utils.cpp* may serve as representative examples.

#### Command or Style names, file names, and keywords (mostly strict)

All user-visible command or style names should be all lower case and should only use letters, numbers, or forward slashes. They should be descriptive and initialisms should be avoided unless they are well established (e.g. lj for Lennard-Jones). For a compute style “some/name” the source files must be called *compute\_some\_name.h* and *compute\_some\_name.cpp*. The “include guard” would then be *LMP\_COMPUTE\_SOME\_NAME\_H* and the class name *ComputeSomeName*.

#### Whitespace and permissions (preferred)

Source files should not contain TAB characters unless required by the syntax (e.g. in makefiles) and no trailing whitespace. Text files should be added with Unix-style line endings (LF-only). Git will automatically convert those in both directions when running on Windows; use *dos2unix* on Linux machines to convert files. Text files should have a line ending on the last line.

All files should have 0644 permissions, i.e. writable to the user only and readable by all and no executable permissions. Executable permissions (0755) should only be on shell scripts or python or similar scripts for interpreted script languages.

You can check for these issues with the python scripts in the “*tools/coding\_standard*” folder. When run normally with a source file or a source folder as argument, they will list all non-conforming lines. By adding the *-f* flag to the command line, they will modify the flagged files to try removing the detected issues.

#### Indentation and Placement of Braces (strongly preferred)

LAMMPS uses 2 characters per indentation level and lines should be kept within 100 characters wide.

For new files added to the “src” tree, a *clang-format* configuration file is provided under the name *.clang-format*. This file is compatible with clang-format version 8 and later. With that file present files can be reformatted according to the configuration with a command like: *clang-format -i new-file.cpp*. Ideally, this is done while writing the code or before a pull request is submitted. Blocks of code where the reformatting from clang-format yields undesirable output may be protected with placing a pair *// clang-format off* and *// clang-format on* comments around that block.

## Error or warning messages and explanations (preferred)

Changed in version 4May2022.

Starting with LAMMPS version 4 May 2022 the LAMMPS developers have agreed on a new policy for error and warning messages.

Previously, all error and warning strings were supposed to be listed in the class header files with an explanation. Those would then be regularly “harvested” and transferred to alphabetically sorted lists in the manual. To avoid excessively long lists and to reduce effort, this came with a requirement to have rather generic error messages (e.g. “Illegal ... command”). To identify the specific cause, the name of the source file and the line number of the error location would be printed, so that one could look up the cause by reading the source code.

The new policy encourages more specific error messages that ideally indicate the cause directly and no further lookup would be needed. This is aided by using the `{fmt}` library to convert the Error class commands so that they take a variable number of arguments and error text will be treated like a `{fmt}` syntax format string. Error messages should still kept to a single line or two lines at the most.

For more complex explanations or errors that have multiple possible reasons, a paragraph should be added to the *Error\_details* page with an error code reference (e.g. `.. _err0001:`) then the utility function `utils::errorurl()` can be used to generate an URL that will directly lead to that paragraph. An error for missing arguments can be easily generated using the `utils::missing_cmd_args()` convenience function.

The transformation of existing LAMMPS code to this new scheme is ongoing and - given the size of the LAMMPS source code - will take a significant amount of time until completion. However, for new code following the new approach is strongly preferred. The expectation is that the new scheme will make it easier for LAMMPS users, developers, and maintainers.

An example for this approach would be the `src/read_data.cpp` and `src/atom.cpp` files that implement the *read\_data* and *atom\_modify* commands and that may create “*Unknown identifier in data file*” errors that seem difficult to debug for users because they may have one of multiple possible reasons, and thus require some additional explanations.

## Programming language standards (required)

The core of LAMMPS is written in C++11 in a style that can be mostly described as “C with classes”. Advanced C++ features like operator overloading or excessive use of templates are avoided with the intent to keep the code readable to programmers that have limited C++ programming experience. C++ constructs are acceptable when they help improving the readability and reliability of the code, e.g. when using the `std::string` class instead of manipulating pointers and calling the string functions of the C library. In addition a collection of convenient *utility functions and classes* for recurring tasks and a collection of *platform neutral functions* for improved portability are provided.

Included Fortran code has to be compatible with the Fortran 2003 standard. Python code must be compatible with Python 3.5. Large parts of LAMMPS (including the *PYTHON package*) are also compatible with Python 2.7. Compatibility with Python 2.7 is desirable, but compatibility with Python 3.5 is **required**.

Compatibility with these older programming language standards is very important to maintain portability and availability of LAMMPS on many platforms. This applies especially to HPC cluster environments, which tend to be running older software stacks and LAMMPS users may be required to use those older tools for access to advanced hardware features or not have the option to install newer compilers or libraries.

## Programming conventions (varied)

The following is a collection of conventions that should be applied when writing code for LAMMPS. Following these steps will make it much easier to integrate your contribution. Please have a look at the existing files in packages in the src directory for examples. As a demonstration for how can be adapted to these conventions you may compare the REAXFF package with the what it looked like when it was called USER-REAXC. If you are uncertain, please ask.

- system headers or from installed libraries are include with angular brackets (example: `#include <vector>`), while local include file use double quotes (example: `#include "atom.h"`).
- **when including system header files from the C library use the C++-style names** (`<cstdlib>` or `<cstring>`) instead of the C-style names (`<stdlib.h>` or `<string.h>`)
- the order of `#include` statements in a file `some_name.cpp` that implements a class `SomeName` defined in a header file `some_name.h` should be as follows:
  - `#include "some_name.h"` followed by an empty line
  - LAMMPS include files e.g. `#include "comm.h"` or `#include "modify.h"` in alphabetical order followed by an empty line
  - system header files from the C++ or C standard library followed by an empty line
  - `using namespace LAMMPS_NS` or other namespace imports.
- I/O is done via the C-style `stdio` library and **not** `iostreams`.
- Output to the screen and the logfile should be using the corresponding `FILE` pointers and only be done on MPI rank 0. Use the `utils::logmsg()` convenience function where possible.
- Usage of C++11 *virtual*, *override*, *final* keywords: Please follow the [C++ Core Guideline C.128](#). That means, you should only use *virtual* to declare a new virtual function, *override* to indicate you are overriding an existing virtual function, and *final* to prevent any further overriding.
- Trivial destructors: Prefer not writing destructors when they are empty and *default*.

```
// don't write destructors for A or B like this
class A : protected Pointers {
public:
    A();
    ~A() override {}
};

class B : protected Pointers {
public:
    B();
    ~B() override = default;
};

// instead, let the compiler create the implicit default destructor by not writing
→it
class A : protected Pointers {
public:
    A();
};

class B : protected Pointers {
public:
```

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```

    B();
};

```

- Header files, especially those defining a “style”, should only use the absolute minimum number of include files and **must not** contain any `using` statements. Typically that would be only the header for the base class. Instead any include statements should be put into the corresponding implementation files and forward declarations be used. For implementation files, the “include what you use” principle should be employed. However, there is the notable exception that when the `pointers.h` header is included (or one of the base classes derived from it) certain headers will always be included and thus do not need to be explicitly specified. These are: `mpi.h`, `cstddef`, `cstdio`, `cstdlib`, `string`, `utils.h`, `vector`, `fmt/format.h`, `climits`, `cinttypes`. This also means any such file can assume that `FILE`, `NULL`, and `INT_MAX` are defined.
- Header files that define a new LAMMPS style (i.e. that have a `SomeStyle(some/name, SomeName);` macro in them) should only use the include file for the base class and otherwise use forward declarations and pointers; when interfacing to a library use the PIMPL (pointer to implementation) approach where you have a pointer to a struct that contains all library specific data (and thus requires the library header) but use a forward declaration and define the struct only in the implementation file. This is a **strict** requirement since this is where type clashes between packages and hard to find bugs have regularly manifested in the past.
- Please use `clang-format` only to reformat files that you have contributed. For header files containing a `SomeStyle(keyword, ClassName)` macros it is required to have this macro embedded with a pair of `// clang-format off, // clang-format on` comments and the line must be terminated with a semi-colon (;). Example:

```

#ifdef COMMAND_CLASS
// clang-format off
CommandStyle(run, Run);
// clang-format on
#else

#ifdef LMP_RUN_H
[...]
```

You may also use `// clang-format on/off` throughout your files to protect individual sections from being reformatted.

- We rarely accept new styles in the core `src` folder. Thus please review the list of [available Packages](#) to see if your contribution could be added to be added to one of them. It should fit into the general purposed of that package. If it does not fit well, it may be added to one of the EXTRA- packages or the MISC package.

### 14.3.9 Contributing a package

If your contribution has several related features that are not covered by one of the existing packages or is dependent on a library (bundled or external), it is best to make it a package directory with a name like FOO. In addition to your new files, the directory should contain a README text file. The README should contain your name and contact information and a brief description of what your new package does.

#### 14.3.10 Build system (strongly preferred)

LAMMPS currently supports two build systems: one that is based on *traditional Makefiles* and one that is based on *CMake*. Thus your contribution must be compatible with and support both.

For a single pair of header and implementation files that are an independent feature, it is usually only required to add them to `src/.gitignore`.

For traditional make, if your contributed files or package depend on other LAMMPS style files or packages also being installed (e.g. because your file is a derived class from the other LAMMPS class), then an `Install.sh` file is also needed to check for those dependencies and modifications to `src/Depend.sh` to trigger the checks. See other README and `Install.sh` files in other directories as examples.

Similarly for CMake support, changes may need to be made to `cmake/CMakeLists.txt`, some of the files in `cmake/presets`, and possibly a file with specific instructions needs to be added to `cmake/Modules/Packages/`. Please check out how this is handled for existing packages and ask the LAMMPS developers if you need assistance.

#### 14.3.11 Citation reminder (suggested)

If there is a paper of yours describing your feature (either the algorithm/science behind the feature itself, or its initial usage, or its implementation in LAMMPS), you can add the citation to the `*.cpp` source file. See `src/DIFFRACTION/compute_saed.cpp` for an example. A BibTeX format citation is stored in a string variable at the top of the file and a single line of code registering this variable is added to the constructor of the class. When your feature is used, by default, LAMMPS will print the brief info and the DOI in the first line to the screen and the full citation to the log file.

If there is additional functionality (which may have been added later) described in a different publication, additional citation descriptions may be added for as long as they are only registered when the corresponding keyword activating this functionality is used. With these options it is possible to have LAMMPS output a specific citation reminder whenever a user invokes your feature from their input script. Please note that you should *only* use this for the *most* relevant paper for a feature and a publication that you or your group authored. E.g. adding a citation in the code for a paper by Nose and Hoover if you write a fix that implements their integrator is not the intended usage. That latter kind of citation should just be included in the documentation page you provide describing your contribution. If you are not sure what the best option would be, please contact the LAMMPS developers for advice.

#### 14.3.12 Testing (optional)

If your contribution contains new utility functions or a supporting class (i.e. anything that does not depend on a LAMMPS object), new unit tests should be added to a suitable folder in the `unittest` tree. When adding a new LAMMPS style computing forces or selected fixes, a `.yaml` file with a test configuration and reference data should be added for the styles where a suitable tester program already exists (e.g. pair styles, bond styles, etc.). Please see *this section in the manual* for more information on how to enable, run, and expand testing.

## 14.4 Atom styles

Classes that define an *atom style* are derived from the AtomVec class and managed by the Atom class. The atom style determines what attributes are associated with an atom and communicated when it is a ghost atom or migrates to a new processor. A new atom style can be created if one of the existing atom styles does not define all the attributes you need to store and communicate with atoms.

The file `atom_vec_atomic.cpp` is the simplest example of an atom style. Examining the code for others will make these instructions more clear.

Note that the *atom style hybrid* command can be used to define atoms or particles which have the union of properties of individual styles. Also the *fix property/atom* command can be used to add a single property (e.g. charge or a molecule ID) to a style that does not have it. It can also be used to add custom properties to an atom, with options to communicate them with ghost atoms or read them from a data file. Other LAMMPS commands can access these custom properties, as can new pair, fix, compute styles that are written to work with these properties. For example, the *set* command can be used to set the values of custom per-atom properties from an input script. All of these methods are less work than writing and testing(!) code for a new atom style.

If you follow these directions your new style will automatically work in tandem with others via the *atom\_style hybrid* command.

The first step is to define a set of string lists in the constructor of the new derived class. Each list will have zero or more comma-separated strings that correspond to the variable names used in the `atom.h` header file for per-atom properties. Note that some represent per-atom vectors (q, molecule) while other are per-atom arrays (x,v). For all but the last two lists you do not need to specify any of (id,type,x,v,f). Those are included automatically as needed in the other lists.

<code>fields_grow</code>	full list of properties which is allocated and stored
<code>fields_copy</code>	list of properties to copy atoms are rearranged on-processor
<code>fields_comm</code>	list of properties communicated to ghost atoms every step
<code>fields_comm_vel</code>	additional properties communicated if <i>comm_modify vel</i> is used
<code>fields_reverse</code>	list of properties summed from ghost atoms every step
<code>fields_border</code>	list of properties communicated with ghost atoms every reneighboring step
<code>fields_border_vel</code>	additional properties communicated if <i>comm_modify vel</i> is used
<code>fields_exchange</code>	list of properties communicated when an atom migrates to another processor
<code>fields_restart</code>	list of properties written/read to/from a restart file
<code>fields_create</code>	list of properties defined when an atom is created by <i>create_atoms</i>
<code>fields_data_atom</code>	list of properties (in order) in the Atoms section of a data file, as read by <i>read_data</i>
<code>fields_data_vel</code>	list of properties (in order) in the Velocities section of a data file, as read by <i>read_data</i>

In these lists you can list variable names which LAMMPS already defines (in some other atom style), or you can create new variable names. You should not re-use a LAMMPS variable in your atom style that is used for something with a different meaning in another atom style. If the meaning is related, but interpreted differently by your atom style, then using the same variable name means a user must not use your style and the other style together in a *atom\_style hybrid* command. Because there will only be one value of the variable and different parts of LAMMPS will then likely use it differently. LAMMPS has no way of checking for this.

If you are defining new variable names then make them descriptive and unique to your new atom style. For example choosing “e” for energy is a bad choice; it is too generic. A better choice would be “e\_foo”, where “foo” is specific to your style.

If any of the variable names in your new atom style do not exist in LAMMPS, you need to add them to the `src/atom.h` and `atom.cpp` files.

Search for the word “customize” or “customization” in these 2 files to see where to add your variable. Adding a flag to the 2nd customization section in `atom.h` is only necessary if your code (e.g. a pair style) needs to check that a per-atom property is defined. These flags should also be set in the constructor of the atom style child class.

In `atom.cpp`, aside from the constructor and destructor, there are 3 methods that a new variable name or flag needs to be added to.

In `Atom::peratom_create()` when using the `Atom::add_peratom()` method, a `cols` argument of 0 is for per-atom vectors, a length > 1 is for per-atom arrays. Note the use of the extra per-thread flag and the `add_peratom_vary()` method when last dimension of the array is variable-length.

Adding the variable name to `Atom::extract()` enables the per-atom data to be accessed through the *LAMMPS library interface* by a calling code, including from *Python*.

The constructor of the new atom style will also typically set a few flags which are defined at the top of `atom_vec.h`. If these are unclear, see how other atom styles use them.

The `grow_pointers()` method is also required to make a copy of peratom data pointers, as explained in the code.

There are a number of other optional methods which your atom style can implement. These are only needed if you need to do something out-of-the-ordinary which the default operation of the `AtomVec` parent class does not take care of. The best way to figure out why they are sometimes useful is to look at how other atom styles use them.

- `process_args` = use if the atom style has arguments
- `init` = called before each run
- `force_clear` = called before force computations each timestep

A few atom styles define “bonus” data associated with some or all of their particles, such as *atom\_style ellipsoid* or *tri*. These methods work with that data:

- `copy_bonus`
- `clear_bonus`
- `pack_comm_bonus`
- `unpack_comm_bonus`
- `pack_border_bonus`
- `unpack_border_bonus`
- `pack_exchange_bonus`
- `unpack_exchange_bonus`
- `size_restart_bonus`
- `pack_restart_bonus`
- `unpack_restart_bonus`
- `data_atom_bonus`
- `memory_usage_bonus`

The *atom\_style body* command can define a particle geometry with an arbitrary number of values. This method reads it from a data file:

- `data_body`

These methods are called before or after operations handled by the parent `AtomVec` class. They allow an atom style to do customized operations on the per-atom values. For example *atom\_style sphere* reads a diameter and density of each particle from a data file. But these need to be converted internally to a radius and mass. That operation is done in the `data_atom_post()` method.

- `pack_restart_pre`
- `pack_restart_post`

- `unpack_restart_init`
- `create_atom_post`
- `data_atom_post`
- `pack_data_pre`
- `pack_data_post`

These methods enable the *compute property/atom* command to access per-atom variables it does not already define as arguments, so that they can be written to a dump file or used by other LAMMPS commands.

- `property_atom`
- `pack_property_atom`

## 14.5 Pair styles

Classes that compute pairwise interactions are derived from the Pair class. In LAMMPS, pairwise calculation include many-body potentials such as EAM or Tersoff where particles interact without a static bond topology. New styles can be created to add new pair potentials to LAMMPS.

`Pair_lj_cut.cpp` is a simple example of a Pair class, though it includes some optional methods to enable its use with rRESPA.

Here is a brief description of the class methods in `pair.h`:

<code>compute</code>	workhorse routine that computes pairwise interactions
<code>settings</code>	reads the input script line with arguments you define
<code>coeff</code>	set coefficients for one i,j type pair
<code>init_one</code>	perform initialization for one i,j type pair
<code>init_style</code>	initialization specific to this pair style
<code>write &amp; read_restart</code>	write/read i,j pair coeffs to restart files
<code>write &amp; read_restart_settings</code>	write/read global settings to restart files
<code>single</code>	force/r and energy of a single pairwise interaction between 2 atoms
<code>compute_inner/middle/outer</code>	versions of compute used by rRESPA

The inner/middle/outer routines are optional.

## 14.6 Bond, angle, dihedral, improper styles

Classes that compute molecular interactions are derived from the Bond, Angle, Dihedral, and Improper classes. New styles can be created to add new potentials to LAMMPS.

`Bond_harmonic.cpp` is the simplest example of a bond style. Ditto for the harmonic forms of the angle, dihedral, and improper style commands.

Here is a brief description of common methods you define in your new derived class. See `bond.h`, `angle.h`, `dihedral.h`, and `improper.h` for details and specific additional methods.



init	check if all coefficients are set, calls <i>init_style</i> (optional)
init_style	check if style specific conditions are met (optional)
compute	compute the molecular interactions (required)
settings	apply global settings for all types (optional)
coeff	set coefficients for one type (required)
equilibrium_distance	length of bond, used by SHAKE (required, bond only)
equilibrium_angle	opening of angle, used by SHAKE (required, angle only)
write & read_restart	writes/reads coeffs to restart files (required)
single	force (bond only) and energy of a single bond or angle (required, bond or angle only)
memory_usage	tally memory allocated by the style (optional)

## 14.7 Compute styles

Classes that compute scalar and vector quantities like temperature and the pressure tensor, as well as classes that compute per-atom quantities like kinetic energy and the centro-symmetry parameter are derived from the Compute class. New styles can be created to add new calculations to LAMMPS.

Compute\_temp.cpp is a simple example of computing a scalar temperature. Compute\_ke\_atom.cpp is a simple example of computing per-atom kinetic energy.

Here is a brief description of methods you define in your new derived class. See compute.h for details.

init	perform one time setup (required)
init_list	neighbor list setup, if needed (optional)
compute_scalar	compute a scalar quantity (optional)
compute_vector	compute a vector of quantities (optional)
compute_peratom	compute one or more quantities per atom (optional)
compute_local	compute one or more quantities per processor (optional)
pack_comm	pack a buffer with items to communicate (optional)
unpack_comm	unpack the buffer (optional)
pack_reverse	pack a buffer with items to reverse communicate (optional)
unpack_reverse	unpack the buffer (optional)
remove_bias	remove velocity bias from one atom (optional)
remove_bias_all	remove velocity bias from all atoms in group (optional)
restore_bias	restore velocity bias for one atom after remove_bias (optional)
restore_bias_all	same as before, but for all atoms in group (optional)
pair_tally_callback	callback function for <i>tally</i> -style computes (optional).
memory_usage	tally memory usage (optional)

Tally-style computes are a special case, as their computation is done in two stages: the callback function is registered with the pair style and then called from the Pair::ev\_tally() function, which is called for each pair after force and energy has been computed for this pair. Then the tallied values are retrieved with the standard compute\_scalar or compute\_vector or compute\_peratom methods. The *compute styles in the TALLY package* provide *examples* for utilizing this mechanism.

## 14.8 Fix styles

In LAMMPS, a “fix” is any operation that is computed during timestepping that alters some property of the system. Essentially everything that happens during a simulation besides force computation, neighbor list construction, and output, is a “fix”. This includes time integration (update of coordinates and velocities), force constraints or boundary conditions (SHAKE or walls), and diagnostics (compute a diffusion coefficient). New styles can be created to add new options to LAMMPS.

Fix\_setforce.cpp is a simple example of setting forces on atoms to prescribed values. There are dozens of fix options already in LAMMPS; choose one as a template that is similar to what you want to implement.

Here is a brief description of methods you can define in your new derived class. See fix.h for details.

setmask	determines when the fix is called during the timestep (required)
init	initialization before a run (optional)
init_list	store pointer to neighbor list; called by neighbor list code (optional)
setup_pre_exchange	called before atom exchange in setup (optional)
setup_pre_force	called before force computation in setup (optional)
setup	called immediately before the first timestep and after forces are computed (optional)
min_setup_pre_force	like setup_pre_force, but for minimizations instead of MD runs (optional)
min_setup	like setup, but for minimizations instead of MD runs (optional)
initial_integrate	called at very beginning of each timestep (optional)
pre_exchange	called before atom exchange on re-neighboring steps (optional)
pre_neighbor	called before neighbor list build (optional)
pre_force	called before pair & molecular forces are computed (optional)
post_force	called after pair & molecular forces are computed and communicated (optional)
final_integrate	called at end of each timestep (optional)
end_of_step	called at very end of timestep (optional)
write_restart	dumps fix info to restart file (optional)
restart	uses info from restart file to re-initialize the fix (optional)
grow_arrays	allocate memory for atom-based arrays used by fix (optional)
copy_arrays	copy atom info when an atom migrates to a new processor (optional)
pack_exchange	store atom’s data in a buffer (optional)
unpack_exchange	retrieve atom’s data from a buffer (optional)
pack_restart	store atom’s data for writing to restart file (optional)
unpack_restart	retrieve atom’s data from a restart file buffer (optional)
size_restart	size of atom’s data (optional)
maxsize_restart	max size of atom’s data (optional)
setup_pre_force_respa	same as setup_pre_force, but for rRESPA (optional)
initial_integrate_respa	same as initial_integrate, but for rRESPA (optional)
post_integrate_respa	called after the first half integration step is done in rRESPA (optional)
pre_force_respa	same as pre_force, but for rRESPA (optional)
post_force_respa	same as post_force, but for rRESPA (optional)
final_integrate_respa	same as final_integrate, but for rRESPA (optional)
min_pre_force	called after pair & molecular forces are computed in minimizer (optional)
min_post_force	called after pair & molecular forces are computed and communicated in minimizer (optional)
min_store	store extra data for linesearch based minimization on a LIFO stack (optional)
min_pushstore	push the minimization LIFO stack one element down (optional)
min_popstore	pop the minimization LIFO stack one element up (optional)
min_clearstore	clear minimization LIFO stack (optional)
min_step	reset or move forward on line search minimization (optional)
min_dof	report number of degrees of freedom <i>added</i> by this fix in minimization (optional)

continues on next page

Table 1 – continued from previous page

max_alpha	report maximum allowed step size during linesearch minimization (optional)
pack_comm	pack a buffer to communicate a per-atom quantity (optional)
unpack_comm	unpack a buffer to communicate a per-atom quantity (optional)
pack_reverse_comm	pack a buffer to reverse communicate a per-atom quantity (optional)
unpack_reverse_comm	unpack a buffer to reverse communicate a per-atom quantity (optional)
dof	report number of degrees of freedom <i>removed</i> by this fix during MD (optional)
compute_scalar	return a global scalar property that the fix computes (optional)
compute_vector	return a component of a vector property that the fix computes (optional)
compute_array	return a component of an array property that the fix computes (optional)
deform	called when the box size is changed (optional)
reset_target	called when a change of the target temperature is requested during a run (optional)
reset_dt	is called when a change of the time step is requested during a run (optional)
modify_param	called when a fix_modify request is executed (optional)
memory_usage	report memory used by fix (optional)
thermo	compute quantities for thermodynamic output (optional)

Typically, only a small fraction of these methods are defined for a particular fix. Setmask is mandatory, as it determines when the fix will be invoked during the timestep. Fixes that perform time integration (*nve*, *nvt*, *npt*) implement `initial_integrate()` and `final_integrate()` to perform velocity Verlet updates. Fixes that constrain forces implement `post_force()`.

Fixes that perform diagnostics typically implement `end_of_step()`. For an `end_of_step` fix, one of your fix arguments must be the variable “nevery” which is used to determine when to call the fix and you must set this variable in the constructor of your fix. By convention, this is the first argument the fix defines (after the ID, group-ID, style).

If the fix needs to store information for each atom that persists from timestep to timestep, it can manage that memory and migrate the info with the atoms as they move from processors to processor by implementing the `grow_arrays`, `copy_arrays`, `pack_exchange`, and `unpack_exchange` methods. Similarly, the `pack_restart` and `unpack_restart` methods can be implemented to store information about the fix in restart files. If you wish an integrator or force constraint fix to work with rRESPA (see the `run_style` command), the `initial_integrate`, `post_force_integrate`, and `final_integrate_respa` methods can be implemented. The `thermo` method enables a fix to contribute values to thermodynamic output, as printed quantities and/or to be summed to the potential energy of the system.

## 14.9 Input script command style

New commands can be added to LAMMPS input scripts by adding new classes that are derived from the `Command` class and thus must have a “command” method. For example, the `create_atoms`, `read_data`, `velocity`, and `run` commands are all implemented in this fashion. When such a command is encountered in the LAMMPS input script, LAMMPS simply creates a class instance with the corresponding name, invokes the “command” method of the class, and passes it the arguments from the input script. The command method can perform whatever operations it wishes on LAMMPS data structures.

The single method your new class must define is as follows:

command	operations performed by the new command
---------	---

Of course, the new class can define other methods and variables as needed.

## 14.10 Dump styles

Classes that dump per-atom info to files are derived from the Dump class. To dump new quantities or in a new format, a new derived dump class can be added, but it is typically simpler to modify the DumpCustom class contained in the dump\_custom.cpp file.

Dump\_atom.cpp is a simple example of a derived dump class.

Here is a brief description of methods you define in your new derived class. See dump.h for details.

write_header	write the header section of a snapshot of atoms
count	count the number of lines a processor will output
pack	pack a proc's output data into a buffer
write_data	write a proc's data to a file

See the *dump* command and its *custom* style for a list of keywords for atom information that can already be dumped by DumpCustom. It includes options to dump per-atom info from Compute classes, so adding a new derived Compute class is one way to calculate new quantities to dump.

Note that new keywords for atom properties are not typically added to the *dump custom* command. Instead they are added to the *compute property/atom* command.

## 14.11 Kspace styles

Classes that compute long-range Coulombic interactions via K-space representations (Ewald, PPPM) are derived from the KSpace class. New styles can be created to add new K-space options to LAMMPS.

Ewald.cpp is an example of computing K-space interactions.

Here is a brief description of methods you define in your new derived class. See kspace.h for details.

init	initialize the calculation before a run
setup	computation before the first timestep of a run
compute	every-timestep computation
memory_usage	tally of memory usage

## 14.12 Minimization styles

Classes that perform energy minimization derived from the Min class. New styles can be created to add new minimization algorithms to LAMMPS.

Min\_cg.cpp is an example of conjugate gradient minimization.

Here is a brief description of methods you define in your new derived class. See min.h for details.

init	initialize the minimization before a run
run	perform the minimization
memory_usage	tally of memory usage

## 14.13 Region styles

Classes that define geometric regions are derived from the Region class. Regions are used elsewhere in LAMMPS to group atoms, delete atoms to create a void, insert atoms in a specified region, etc. New styles can be created to add new region shapes to LAMMPS.

Region\_sphere.cpp is an example of a spherical region.

Here is a brief description of methods you define in your new derived class. See region.h for details.

inside	determine whether a point is in the region
surface_interior	determine if a point is within a cutoff distance inside of surface
surface_exterior	determine if a point is within a cutoff distance outside of surface
shape_update	change region shape if set by time-dependent variable

## 14.14 Body styles

Classes that define body particles are derived from the Body class. Body particles can represent complex entities, such as surface meshes of discrete points, collections of sub-particles, deformable objects, etc.

See the [Howto body](#) page for an overview of using body particles and the various body styles LAMMPS supports. New styles can be created to add new kinds of body particles to LAMMPS.

Body\_nparticle.cpp is an example of a body particle that is treated as a rigid body containing N sub-particles.

Here is a brief description of methods you define in your new derived class. See body.h for details.

data_body	process a line from the Bodies section of a data file
noutrow	number of sub-particles output is generated for
noutcol	number of values per-sub-particle output is generated for
output	output values for the Mth sub-particle
pack_comm_body	body attributes to communicate every timestep
unpack_comm_body	unpacking of those attributes
pack_border_body	body attributes to communicate when reneighboring is done
unpack_border_body	unpacking of those attributes

## 14.15 Thermodynamic output options

The Thermo class computes and prints thermodynamic information to the screen and log file; see the files thermo.cpp and thermo.h.

There are four styles defined in thermo.cpp: “one”, “multi”, “yaml”, and “custom”. The “custom” style allows the user to explicitly list keywords for individual quantities to print when thermodynamic output is generated. The others have a fixed list of keywords. See the [thermo\\_style](#) command for a list of available quantities. The formatting of the “custom” style defaults to the “one” style, but can be adapted using [thermo\\_modify line](#).

The thermo styles (one, multi, etc) are defined by lists of keywords with associated formats for integer and floating point numbers and identified but an enumerator constant. Adding a new style thus mostly requires defining a new list of keywords and the associated formats and then inserting the required output processing where the enumerators are identified. Search for the word “CUSTOMIZATION” with references to “thermo style” in the thermo.cpp file to see the locations where code will need to be added. The member function Thermo::header() prints output at the very beginning of a thermodynamic output block and can be used to print column headers or other front matter. The

member function `Thermo::footer()` prints output at the end of a thermodynamic output block. The formatting of the output is done by assembling a “line” (which may span multiple lines if the style inserts newline characters (“n” as in the “multi” style).

New thermodynamic keywords can also be added to `thermo.cpp` to compute new quantities for output. Search for the word “CUSTOMIZATION” with references to “keyword” in `thermo.cpp` to see the several locations where code will need to be added. Effectively, you need to define a member function that computes the property, add an if statement in `Thermo::parse_fields()` where the corresponding header string for the keyword and the function pointer is registered by calling the `Thermo::addfield()` method, and add an if statement in `Thermo::evaluate_keyword()` which is called from the `Variable` class when a thermo keyword is encountered.

---

**Note:** The third argument to `Thermo::addfield()` is a flag indicating whether the function for the keyword computes a floating point (FLOAT), regular integer (INT), or big integer (BIGINT) value. This information is used for formatting the thermodynamic output. Inside the function the result must then be stored either in the `dvalue`, `ivalue` or `bivalue` member variable, respectively.

---

Since the `thermo_style custom` command allows to use output of quantities calculated by `fixes`, `computes`, and `variables`, it may often be simpler to compute what you wish via one of those constructs, rather than by adding a new keyword to the `thermo_style` command.

## 14.16 Variable options

The `Variable` class computes and stores *variable* information in LAMMPS; see the file `variable.cpp`. The value associated with a variable can be periodically printed to the screen via the `print`, `fix print`, or `thermo_style custom` commands. Variables of style “equal” can compute complex equations that involve the following types of arguments:

```
thermo keywords = ke, vol, atoms, ...
other variables = v_a, v_myvar, ...
math functions = div(x,y), mult(x,y), add(x,y), ...
group functions = mass(group), xcm(group,x), ...
atom values = x[123], y[3], vx[34], ...
compute values = c_mytemp[0], c_thermo_press[3], ...
```

Adding keywords for the `thermo_style custom` command (which can then be accessed by variables) is discussed in the [Modify thermo](#) documentation.

Adding a new math function of one or two arguments can be done by editing one section of the `Variable::evaluate()` method. Search for the word “customize” to find the appropriate location.

Adding a new group function can be done by editing one section of the `Variable::evaluate()` method. Search for the word “customize” to find the appropriate location. You may need to add a new method to the `Group` class as well (see the `group.cpp` file).

Accessing a new atom-based vector can be done by editing one section of the `Variable::evaluate()` method. Search for the word “customize” to find the appropriate location.

Adding new *compute styles* (whose calculated values can then be accessed by variables) is discussed in the [Modify compute](#) documentation.

## INFORMATION FOR DEVELOPERS

This section describes the internal structure and basic algorithms of the LAMMPS code. This is a work in progress and additional information will be added incrementally depending on availability of time and requests from the LAMMPS user community.

### 15.1 Source files

The source files of the LAMMPS code are found in two directories of the distribution: `src` and `lib`. Most of the code is written in C++ but there are small a number of files in several other languages like C, Fortran, Shell script, or Python.

The core of the code is located in the `src` folder and its sub-directories. A sizable number of these files are in the `src` directory itself, but there are plenty of *packages*, which can be included or excluded when LAMMPS is built. See the *Include packages in build* section of the manual for more information about that part of the build process. LAMMPS currently supports building with *conventional makefiles* and through *CMake*. Those procedures differ in how packages are enabled or disabled for inclusion into a LAMMPS binary so they cannot be mixed. The source files for each package are in all-uppercase sub-directories of the `src` folder, for example `src/MOLECULE` or `src/EXTRA-MOLECULE`. The `src/STUBS` sub-directory is not a package but contains a dummy MPI library, that is used when building a serial version of the code. The `src/MAKE` directory and its sub-directories contain makefiles with settings and flags for a variety of configuration and machines for the build process with traditional makefiles.

The `lib` directory contains the source code for several supporting libraries or files with configuration settings to use globally installed libraries, that are required by some of the optional packages. They may include python scripts that can transparently download additional source code on request. Each sub-directory, like `lib/poems` or `lib/gpu`, contains the source files, some of which are in different languages such as Fortran or CUDA. These libraries included in the LAMMPS build, if the corresponding package is installed.

LAMMPS C++ source files almost always come in pairs, such as `src/run.cpp` (implementation file) and `src/run.h` (header file). Each pair of files defines a C++ class, for example the `LAMMPS_NS::Run` class which contains the code invoked by the `run` command in a LAMMPS input script. As this example illustrates, source file and class names often have a one-to-one correspondence with a command used in a LAMMPS input script. Some source files and classes do not have a corresponding input script command, e.g. `src/force.cpp` and the `LAMMPS_NS::Force` class. They are discussed in the next section.

The names of all source files are in lower case and may use the underscore character ‘`_`’ to separate words. Outside of bundled libraries which may have different conventions, all C and C++ header files have a `.h` extension, all C++ files have a `.cpp` extension, and C files a `.c` extension. A small number of C++ classes and utility functions are implemented with only a `.h` file. Examples are the Pointers and Commands classes or the MathVec functions.

## 15.2 Class topology

Though LAMMPS has a lot of source files and classes, its class topology is not very deep, which can be seen from the *LAMMPS class topology* figure. In that figure, each name refers to a class and has a pair of associated source files in the `src` folder, for example the class `LAMMPS_NS::Memory` corresponds to the files `memory.cpp` and `memory.h`, or the class `LAMMPS_NS::AtomVec` corresponds to the files `atom_vec.cpp` and `atom_vec.h`. Full lines in the figure represent compositing: that is the class at the base of the arrow holds a pointer to an instance of the class at the tip. Dashed lines instead represent inheritance: the class to the tip of the arrow is derived from the class at the base. Classes with a red boundary are not instantiated directly, but they represent the base classes for “styles”. Those “styles” make up the bulk of the LAMMPS code and only a few representative examples are included in the figure so it remains readable.

The `LAMMPS_NS::LAMMPS` class is the topmost class and represents what is generally referred to an “instance” of LAMMPS. It is a composite holding pointers to instances of other core classes providing the core functionality of the MD engine in LAMMPS and through them abstractions of the required operations. The constructor of the LAMMPS class will instantiate those instances, process the command line flags, initialize MPI (if not already done) and set up file pointers for input and output. The destructor will shut everything down and free all associated memory. Thus code for the standalone LAMMPS executable in `main.cpp` simply initializes MPI, instantiates a single instance of LAMMPS while passing it the command line flags and input script. It deletes the LAMMPS instance after the method reading the input returns and shuts down the MPI environment before it exits the executable.

The `LAMMPS_NS::Pointers` is not shown in the *LAMMPS class topology* figure for clarity. It holds references to many of the members of the `LAMMPS_NS::LAMMPS`, so that all classes derived from `LAMMPS_NS::Pointers` have direct access to those reference. From the class topology all classes with blue boundary are referenced in the Pointers class and all classes in the second and third columns, that are not listed as derived classes are instead derived from `LAMMPS_NS::Pointers`. To initialize the pointer references in Pointers, a pointer to the LAMMPS class instance needs to be passed to the constructor and thus all constructors for classes derived from it must do so and pass this pointer to the constructor for Pointers.

Since all storage is supposed to be encapsulated (there are a few exceptions), the LAMMPS class can also be instantiated multiple times by a calling code. Outside of the aforementioned exceptions, those LAMMPS instances can be used alternately. As of the time of this writing (early 2021) LAMMPS is not yet sufficiently thread-safe for concurrent execution. When running in parallel with MPI, care has to be taken, that suitable copies of communicators are used to not create conflicts between different instances.

The LAMMPS class currently (early 2021) holds instances of 19 classes representing the core functionality. There are a handful of virtual parent classes in LAMMPS that define what LAMMPS calls *styles*. They are shaded red in the *LAMMPS class topology* figure. Each of these are parents of a number of child classes that implement the interface defined by the parent class. There are two main categories of these *styles*: some may only have one instance active at a time (e.g. `atom`, `pair`, `bond`, `angle`, `dihedral`, `improper`, `kpace`, `comm`) and there is a dedicated pointer variable for each of them in the composite class. Setups that require a mix of different such styles have to use a *hybrid* class that takes the place of the one allowed instance and then manages and forwards calls to the corresponding sub-styles for the designated subset of atoms or data. The composite class may also have lists of class instances, e.g. `Modify` handles lists of compute and fix styles, while `Output` handles a list of dump class instances.

The exception to this scheme are the `command` style classes. These implement specific commands that can be invoked before, after, or in between runs. For these an instance of the class is created, its `command()` method called and then, after completion, the class instance deleted. Examples for this are the `create_box`, `create_atoms`, `minimize`, `run`, `set`, or `velocity` command styles.

For all those *styles* certain naming conventions are employed: for the `fix nve` command the class is called `FixNVE` and the source files are `fix_nve.h` and `fix_nve.cpp`. Similarly for `fix ave/time` we have `FixAveTime` and `fix_ave_time.h` and `fix_ave_time.cpp`. Style names are lower case and without spaces or special characters. A suffix or words are appended with a forward slash `'/'` which denotes a variant of the corresponding class without the suffix. To connect the style name and the class name, LAMMPS uses macros like: `AtomStyle()`, `PairStyle()`, `BondStyle()`, `RegionStyle()`, and so on in the corresponding header file. During configuration or compilation files



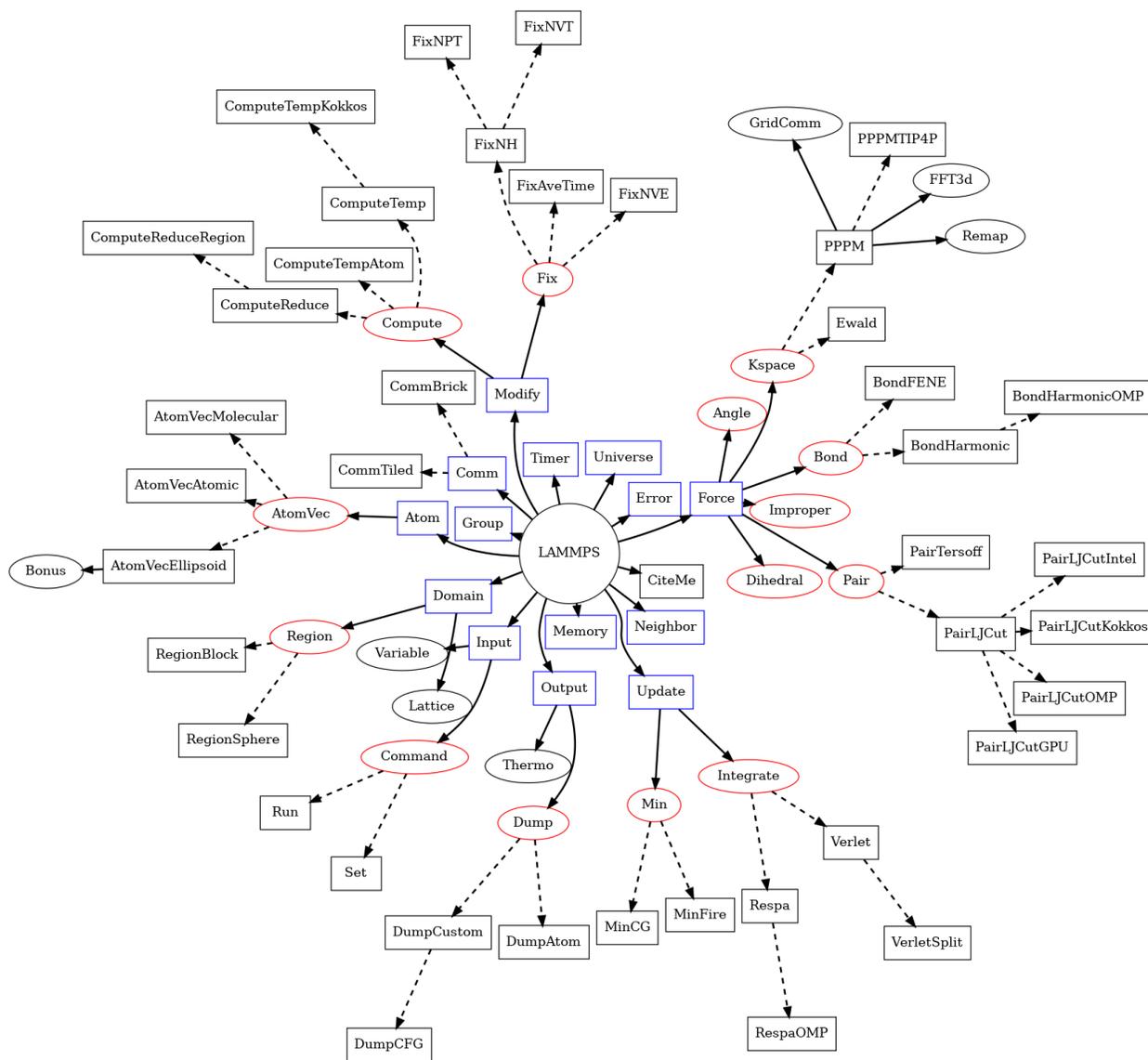


Fig. 1: LAMMPS class topology

This figure shows some of the relations of the base classes of the LAMMPS simulation package. Full lines indicate that a class holds an instance of the class it is pointing to; dashed lines point to derived classes that are given as examples of what classes may be instantiated during a LAMMPS run based on the input commands and accessed through the API define by their respective base classes. At the core is the *LAMMPS* class, which holds pointers to class instances with specific purposes. Those may hold instances of other classes, sometimes directly, or only temporarily, sometimes as derived classes or derived classes of derived classes, which may also hold instances of other classes.

with the pattern `style_<name>.h` are created that consist of a list of include statements including all headers of all styles of a given type that are currently active (or “installed”).

More details on individual classes in the *LAMMPS class topology* are as follows:

- The Memory class handles allocation of all large vectors and arrays.
- The Error class prints all (terminal) error and warning messages.
- The Universe class sets up one or more partitions of processors so that one or multiple simulations can be run, on the processors allocated for a run, e.g. by the `mpirun` command.
- The Input class reads and processes input strings and files, stores variables, and invokes *commands*.
- Command style classes are derived from the Command class. They provide input script commands that perform one-time operations before/after/between simulations or which invoke a simulation. They are usually instantiated from within the Input class, its `command` method invoked, and then immediately destructed.
- The Finish class is instantiated to print statistics to the screen after a simulation is performed, by commands like `run` and `minimize`.
- The Special class walks the bond topology of a molecular system to find first, second, third neighbors of each atom. It is invoked by several commands, like `read_data`, `read_restart`, or `replicate`.
- The Atom class stores per-atom properties associated with atom styles. More precisely, they are allocated and managed by a class derived from the AtomVec class, and the Atom class simply stores pointers to them. The classes derived from AtomVec represent the different atom styles and they are instantiated through the `atom_style` command.
- The Update class holds instances of an integrator and a minimizer class. The Integrate class is a parent style for the Verlet and r-RESPA time integrators, as defined by the `run_style` command. The Min class is a parent style for various energy minimizers.
- The Neighbor class builds and stores neighbor lists. The NeighList class stores a single list (for all atoms). A NeighRequest class instance is created by pair, fix, or compute styles when they need a particular kind of neighbor list and use the NeighRequest properties to select the neighbor list settings for the given request. There can be multiple instances of the NeighRequest class and the Neighbor class will try to optimize how they are computed by creating copies or sub-lists where possible.
- The Comm class performs inter-processor communication, typically of ghost atom information. This usually involves MPI message exchanges with 6 neighboring processors in the 3d logical grid of processors mapped to the simulation box. There are two *communication styles* enabling different ways to do the domain decomposition. Sometimes the Irregular class is used, when atoms may migrate to arbitrary processors.
- The Domain class stores the simulation box geometry, as well as geometric Regions and any user definition of a Lattice. The latter are defined by the `region` and `lattice` commands in an input script.
- The Force class computes various forces between atoms. The Pair parent class is for non-bonded or pairwise forces, which in LAMMPS also includes many-body forces such as the Tersoff 3-body potential if those are computed by walking pairwise neighbor lists. The Bond, Angle, Dihedral, Improper parent classes are styles for bonded interactions within a static molecular topology. The KSpace parent class is for computing long-range Coulombic interactions. One of its child classes, PPPM, uses the FFT3D and Remap classes to redistribute and communicate grid-based information across the parallel processors.
- The Modify class stores lists of class instances derived from the `Fix` and `Compute` base classes.
- The Group class manipulates groups that atoms are assigned to via the `group` command. It also has functions to compute various attributes of groups of atoms.
- The Output class is used to generate 3 kinds of output from a LAMMPS simulation: thermodynamic information printed to the screen and log file, dump file snapshots, and restart files. These correspond to the *Thermo*, *Dump*,

and *WriteRestart* classes respectively. The Dump class is a base class with several derived classes implementing various dump style variants.

- The Timer class logs timing information, output at the end of a run.

## 15.3 Code design

This section explains some of the code design choices in LAMMPS with the goal of helping developers write new code similar to the existing code. Please see the section on *Requirements for contributed code* for more specific recommendations and guidelines. While that section is organized more in the form of a checklist for code contributors, the focus here is on overall code design strategy, choices made between possible alternatives, and discussing some relevant C++ programming language constructs.

Historically, the basic design philosophy of the LAMMPS C++ code was a “C with classes” style. The motivation was to make it easy to modify LAMMPS for people without significant training in C++ programming. Data structures and code constructs were used that resemble the previous implementation(s) in Fortran. A contributing factor to this choice also was that at the time, C++ compilers were often not mature and some of the advanced features contained bugs or did not function as the standard required. There were also disagreements between compiler vendors as to how to interpret the C++ standard documents.

However, C++ compilers have now advanced significantly. In 2020 we decided to require the C++11 standard as the minimum C++ language standard for LAMMPS. Since then we have begun to also replace some of the C-style constructs with equivalent C++ functionality, either from the C++ standard library or as custom classes or functions, in order to improve readability of the code and to increase code reuse through abstraction of commonly used functionality.

---

**Note:** Please note that as of spring 2022 there is still a sizable chunk of legacy code in LAMMPS that has not yet been refactored to reflect these style conventions in full. LAMMPS has a large code base and many different contributors and there also is a hierarchy of precedence in which the code is adapted. Highest priority has been the code in the `src` folder, followed by code in packages in order of their popularity and complexity (simpler code is adapted sooner), followed by code in the `lib` folder. Source code that is downloaded from external packages or libraries during compilation is not subject to the conventions discussed here.

---

### 15.3.1 Object oriented code

LAMMPS is designed to be an object oriented code. Each simulation is represented by an instance of the LAMMPS class. When running in parallel each MPI process creates such an instance. This can be seen in the `main.cpp` file where the core steps of running a LAMMPS simulation are the following 3 lines of code:

```
LAMMPS *lammps = new LAMMPS(argc, argv, lammps_comm);
lammps->input->file();
delete lammps;
```

The first line creates a LAMMPS class instance and passes the command line arguments and the global communicator to its constructor. The second line triggers the LAMMPS instance to process the input (either from standard input or a provided input file) until the simulation ends. The third line deletes the LAMMPS instance. The remainder of the `main.cpp` file has code for error handling, MPI configuration, and other special features.

The basic LAMMPS class hierarchy which is created by the LAMMPS class constructor is shown in *LAMMPS class topology*. When input commands are processed, additional class instances are created, or deleted, or replaced. Likewise specific member functions of specific classes are called to trigger actions such as creating atoms, computing forces, computing properties, time-propagating the system, or writing output.

## Compositing and Inheritance

LAMMPS makes extensive use of the object oriented programming (OOP) principles of *compositing* and *inheritance*. Classes like the LAMMPS class are a **composite** containing pointers to instances of other classes like `Atom`, `Comm`, `Force`, `Neighbor`, `Modify`, and so on. Each of these classes implement certain functionality by storing and manipulating data related to the simulation and providing member functions that trigger certain actions. Some of those classes like `Force` are themselves composites, containing instances of classes describing different force interactions. Similarly the `Modify` class contains a list of `Fix` and `Compute` classes. If the input commands that correspond to these classes include the word *style*, then LAMMPS stores only a single instance of that class. E.g. *atom\_style*, *comm\_style*, *pair\_style*, *bond\_style*. If the input command does not include the word *style*, there can be many instances of that class defined. E.g. *region*, *fix*, *compute*, *dump*.

**Inheritance** enables creation of *derived* classes that can share common functionality in their base class while providing a consistent interface. The derived classes replace (dummy or pure) functions in the base class. The higher level classes can then call those methods of the instantiated classes without having to know which specific derived class variant was instantiated. In LAMMPS these derived classes are often referred to as “styles”, e.g. pair styles, fix styles, atom styles and so on.

This is the origin of the flexibility of LAMMPS. For example pair styles implement a variety of different non-bonded interatomic potentials functions. All details for the implementation of a potential are stored and executed in a single class.

As mentioned above, there can be multiple instances of classes derived from the `Fix` or `Compute` base classes. They represent a different facet of LAMMPS flexibility as they provide methods which can be called at different points in time within a timestep, as explained in *Developer\_flow*. This allows the input script to tailor how a specific simulation is run, what diagnostic computations are performed, and how the output of those computations is further processed or output.

Additional code sharing is possible by creating derived classes from the derived classes (e.g., to implement an accelerated version of a pair style) where only a subset of the derived class methods are replaced with accelerated versions.

## Polymorphism

Polymorphism and dynamic dispatch are another OOP feature that play an important role in how LAMMPS selects what code to execute. In a nutshell, this is a mechanism where the decision of which member function to call from a class is determined at runtime and not when the code is compiled. To enable it, the function has to be declared as `virtual` and all corresponding functions in derived classes should use the `override` property. Below is a brief example.

```
class Base {
public:
    virtual ~Base() = default;
    void call();
    void normal();
    virtual void poly();
};

void Base::call() {
    normal();
    poly();
}

class Derived : public Base {
public:
    ~Derived() override = default;
```

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```

void normal();
void poly() override;
};

// [...]

Base *base1 = new Base();
Base *base2 = new Derived();

base1->call();
base2->call();

```

The difference in behavior of the `normal()` and the `poly()` member functions is which of the two member functions is called when executing `base1->call()` versus `base2->call()`. Without polymorphism, a function within the base class can only call member functions within the same scope, that is `Base::call()` will always call `Base::normal()`. But for the `base2->call()` case the call of the virtual member function will be dispatched to `Derived::poly()` instead. This mechanism means that functions are called within the scope of the class type that was used to *create* the class instance are invoked; even if they are assigned to a pointer using the type of a base class. This is the desired behavior and this way LAMMPS can even use styles that are loaded at runtime from a shared object file with the *plugin command*.

A special case of virtual functions are so-called pure functions. These are virtual functions that are initialized to 0 in the class declaration (see example below).

```

class Base {
public:
    virtual void pure() = 0;
};

```

This has the effect that an instance of the base class cannot be created and that derived classes **must** implement these functions. Many of the functions listed with the various class styles in the section *Modifying & extending LAMMPS* are pure functions. The motivation for this is to define the interface or API of the functions but defer their implementation to the derived classes.

However, there are downsides to this. For example, calls to virtual functions from within a constructor, will not be in the scope of the derived class and thus it is good practice to either avoid calling them or to provide an explicit scope such as `Base::poly()` or `Derived::poly()`. Furthermore, any destructors in classes containing virtual functions should be declared virtual too, so they will be processed in the expected order before types are removed from dynamic dispatch.

---

### Important Notes

In order to be able to detect incompatibilities at compile time and to avoid unexpected behavior, it is crucial that all member functions that are intended to replace a virtual or pure function use the `override` property keyword. For the same reason, the use of overloads or default arguments for virtual functions should be avoided as they lead to confusion over which function is supposed to override which and which arguments need to be declared.

---

## Style Factories

In order to create class instances for different styles, LAMMPS often uses a programming pattern called *Factory*. Those are functions that create an instance of a specific derived class, say `PairLJCut` and return a pointer to the type of the common base class of that style, `Pair` in this case. To associate the factory function with the style keyword, an `std::map` class is used with function pointers indexed by their keyword (for example “lj/cut” for `PairLJCut` and “morse” for `PairMorse`). A couple of typedefs help keep the code readable and a template function is used to implement the actual factory functions for the individual classes. Below is an example of such a factory function from the `Force` class as declared in `force.h` and implemented in `force.cpp`. The file `style_pair.h` is generated during compilation and includes all main header files (i.e. those starting with `pair_`) of pair styles and then the macro `PairStyle()` will associate the style name “lj/cut” with a factory function creating an instance of the `PairLJCut` class.

```
// from force.h
typedef Pair *(*PairCreator)(LAMMPS *);
typedef std::map<std::string, PairCreator> PairCreatorMap;
PairCreatorMap *pair_map;

// from force.cpp
template <typename S, typename T> static S *style_creator(LAMMPS *lmp)
{
    return new T(lmp);
}

// [...]

pair_map = new PairCreatorMap();

#define PAIR_CLASS
#define PairStyle(key, Class) (*pair_map)[#key] = &style_creator<Pair, Class>;
#include "style_pair.h"
#undef PairStyle
#undef PAIR_CLASS

// from pair_lj_cut.h

#ifdef PAIR_CLASS
PairStyle(lj/cut,PairLJCut);
#else
// [...]
```

Similar code constructs are present in other files like `modify.cpp` and `modify.h` or `neighbor.cpp` and `neighbor.h`. Those contain similar macros and include `style_*.h` files for creating class instances of styles they manage.

## 15.3.2 I/O and output formatting

### C-style stdio versus C++ style iostreams

LAMMPS uses the “stdio” library of the standard C library for reading from and writing to files and console instead of C++ “iostreams”. This is mainly motivated by better performance, better control over formatting, and less effort to achieve specific formatting.

Since mixing “stdio” and “iostreams” can lead to unexpected behavior, use of the latter is strongly discouraged. Also output to the screen should not use the predefined `stdout` FILE pointer, but rather the `screen` and `logfile` FILE pointers managed by the LAMMPS class. Furthermore, output should generally only be done by MPI rank 0 (`comm->me == 0`). Output that is sent to both `screen` and `logfile` should use the `utils::logmesg()` convenience function.

We also discourage the use of stringstream because the bundled `{fmt}` library and the customized tokenizer classes can provide the same functionality in a cleaner way with better performance. This also helps maintain a consistent programming syntax with code from many different contributors.

### Formatting with the {fmt} library

The LAMMPS source code includes a copy of the `{fmt}` library which is preferred over formatting with the “printf()” family of functions. The primary reason is that it allows a typesafe default format for any type of supported data. This is particularly useful for formatting integers of a given size (32-bit or 64-bit) which may require different format strings depending on compile time settings or compilers/operating systems. Furthermore, `{fmt}` gives better performance, has more functionality, a familiar formatting syntax that has similarities to `format()` in Python, and provides a facility that can be used to integrate format strings and a variable number of arguments into custom functions in a much simpler way than the `varargs` mechanism of the C library. Finally, `{fmt}` has been included into the C++20 language standard, so changes to adopt it are future-proof.

Formatted strings are frequently created by calling the `fmt::format()` function which will return a string as a `std::string` class instance. In contrast to the `%` placeholder in `printf()`, the `{fmt}` library uses `{}` to embed format descriptors. In the simplest case, no additional characters are needed as `{fmt}` will choose the default format based on the data type of the argument. Otherwise the `fmt::print()` function may be used instead of `printf()` or `fprintf()`. In addition, several LAMMPS output functions, that originally accepted a single string as argument have been overloaded to accept a format string with optional arguments as well (e.g., `Error::all()`, `Error::one()`, `utils::logmesg()`).

### Summary of the {fmt} format syntax

The syntax of the format string is “`{[<argument id>]:<format spec>}`”, where either the argument id or the format spec (separated by a colon ‘:’) is optional. The argument id is usually a number starting from 0 that is the index to the arguments following the format string. By default these are assigned in order (i.e. 0, 1, 2, 3, 4 etc.). The most common case for using argument id would be to use the same argument in multiple places in the format string without having to provide it as an argument multiple times. In LAMMPS the argument id is rarely used.

More common is the use of a format specifier, which starts with a colon. This may optionally be followed by a fill character (default is ‘ ’). If provided, the fill character **must** be followed by an alignment character (‘<’, ‘^’, ‘>’ for left, centered, or right alignment (default)). The alignment character may be used without a fill character. The next important format parameter would be the minimum width, which may be followed by a dot ‘.’ and a precision for floating point numbers. The final character in the format string would be an indicator for the “presentation”, i.e. ‘d’ for decimal presentation of integers, ‘x’ for hexadecimal, ‘o’ for octal, ‘c’ for character etc. This mostly follows the “printf()” scheme but without requiring an additional length parameter to distinguish between different integer widths. The `{fmt}` library will detect those and adapt the formatting accordingly. For floating point numbers there are correspondingly, ‘g’ for generic presentation, ‘e’ for exponential presentation, and ‘f’ for fixed point presentation.

Thus “{:8}” would represent *any* type argument using at least 8 characters; “{:<8}” would do this as left aligned, “{:^8}” as centered, “{:>8}” as right aligned. If a specific presentation is selected, the argument type must be compatible or else the {fmt} formatting code will throw an exception. Some format string examples are given below:

```
auto mesg = fmt::format(" CPU time: {:4d}:{:02d}:{:02d}\n", cpuh, cpum, cpus);
mesg = fmt::format("{:<8s}| {:<10.5g} | {:<10.5g} | {:<10.5g} |{:6.1f} |{:6.2f}\n",
                  label, time_min, time, time_max, time_sq, tmp);
utils::logmesg(lmp, "{:>6} = max # of 1-2 neighbors\n", maxall);
utils::logmesg(lmp, "Lattice spacing in x,y,z = {:.8} {:.8} {:.8}\n",
               xlattice, ylattice, zlattice);
```

which will create the following output lines:

```
CPU time:    0:02:16
Pair      | 2.0133      | 2.0133      | 2.0133      | 0.0 | 84.21
      4 = max # of 1-2 neighbors
Lattice spacing in x,y,z = 1.6795962 1.6795962 1.6795962
```

Finally, a special feature of the {fmt} library is that format parameters like the width or the precision may be also provided as arguments. In that case a nested format is used where a pair of curly braces (with an optional argument id) “{ }” are used instead of the value, for example “{:{}d}” will consume two integer arguments, the first will be the value shown and the second the minimum width.

For more details and examples, please consult the [{fmt} syntax documentation website](#).

### 15.3.3 Memory management

Dynamical allocation of small data and objects can be done with the the C++ commands “new” and “delete/delete[]”. Large data should use the member functions of the `Memory` class, most commonly, `Memory::create()`, `Memory::grow()`, and `Memory::destroy()`, which provide variants for vectors, 2d arrays, 3d arrays, etc. These can also be used for small data.

The use of `malloc()`, `calloc()`, `realloc()` and `free()` directly is strongly discouraged. To simplify adapting legacy code into the LAMMPS code base the member functions `Memory::smalloc()`, `Memory::srealloc()`, and `Memory::sfree()` are available, which perform additional error checks for safety.

Use of these custom memory allocation functions is motivated by the following considerations:

- memory allocation failures on *any* MPI rank during a parallel run will trigger an immediate abort of the entire parallel calculation instead of stalling it
- a failing “new” will trigger an exception which is also captured by LAMMPS and triggers a global abort
- allocation of multi-dimensional arrays will be done in a C compatible fashion but so that the storage of the actual data is stored in one large contiguous block. Thus when MPI communication is needed, the data can be communicated directly (similar to Fortran arrays).
- the “destroy()” and “sfree()” functions may safely be called on NULL pointers
- the “destroy()” functions will nullify the pointer variables making “use after free” errors easy to detect
- it is possible to use a larger than default memory alignment (not on all operating systems, since the allocated storage pointers must be compatible with `free()` for technical reasons)

In the practical implementation of code this means that any pointer variables that are class members should be initialized to a `nullptr` value in their respective constructors. That way it is safe to call `Memory::destroy()` or `delete[]` on them before *any* allocation outside the constructor. This helps prevent memory leaks.



## 15.4 Parallel algorithms

LAMMPS is designed to enable running simulations in parallel using the MPI parallel communication standard with distributed data via domain decomposition. The parallelization aims to be efficient result in good strong scaling (= good speedup for the same system) and good weak scaling (= the computational cost of enlarging the system is proportional to the system size). Additional parallelization using GPUs or OpenMP can also be applied within the sub-domain assigned to an MPI process. For clarity, most of the following illustrations show the 2d simulation case. The underlying algorithms in those cases, however, apply to both 2d and 3d cases equally well.

---

**Note:** The text and most of the figures in this chapter were adapted for the manual from the section on parallel algorithms in the *new LAMMPS paper*.

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### 15.4.1 Partitioning

The underlying spatial decomposition strategy used by LAMMPS for distributed-memory parallelism is set with the *comm\_style* command and can be either “brick” (a regular grid) or “tiled”.

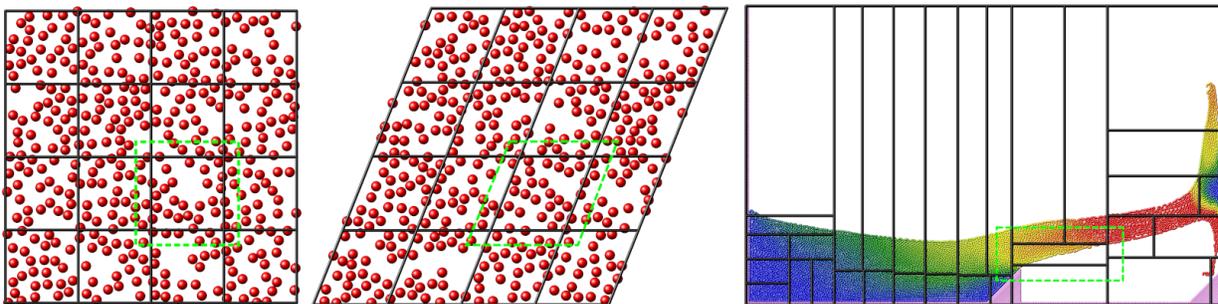


Fig. 2: domain decomposition

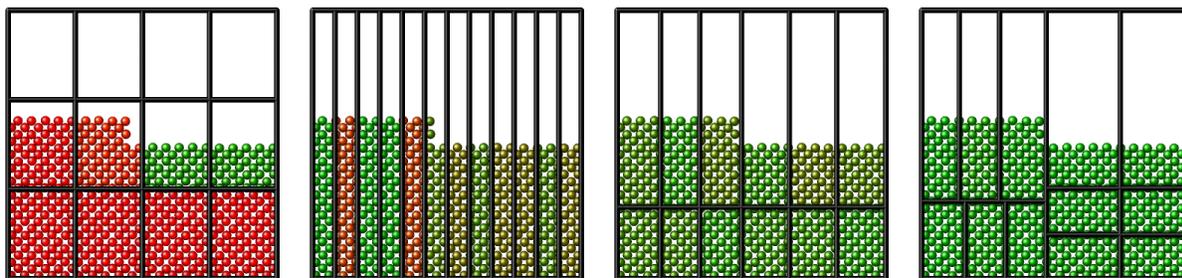
This figure shows the different kinds of domain decomposition used for MPI parallelization: “brick” on the left with an orthogonal (left) and a triclinic (middle) simulation domain, and a “tiled” decomposition (right). The black lines show the division into sub-domains and the contained atoms are “owned” by the corresponding MPI process. The green dashed lines indicate how sub-domains are extended with “ghost” atoms up to the communication cutoff distance.

The LAMMPS simulation box is a 3d or 2d volume, which can be orthogonal or triclinic in shape, as illustrated in the *domain decomposition* figure for the 2d case. Orthogonal means the box edges are aligned with the  $x$ ,  $y$ ,  $z$  Cartesian axes, and the box faces are thus all rectangular. Triclinic allows for a more general parallelepiped shape in which edges are aligned with three arbitrary vectors and the box faces are parallelograms. In each dimension box faces can be periodic, or non-periodic with fixed or shrink-wrapped boundaries. In the fixed case, atoms which move outside the face are deleted; shrink-wrapped means the position of the box face adjusts continuously to enclose all the atoms.

For distributed-memory MPI parallelism, the simulation box is spatially decomposed (partitioned) into non-overlapping sub-domains which fill the box. The default partitioning, “brick”, is most suitable when atom density is roughly uniform, as shown in the left-side images of the *domain decomposition* figure. The sub-domains comprise a regular grid and all sub-domains are identical in size and shape. Both the orthogonal and triclinic boxes can deform continuously during a simulation, e.g. to compress a solid or shear a liquid, in which case the processor sub-domains likewise deform.

For models with non-uniform density, the number of particles per processor can be load-imbalanced with the default partitioning. This reduces parallel efficiency, as the overall simulation rate is limited by the slowest processor, i.e. the one with the largest computational load. For such models, LAMMPS supports multiple strategies to reduce the load imbalance:

- The processor grid decomposition is by default based on the simulation cell volume and tries to optimize the volume to surface ratio for the sub-domains. This can be changed with the *processors command*.
- The parallel planes defining the size of the sub-domains can be shifted with the *balance command*. Which can be done in addition to choosing a more optimal processor grid.
- The recursive bisectioning algorithm in combination with the “tiled” communication style can produce a partitioning with equal numbers of particles in each sub-domain.



The pictures above demonstrate different decompositions for a 2d system with 12 MPI ranks. The atom colors indicate the load imbalance of each sub-domain with green being optimal and red the least optimal.

Due to the vacuum in the system, the default decomposition is unbalanced with several MPI ranks without atoms (left). By forcing a 1x12x1 processor grid, every MPI rank does computations now, but number of atoms per sub-domain is still uneven and the thin slice shape increases the amount of communication between sub-domains (center left). With a 2x6x1 processor grid and shifting the sub-domain divisions, the load imbalance is further reduced and the amount of communication required between sub-domains is less (center right). And using the recursive bisectioning leads to further improved decomposition (right).

## 15.4.2 Communication

Following the partitioning scheme in use all per-atom data is distributed across the MPI processes, which allows LAMMPS to handle very large systems provided it uses a correspondingly large number of MPI processes. Since the per-atom data (atom IDs, positions, velocities, types, etc.) To be able to compute the short-range interactions MPI processes need not only access to data of atoms they “own” but also information about atoms from neighboring sub-domains, in LAMMPS referred to as “ghost” atoms. These are copies of atoms storing required per-atom data for up to the communication cutoff distance. The green dashed-line boxes in the *domain decomposition* figure illustrate the extended ghost-atom sub-domain for one processor.

This approach is also used to implement periodic boundary conditions: atoms that lie within the cutoff distance across a periodic boundary are also stored as ghost atoms and taken from the periodic replication of the sub-domain, which may be the same sub-domain, e.g. if running in serial. As a consequence of this, force computation in LAMMPS is not subject to minimum image conventions and thus cutoffs may be larger than half the simulation domain.

Efficient communication patterns are needed to update the “ghost” atom data, since that needs to be done at every MD time step or minimization step. The diagrams of the *ghost-atom-comm* figure illustrate how ghost atom communication is performed in two stages for a 2d simulation (three in 3d) for both a regular and irregular partitioning of the simulation box. For the regular case (left) atoms are exchanged first in the *x*-direction, then in *y*, with four neighbors in the grid of processor sub-domains.

In the *x* stage, processor ranks 1 and 2 send owned atoms in their red-shaded regions to rank 0 (and vice versa). Then in the *y* stage, ranks 3 and 4 send atoms in their blue-shaded regions to rank 0, which includes ghost atoms they received in the *x* stage. Rank 0 thus acquires all its ghost atoms; atoms in the solid blue corner regions are communicated twice before rank 0 receives them.

For the irregular case (right) the two stages are similar, but a processor can have more than one neighbor in each direction. In the *x* stage, MPI ranks 1,2,3 send owned atoms in their red-shaded regions to rank 0 (and vice versa).

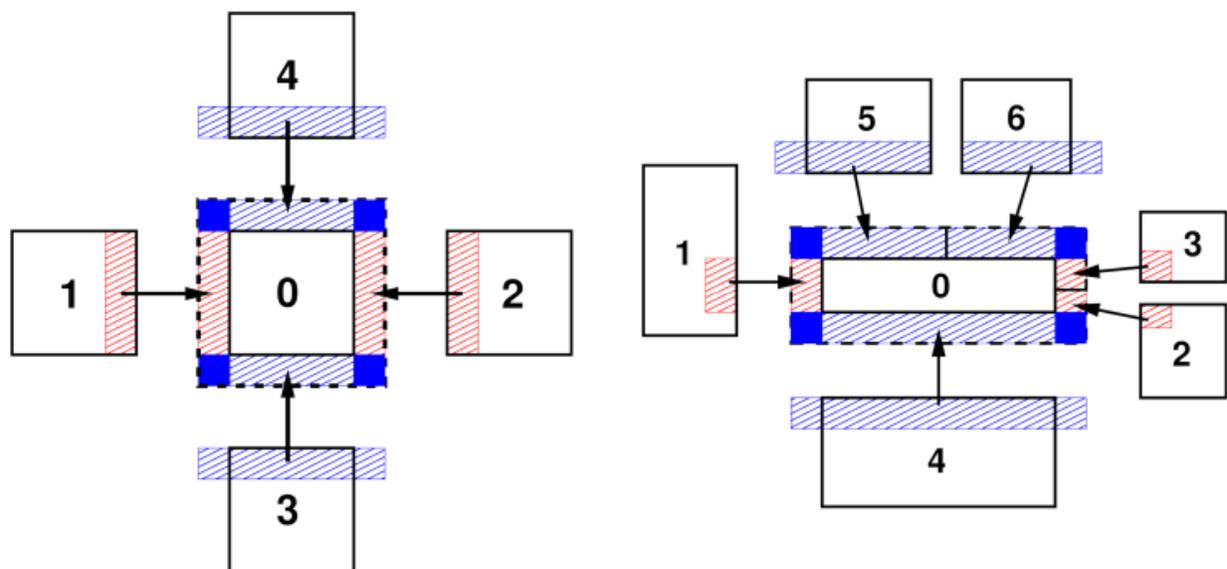


Fig. 3: ghost atom communication

This figure shows the ghost atom communication patterns between sub-domains for “brick” (left) and “tiled” communication styles for 2d simulations. The numbers indicate MPI process ranks. Here the sub-domains are drawn spatially separated for clarity. The dashed-line box is the extended sub-domain of processor 0 which includes its ghost atoms. The red- and blue-shaded boxes are the regions of communicated ghost atoms.

These include only atoms between the lower and upper  $y$ -boundary of rank 0’s sub-domain. In the  $y$  stage, ranks 4,5,6 send atoms in their blue-shaded regions to rank 0. This may include ghost atoms they received in the  $x$  stage, but only if they are needed by rank 0 to fill its extended ghost atom regions in the  $\pm y$  directions (blue rectangles). Thus in this case, ranks 5 and 6 do not include ghost atoms they received from each other (in the  $x$  stage) in the atoms they send to rank 0. The key point is that while the pattern of communication is more complex in the irregular partitioning case, it can still proceed in two stages (three in 3d) via atom exchanges with only neighboring processors.

When attributes of owned atoms are sent to neighboring processors to become attributes of their ghost atoms, LAMMPS calls this a “forward” communication. On timesteps when atoms migrate to new owning processors and neighbor lists are rebuilt, each processor creates a list of its owned atoms which are ghost atoms in each of its neighbor processors. These lists are used to pack per-atom coordinates (for example) into message buffers in subsequent steps until the next reneighboring.

A “reverse” communication is when computed ghost atom attributes are sent back to the processor who owns the atom. This is used (for example) to sum partial forces on ghost atoms to the complete force on owned atoms. The order of the two stages described in the *ghost atom communication* figure is inverted and the same lists of atoms are used to pack and unpack message buffers with per-atom forces. When a received buffer is unpacked, the ghost forces are summed to owned atom forces. As in forward communication, forces on atoms in the four blue corners of the diagrams are sent, received, and summed twice (once at each stage) before owning processors have the full force.

These two operations are used many places within LAMMPS aside from exchange of coordinates and forces, for example by manybody potentials to share intermediate per-atom values, or by rigid-body integrators to enable each atom in a body to access body properties. Here are additional details about how these communication operations are performed in LAMMPS:

- When exchanging data with different processors, forward and reverse communication is done using `MPI_Send()` and `MPI_IRecv()` calls. If a processor is “exchanging” atoms with itself, only the pack and unpack operations are performed, e.g. to create ghost atoms across periodic boundaries when running on a single processor.
- For forward communication of owned atom coordinates, periodic box lengths are added and subtracted when the receiving processor is across a periodic boundary from the sender. There is then no need to apply a minimum

image convention when calculating distances between atom pairs when building neighbor lists or computing forces.

- The cutoff distance for exchanging ghost atoms is typically equal to the neighbor cutoff. But it can also be chosen to be longer if needed, e.g. half the diameter of a rigid body composed of multiple atoms or over 3x the length of a stretched bond for dihedral interactions. It can also exceed the periodic box size. For the regular communication pattern (left), if the cutoff distance extends beyond a neighbor processor's sub-domain, then multiple exchanges are performed in the same direction. Each exchange is with the same neighbor processor, but buffers are packed/unpacked using a different list of atoms. For forward communication, in the first exchange a processor sends only owned atoms. In subsequent exchanges, it sends ghost atoms received in previous exchanges. For the irregular pattern (right) overlaps of a processor's extended ghost-atom sub-domain with all other processors in each dimension are detected.

### 15.4.3 Neighbor lists

To compute forces efficiently, each processor creates a Verlet-style neighbor list which enumerates all pairs of atoms  $i, j$  ( $i$  = owned,  $j$  = owned or ghost) with separation less than the applicable neighbor list cutoff distance. In LAMMPS the neighbor lists are stored in a multiple-page data structure; each page is a contiguous chunk of memory which stores vectors of neighbor atoms  $j$  for many  $i$  atoms. This allows pages to be incrementally allocated or deallocated in blocks as needed. Neighbor lists typically consume the most memory of any data structure in LAMMPS. The neighbor list is rebuilt (from scratch) once every few timesteps, then used repeatedly each step for force or other computations. The neighbor cutoff distance is  $R_n = R_f + \Delta_s$ , where  $R_f$  is the (largest) force cutoff defined by the interatomic potential for computing short-range pairwise or manybody forces and  $\Delta_s$  is a "skin" distance that allows the list to be used for multiple steps assuming that atoms do not move very far between consecutive time steps. Typically the code triggers reneighboring when any atom has moved half the skin distance since the last reneighboring; this and other options of the neighbor list rebuild can be adjusted with the `neigh_modify` command.

On steps when reneighboring is performed, atoms which have moved outside their owning processor's sub-domain are first migrated to new processors via communication. Periodic boundary conditions are also (only) enforced on these steps to ensure each atom is re-assigned to the correct processor. After migration, the atoms owned by each processor are stored in a contiguous vector. Periodically each processor spatially sorts owned atoms within its vector to reorder it for improved cache efficiency in force computations and neighbor list building. For that atoms are spatially binned and then reordered so that atoms in the same bin are adjacent in the vector. Atom sorting can be disabled or its settings modified with the `atom_modify` command.

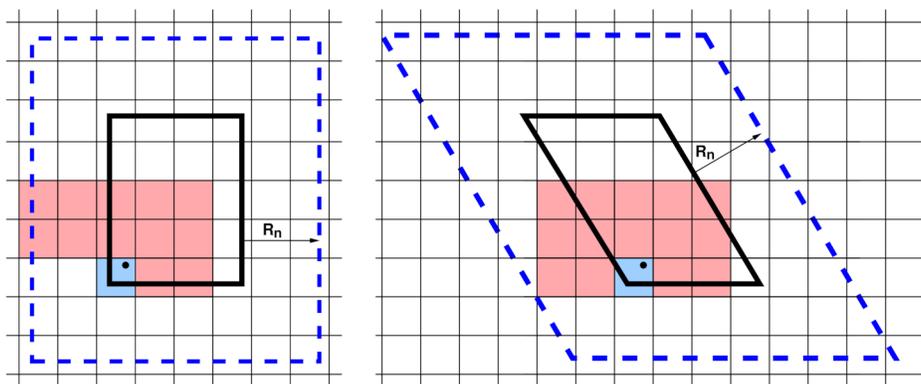


Fig. 4: neighbor list stencils

A 2d simulation sub-domain (thick black line) and the corresponding ghost atom cutoff region (dashed blue line) for both orthogonal (left) and triclinic (right) domains. A regular grid of neighbor bins (thin lines) overlays the entire simulation domain and need not align with sub-domain boundaries; only the portion overlapping the augmented sub-domain is shown. In the triclinic case it overlaps the bounding box of the tilted rectangle. The blue- and red-shaded bins represent a stencil of bins searched to find neighbors of a particular atom (black dot).

To build a local neighbor list in linear time, the simulation domain is overlaid (conceptually) with a regular 3d (or 2d) grid of neighbor bins, as shown in the *neighbor list stencils* figure for 2d models and a single MPI processor's sub-domain. Each processor stores a set of neighbor bins which overlap its sub-domain extended by the neighbor cutoff distance  $R_n$ . As illustrated, the bins need not align with processor boundaries; an integer number in each dimension is fit to the size of the entire simulation box.

Most often LAMMPS builds what it calls a “half” neighbor list where each  $i,j$  neighbor pair is stored only once, with either atom  $i$  or  $j$  as the central atom. The build can be done efficiently by using a pre-computed “stencil” of bins around a central origin bin which contains the atom whose neighbors are being searched for. A stencil is simply a list of integer offsets in  $x,y,z$  of nearby bins surrounding the origin bin which are close enough to contain any neighbor atom  $j$  within a distance  $R_n$  from any atom  $i$  in the origin bin. Note that for a half neighbor list, the stencil can be asymmetric since each atom only need store half its nearby neighbors.

These stencils are illustrated in the figure for a half list and a bin size of  $\frac{1}{2}R_n$ . There are 13 red+blue stencil bins in 2d (for the orthogonal case, 15 for triclinic). In 3d there would be 63, 13 in the plane of bins that contain the origin bin and 25 in each of the two planes above it in the  $z$  direction (75 for triclinic). The reason the triclinic stencil has extra bins is because the bins tile the bounding box of the entire triclinic domain and thus are not periodic with respect to the simulation box itself. The stencil and logic for determining which  $i,j$  pairs to include in the neighbor list are altered slightly to account for this.

To build a neighbor list, a processor first loops over its “owned” plus “ghost” atoms and assigns each to a neighbor bin. This uses an integer vector to create a linked list of atom indices within each bin. It then performs a triply-nested loop over its owned atoms  $i$ , the stencil of bins surrounding atom  $i$ 's bin, and the  $j$  atoms in each stencil bin (including ghost atoms). If the distance  $r_{ij} < R_n$ , then atom  $j$  is added to the vector of atom  $i$ 's neighbors.

Here are additional details about neighbor list build options LAMMPS supports:

- The choice of bin size is an option; a size half of  $R_n$  has been found to be optimal for many typical cases. Smaller bins incur additional overhead to loop over; larger bins require more distance calculations. Note that for smaller bin sizes, the 2d stencil in the figure would be more semi-circular in shape (hemispherical in 3d), with bins near the corners of the square eliminated due to their distance from the origin bin.
- Depending on the interatomic potential(s) and other commands used in an input script, multiple neighbor lists and stencils with different attributes may be needed. This includes lists with different cutoff distances, e.g. for force computation versus occasional diagnostic computations such as a radial distribution function, or for the r-RESPA time integrator which can partition pairwise forces by distance into subsets computed at different time intervals. It includes “full” lists (as opposed to half lists) where each  $i,j$  pair appears twice, stored once with  $i$  and  $j$ , and which use a larger symmetric stencil. It also includes lists with partial enumeration of ghost atom neighbors. The full and ghost-atom lists are used by various manybody interatomic potentials. Lists may also use different criteria for inclusion of a pair interaction. Typically this simply depends only on the distance between two atoms and the cutoff distance. But for finite-size coarse-grained particles with individual diameters (e.g. polydisperse granular particles), it can also depend on the diameters of the two particles.
- When using *pair style hybrid* multiple sub-lists of the master neighbor list for the full system need to be generated, one for each sub-style, which contains only the  $i,j$  pairs needed to compute interactions between subsets of atoms for the corresponding potential. This means not all  $i$  or  $j$  atoms owned by a processor are included in a particular sub-list.
- Some models use different cutoff lengths for pairwise interactions between different kinds of particles which are stored in a single neighbor list. One example is a solvated colloidal system with large colloidal particles where colloid/colloid, colloid/solvent, and solvent/solvent interaction cutoffs can be dramatically different. Another is a model of polydisperse finite-size granular particles; pairs of particles interact only when they are in contact with each other. Mixtures with particle size ratios as high as 10-100x may be used to model realistic systems. Efficient neighbor list building algorithms for these kinds of systems are available in LAMMPS. They include a method which uses different stencils for different cutoff lengths and trims the stencil to only include bins that straddle the cutoff sphere surface. More recently a method which uses both multiple stencils and multiple bin sizes was developed; it builds neighbor lists efficiently for systems with particles of any size ratio, though other considerations (timestep size, force computations) may limit the ability to model systems with huge polydispersity.

- For small and sparse systems and as a fallback method, LAMMPS also supports neighbor list construction without binning by using a full  $O(N^2)$  loop over all  $i,j$  atom pairs in a sub-domain when using the *neighbor nsq* command.
- Dependent on the “pair” setting of the *newton* command, the “half” neighbor lists may contain **all** pairs of atoms where atom  $j$  is a ghost atom (i.e. when the *newton* pair setting is *off*) For the *newton* pair *on* setting the atom  $j$  is only added to the list if its  $z$  coordinate is larger, or if equal the  $y$  coordinate is larger, and that is equal, too, the  $x$  coordinate is larger. For homogeneously dense systems that will result in picking neighbors from a same size sector in always the same direction relative to the “owned” atom and thus it should lead to similar length neighbor lists and thus reduce the chance of a load imbalance.

### 15.4.4 Long-range interactions

For charged systems, LAMMPS can compute long-range Coulombic interactions via the FFT-based particle-particle/particle-mesh (PPPM) method implemented in *k-space style pppm and its variants*. For that Coulombic interactions are partitioned into short- and long-range components. The short-ranged portion is computed in real space as a loop over pairs of charges within a cutoff distance, using neighbor lists. The long-range portion is computed in reciprocal space using a *k-space* style. For the PPPM implementation the simulation cell is overlaid with a regular FFT grid in 3d. It proceeds in several stages:

- a) each atom’s point charge is interpolated to nearby FFT grid points,
- b) a forward 3d FFT is performed,
- c) a convolution operation is performed in reciprocal space,
- d) one or more inverse 3d FFTs are performed, and
- e) electric field values from grid points near each atom are interpolated to compute its forces.

For any of the spatial-decomposition partitioning schemes each processor owns the brick-shaped portion of FFT grid points contained within its sub-domain. The two interpolation operations use a stencil of grid points surrounding each atom. To accommodate the stencil size, each processor also stores a few layers of ghost grid points surrounding its brick. Forward and reverse communication of grid point values is performed similar to the corresponding *atom data communication*. In this case, electric field values on owned grid points are sent to neighboring processors to become ghost point values. Likewise charge values on ghost points are sent and summed to values on owned points.

For triclinic simulation boxes, the FFT grid planes are parallel to the box faces, but the mapping of charge and electric field values to/from grid points is done in reduced coordinates where the tilted box is conceptually a unit cube, so that the stencil and FFT operations are unchanged. However the FFT grid size required for a given accuracy is larger for triclinic domains than it is for orthogonal boxes.

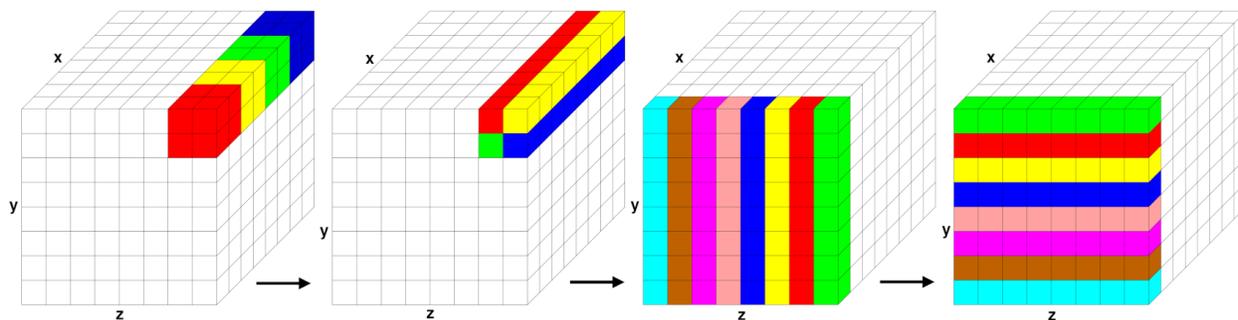


Fig. 5: parallel FFT in PPPM

Stages of a parallel FFT for a simulation domain overlaid with an  $8 \times 8 \times 8$  3d FFT grid, partitioned across 64 processors. Within each of the 4 diagrams, grid cells of the same color are owned by a single processor; for simplicity only cells owned by 4 or 8 of the 64 processors are colored. The two images on the left illustrate brick-to-pencil communication. The two images on the right illustrate pencil-to-pencil communication, which in this case transposes the  $y$  and  $z$  dimensions of the grid.

Parallel 3d FFTs require substantial communication relative to their computational cost. A 3d FFT is implemented by a series of 1d FFTs along the  $x$ -,  $y$ -, and  $z$ -direction of the FFT grid. Thus the FFT grid cannot be decomposed like atoms into 3 dimensions for parallel processing of the FFTs but only in 1 (as planes) or 2 (as pencils) dimensions and in between the steps the grid needs to be transposed to have the FFT grid portion “owned” by each MPI process complete in the direction of the 1d FFTs it has to perform. LAMMPS uses the pencil-decomposition algorithm as shown in the *parallel FFT in PPPM* figure.

Initially (far left), each processor owns a brick of same-color grid cells (actually grid points) contained within its sub-domain. A brick-to-pencil communication operation converts this layout to 1d pencils in the  $x$ -dimension (center left). Again, cells of the same color are owned by the same processor. Each processor can then compute a 1d FFT on each pencil of data it wholly owns using a call to the configured FFT library. A pencil-to-pencil communication then converts this layout to pencils in the  $y$  dimension (center right) which effectively transposes the  $x$  and  $y$  dimensions of the grid, followed by 1d FFTs in  $y$ . A final transpose of pencils from  $y$  to  $z$  (far right) followed by 1d FFTs in  $z$  completes the forward FFT. The data is left in a  $z$ -pencil layout for the convolution operation. One or more inverse FFTs then perform the sequence of 1d FFTs and communication steps in reverse order; the final layout of resulting grid values is the same as the initial brick layout.

Each communication operation within the FFT (brick-to-pencil or pencil-to-pencil or pencil-to-brick) converts one tiling of the 3d grid to another, where a tiling in this context means an assignment of a small brick-shaped subset of grid points to each processor, the union of which comprise the entire grid. The parallel *fftMPI* library written for LAMMPS allows arbitrary definitions of the tiling so that an irregular partitioning of the simulation domain can use it directly. Transforming data from one tiling to another is implemented in *fftMPI* using point-to-point communication, where each processor sends data to a few other processors, since each tile in the initial tiling overlaps with a handful of tiles in the final tiling.

The transformations could also be done using collective communication across all  $P$  processors with a single call to `MPI_Alltoall()`, but this is typically much slower. However, for the specialized brick and pencil tiling illustrated in *parallel FFT in PPPM* figure, collective communication across the entire MPI communicator is not required. In the example an  $8^3$  grid with 512 grid cells is partitioned across 64 processors; each processor owns a  $2 \times 2 \times 2$  3d brick of grid cells. The initial brick-to-pencil communication (upper left to upper right) only requires collective communication within subgroups of 4 processors, as illustrated by the 4 colors. More generally, a brick-to-pencil communication can be performed by partitioning  $P$  processors into  $P^{\frac{2}{3}}$  subgroups of  $P^{\frac{1}{3}}$  processors each. Each subgroup performs collective communication only within its subgroup. Similarly, pencil-to-pencil communication can be performed by partitioning  $P$  processors into  $P^{\frac{1}{2}}$  subgroups of  $P^{\frac{1}{2}}$  processors each. This is illustrated in the figure for the  $y \Rightarrow z$  communication (center). An eight-processor subgroup owns the front  $yz$  plane of data and performs collective communication within the subgroup to transpose from a  $y$ -pencil to  $z$ -pencil layout.

LAMMPS invokes point-to-point communication by default, but also provides the option of partitioned collective communication when using a *kpspace\_modify collective yes* command to switch to that mode. In the latter case, the code detects the size of the disjoint subgroups and partitions the single  $P$ -size communicator into multiple smaller communicators, each of which invokes collective communication. Testing on a large IBM Blue Gene/Q machine at Argonne National Labs showed a significant improvement in FFT performance for large processor counts; partitioned collective communication was faster than point-to-point communication or global collective communication involving all  $P$  processors.

Here are some additional details about FFTs for long-range and related grid/particle operations that LAMMPS supports:

- The *fftMPI* library allows each grid dimension to be a multiple of small prime factors (2,3,5), and allows any number of processors to perform the FFT. The resulting brick and pencil decompositions are thus not always as well-aligned but the size of subgroups of processors for the two modes of communication (brick/pencil and pencil/pencil) still scale as  $O(P^{\frac{1}{3}})$  and  $O(P^{\frac{1}{2}})$ .
- For efficiency in performing 1d FFTs, the grid transpose operations illustrated in Figure [ref{fig:fft}](#) also involve reordering the 3d data so that a different dimension is contiguous in memory. This reordering can be done during the packing or unpacking of buffers for MPI communication.
- For large systems and particularly a large number of MPI processes, the dominant cost for parallel FFTs is often the communication, not the computation of 1d FFTs, even though the latter scales as  $N \log(N)$  in the number of

grid points  $N$  per grid direction. This is due to the fact that only a 2d decomposition into pencils is possible while atom data (and their corresponding short-range force and energy computations) can be decomposed efficiently in 3d.

This can be addressed by reducing the number of MPI processes involved in the MPI communication by using *hybrid MPI + OpenMP parallelization*. This will use OpenMP parallelization inside the MPI domains and while that may have a lower parallel efficiency, it reduces the communication overhead.

As an alternative it is also possible to start a *multi-partition* calculation and then use the *verlet/split integrator* to perform the PPPM computation on a dedicated, separate partition of MPI processes. This uses an integer “1: $p$ ” mapping of  $p$  sub-domains of the atom decomposition to one sub-domain of the FFT grid decomposition and where pairwise non-bonded and bonded forces and energies are computed on the larger partition and the PPPM kspace computation concurrently on the smaller partition.

- LAMMPS also implements PPPM-based solvers for other long-range interactions, dipole and dispersion (Lennard-Jones), which can be used in conjunction with long-range Coulombics for point charges.
- LAMMPS implements a `GridComm` class which overlays the simulation domain with a regular grid, partitions it across processors in a manner consistent with processor sub-domains, and provides methods for forward and reverse communication of owned and ghost grid point values. It is used for PPPM as an FFT grid (as outlined above) and also for the MSM algorithm which uses a cascade of grid sizes from fine to coarse to compute long-range Coulombic forces. The `GridComm` class is also useful for models where continuum fields interact with particles. For example, the two-temperature model (TTM) defines heat transfer between atoms (particles) and electrons (continuum gas) where spatial variations in the electron temperature are computed by finite differences of a discretized heat equation on a regular grid. The `fix ttm/grid` command uses the `GridComm` class internally to perform its grid operations on a distributed grid instead of the original `fix ttm` which uses a replicated grid.

### 15.4.5 OpenMP Parallelism

The styles in the INTEL, KOKKOS, and OPENMP package offer to use OpenMP thread parallelism to predominantly distribute loops over local data and thus follow an orthogonal parallelization strategy to the decomposition into spatial domains used by the *MPI partitioning*. For clarity, this section discusses only the implementation in the OPENMP package as it is the simplest. The INTEL and KOKKOS package offer additional options and are more complex since they support more features and different hardware like co-processors or GPUs.

One of the key decisions when implementing the OPENMP package was to keep the changes to the source code small, so that it would be easier to maintain the code and keep it in sync with the non-threaded standard implementation. This is achieved by a) making the OPENMP version a derived class from the regular version (e.g. `PairLJCutOMP` from `PairLJCut`) and overriding only methods that are multi-threaded or need to be modified to support multi-threading (similar to what was done in the OPT package), b) keeping the structure in the modified code very similar so that side-by-side comparisons are still useful, and c) offloading additional functionality and multi-thread support functions into three separate classes `ThrOMP`, `ThrData`, and `FixOMP`. `ThrOMP` provides additional, multi-thread aware functionality not available in the corresponding base class (e.g. `Pair` for `PairLJCutOMP`) like multi-thread aware variants of the “tally” functions. Those functions are made available through multiple inheritance so those new functions have to have unique names to avoid ambiguities; typically `_thr` is appended to the name of the function. `ThrData` is a classes that manages per-thread data structures. It is used instead of extending the corresponding storage to per-thread arrays to avoid slowdowns due to “false sharing” when multiple threads update adjacent elements in an array and thus force the CPU cache lines to be reset and re-fetched. `FixOMP` finally manages the “multi-thread state” like settings and access to per-thread storage, it is activated by the *package omp* command.



## Avoiding data races

A key problem when implementing thread parallelism in an MD code is to avoid data races when updating accumulated properties like forces, energies, and stresses. When interactions are computed, they always involve multiple atoms and thus there are race conditions when multiple threads want to update per-atom data of the same atoms. Five possible strategies have been considered to avoid this:

- 1) restructure the code so that there is no overlapping access possible when computing in parallel, e.g. by breaking lists into multiple parts and synchronizing threads in between.
- 2) have each thread be “responsible” for a specific group of atoms and compute these interactions multiple times, once on each thread that is responsible for a given atom and then have each thread only update the properties of this atom.
- 3) use mutexes around functions and regions of code where the data race could happen
- 4) use atomic operations when updating per-atom properties
- 5) use replicated per-thread data structures to accumulate data without conflicts and then use a reduction to combine those results into the data structures used by the regular style.

Option 5 was chosen for the OPENMP package because it would retain the performance for the case of 1 thread and the code would be more maintainable. Option 1 would require extensive code changes, particularly to the neighbor list code; options 2 would have incurred a 2x or more performance penalty for the serial case; option 3 causes significant overhead and would enforce serialization of operations in inner loops and thus defeat the purpose of multi-threading; option 4 slows down the serial case although not quite as bad as option 2. The downside of option 5 is that the overhead of the reduction operations grows with the number of threads used, so there would be a crossover point where options 2 or 4 would result in faster executing. That is why option 2 for example is used in the GPU package because a GPU is a processor with a massive number of threads. However, since the MPI parallelization is generally more effective for typical MD systems, the expectation is that thread parallelism is only used for a smaller number of threads (2-8). At the time of its implementation, that number was equivalent to the number of CPU cores per CPU socket on high-end supercomputers.

Thus arrays like the force array are dimensioned to the number of atoms times the number of threads when enabling OpenMP support and inside the compute functions a pointer to a different chunk is obtained by each thread. Similarly, accumulators like potential energy or virial are kept in per-thread instances of the ThrData class and then only reduced and stored in their global counterparts at the end of the force computation.

## Loop scheduling

Multi-thread parallelization is applied by distributing (outer) loops statically across threads. Typically this would be the loop over local atoms  $i$  when processing  $i,j$  pairs of atoms from a neighbor list. The design of the neighbor list code results in atoms having a similar number of neighbors for homogeneous systems and thus load imbalances across threads are not common and typically happen for systems where also the MPI parallelization would be unbalanced, which would typically have a more pronounced impact on the performance. This same loop scheduling scheme can also be applied to the reduction operations on per-atom data to try and reduce the overhead of the reduction operation.

## Neighbor list parallelization

In addition to the parallelization of force computations, also the generation of the neighbor lists is parallelized. As explained previously, neighbor lists are built by looping over “owned” atoms and storing the neighbors in “pages”. In the OPENMP variants of the neighbor list code, each thread operates on a different chunk of “owned” atoms and allocates and fills its own set of pages with neighbor list data. This is achieved by each thread keeping its own instance of the *MyPage* page allocator class.

## 15.5 Communication patterns

This page describes various inter-processor communication operations provided by LAMMPS, mostly in the core *Comm* class. These are operations for common tasks implemented using MPI library calls. They are used by other classes to perform communication of different kinds. These operations are useful to know about when writing new code for LAMMPS that needs to communicate data between processors.

### 15.5.1 Owned and ghost atoms

As described on the *parallel partitioning algorithms* page, LAMMPS spatially decomposes the simulation domain, either in a *brick* or *tiled* manner. Each processor (MPI task) owns atoms within its sub-domain and additionally stores ghost atoms within a cutoff distance of its sub-domain.

#### Forward and reverse communication

As described on the *parallel communication algorithms* page, the most common communication operations are first, *forward communication* which sends owned atom information from each processor to nearby processors to store with their ghost atoms. The need to do this communication arises when data from the owned atoms is updated (e.g. their positions) and this updated information needs to be **copied** to the corresponding ghost atoms.

And second, *reverse communication* which sends ghost atom information from each processor to the owning processor to **accumulate** (sum) the values with the corresponding owned atoms. The need for this arises when data is computed and also stored with ghost atoms (e.g. forces when using a “half” neighbor list) and thus those terms need to be added to their corresponding atoms on the process where they are “owned” atoms. Please note, that with the *newton off* setting this does not happen and the neighbor lists are constructed so that these interactions are computed on both MPI processes containing one of the atoms and only the data pertaining to the local atom is stored.

The time-integration classes in LAMMPS invoke these operations each timestep via the *forward\_comm()* and *reverse\_comm()* methods in the *Comm* class. Which per-atom data is communicated depends on the currently used *atom style* and whether *comm\_modify vel* setting is “no” (default) or “yes”.

Similarly, *Pair* style classes can invoke the *forward\_comm(this)* and *reverse\_comm(this)* methods in the *Comm* class to perform the same operations on per-atom data that is generated and stored within the pair style class. Note that this function requires passing the `this` pointer as the first argument to enable the *Comm* class to call the “pack” and “unpack” functions discussed below. An example of the use of these functions are many-body pair styles like the embedded-atom method (EAM) which compute intermediate values in the first part of the *compute()* function that need to be stored by both owned and ghost atoms for the second part of the force computation. The *Comm* class methods perform the MPI communication for buffers of per-atom data. They “call back” to the *Pair* class so it can *pack* or *unpack* the buffer with data the *Pair* class owns. There are 4 such methods that the *Pair* class must define, assuming it uses both forward and reverse communication:

- `pack_forward_comm()`
- `unpack_forward_comm()`
- `pack_reverse_comm()`

- `unpack_reverse_comm()`

The arguments to these methods include the buffer and a list of atoms to pack or unpack. The *Pair* class also must set the `comm_forward` and `comm_reverse` variables which store the number of values stored in the communication buffers for each operation. This means, if desired, it can choose to store multiple per-atom values in the buffer, and they will be communicated together to minimize communication overhead. The communication buffers are defined vectors containing double values. To correctly store integers that may be 64-bit (bigint, tagint, imageint) in the buffer, you need to use the `ubuf union` construct.

The *Fix*, *Compute*, and *Dump* classes can also invoke the same kind of forward and reverse communication operations using the same *Comm* class methods. Likewise the same pack/unpack methods and `comm_forward/comm_reverse` variables must be defined by the calling *Fix*, *Compute*, or *Dump* class.

For *Fix* classes there is an optional second argument to the `forward_comm()` and `reverse_comm()` call which can be used when the fix performs multiple modes of communication, with different numbers of values per atom. The fix should set the `comm_forward` and `comm_reverse` variables to the maximum value, but can invoke the communication for a particular mode with a smaller value. For this to work, the `pack_forward_comm()`, etc methods typically use a class member variable to choose which values to pack/unpack into/from the buffer.

Finally, for reverse communications in *Fix* classes there is also the `reverse_comm_variable()` method that allows the communication to have a different amount of data per-atom. It invokes these corresponding callback methods:

- `pack_reverse_comm_size()`
- `unpack_reverse_comm_size()`

which have extra arguments to specify the amount of data stored in the buffer for each atom.

## 15.5.2 Higher level communication

There are also several higher-level communication operations provided in LAMMPS which work for either *brick* or *tiled* decompositions. They may be useful for a new class to invoke if it requires more sophisticated communication than the *forward* and *reverse* methods provide. The 3 communication operations described here are

- ring
- irregular
- rendezvous

You can invoke these *grep* command in the LAMMPS src directory, to see a list of classes that invoke the 3 operations.

- `grep "\->ring" *.cpp */*.cpp`
- `grep "irregular\->" *.cpp`
- `grep "\->rendezvous" *.cpp */*.cpp`

### Ring operation

The *ring* operation is invoked via the `ring()` method in the *Comm* class.

Each processor first creates a buffer with a list of values, typically associated with a subset of the atoms it owns. Now think of the  $P$  processors as connected to each other in a *ring*. Each processor  $M$  sends data to the next  $M+1$  processor. It receives data from the preceding  $M-1$  processor. The ring is periodic so that the last processor sends to the first processor, and the first processor receives from the last processor.

Invoking the `ring()` method passes each processor's buffer in  $P$  steps around the ring. At each step a *callback* method, provided as an argument to `ring()`, in the caller is invoked. This allows each processor to examine the data buffer provided by every other processor. It may extract values needed by its atoms from the buffers, or it may alter placeholder

values in the buffer. In the latter case, when the *ring* operation is complete, each processor can examine its original buffer to extract modified values.

Note that the *ring* operation is similar to an `MPI_Alltoall()` operation where every processor effectively sends and receives data to every other processor. The difference is that the *ring* operation does it one step at a time, so the total volume of data does not need to be stored by every processor. However, the *ring* operation is also less efficient than `MPI_Alltoall()` because of the  $P$  stages required. So it is typically only suitable for small data buffers and occasional operations that are not time-critical.

## Irregular operation

The *irregular* operation is provided by the *Irregular* class. What LAMMPS terms irregular communication is when each processor knows what data it needs to send to what processor, but does not know what processors are sending it data. An example is when load-balancing is performed and each processor needs to send some of its atoms to new processors.

The *Irregular* class provides 5 high-level methods useful in this context:

- `create_data()`
- `exchange_data()`
- `create_atom()`
- `exchange_atom()`
- `migrate_atoms()`

For the `create_data()` method, each processor specifies a list of  $N$  datums to send, each to a specified processor. Internally, the method creates efficient data structures for performing the communication. The `exchange_data()` method triggers the communication to be performed. Each processor provides the vector of  $N$  datums to send, and the size of each datum. All datums must be the same size.

The `create_atom()` and `exchange_atom()` methods are similar except that the size of each datum can be different. Typically this is used to communicate atoms, each with a variable amount of per-atom data, to other processors.

The `migrate_atoms()` method is a convenience wrapper on the `create_atom()` and `exchange_atom()` methods to simplify communication of all the per-atom data associated with an atom so that the atom can effectively migrate to a new owning processor. It is similar to the `exchange()` method in the *Comm* class invoked when atoms move to neighboring processors (in the regular or tiled decomposition) during timestepping, except that it allows atoms to have moved arbitrarily long distances and still be properly communicated to a new owning processor.

## Rendezvous operation

Finally, the *rendezvous* operation is invoked via the `rendezvous()` method in the *Comm* class. Depending on how much communication is needed and how many processors a LAMMPS simulation is running on, it can be a much more efficient choice than the *ring()* method. It uses the *irregular* operation internally once or twice to do its communication. The rendezvous algorithm is described in detail in (*Plimpton*), including some LAMMPS use cases.

For the `rendezvous()` method, each processor specifies a list of  $N$  datums to send and which processor to send each of them to. Internally, this communication is performed as an irregular operation. The received datums are returned to the caller via invocation of *callback* function, provided as an argument to `rendezvous()`. The caller can then process the received datums and (optionally) assemble a new list of datums to communicate to a new list of specific processors. When the callback function exits, the `rendezvous()` method performs a second irregular communication on the new list of datums.

Examples in LAMMPS of use of the *rendezvous* operation are the *fix rigid/small* and *fix shake* commands (for one-time identification of the rigid body atom clusters) and the identification of special\_bond 1-2, 1-3 and 1-4 neighbors within molecules. See the *special\_bonds* command for context.

(Plimpton) Plimpton and Knight, JPDC, 147, 184-195 (2021).

## 15.6 How a timestep works

The first and most fundamental operation within LAMMPS to understand is how a timestep is structured. Timestepping is performed by calling methods of the Integrate class instance within the Update class. Since Integrate is a base class, it will point to an instance of a derived class corresponding to what is selected by the *run\_style* input script command.

In this section, the timestep implemented by the Verlet class is described. A similar timestep protocol is implemented by the Respa class, for the r-RESPA hierarchical timestepping method.

The Min base class performs energy minimization, so does not perform a literal timestep. But it has logic similar to what is described here, to compute forces and invoke fixes at each iteration of a minimization. Differences between time integration and minimization are highlighted at the end of this section.

The Verlet class is encoded in the `src/verlet.cpp` and `verlet.h` files. It implements the velocity-Verlet timestepping algorithm. The workhorse method is `Verlet::run()`, but first we highlight several other methods in the class.

- The `init()` method is called at the beginning of each dynamics run. It simply sets some internal flags, based on user settings in other parts of the code.
- The `setup()` or `setup_minimal()` methods are also called before each run. The velocity-Verlet method requires current forces be calculated before the first timestep, so these routines compute forces due to all atomic interactions, using the same logic that appears in the timestepping described next. A few fixes are also invoked, using the mechanism described in the next section. Various counters are also initialized before the run begins. The `setup_minimal()` method is a variant that has a flag for performing less setup. This is used when runs are continued and information from the previous run is still valid. For example, if repeated short LAMMPS runs are being invoked, interleaved by other commands, via the *pre no* and *every* options of the run command, the `setup_minimal()` method is used.
- The `force_clear()` method initializes force and other arrays to zero before each timestep, so that forces (torques, etc) can be accumulated.

Now for the `Verlet::run()` method. Its basic structure in hi-level pseudo code is shown below. In the actual code in `src/verlet.cpp` some of these operations are conditionally invoked.

```

loop over N timesteps:
  if timeout condition: break
  ev_set()

  fix->initial_integrate()
  fix->post_integrate()

  nflag = neighbor->decide()
  if nflag:
    fix->pre_exchange()
    domain->pbcb()
    domain->reset_box()
    comm->setup()
    neighbor->setup_bins()
    comm->exchange()
    comm->borders()
    fix->pre_neighbor()

```

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```

neighbor->build()
fix->post_neighbor()
else:
  comm->forward_comm()

force_clear()
fix->pre_force()

pair->compute()
bond->compute()
angle->compute()
dihedral->compute()
improper->compute()
kspace->compute()

fix->pre_reverse()
comm->reverse_comm()

fix->post_force()
fix->final_integrate()
fix->end_of_step()

if any output on this step:
  output->write()

# after loop
fix->post_run()

```

The `ev_set()` method (in the parent `Integrate` class), sets two flags (*eflag* and *vflag*) for energy and virial computation. Each flag encodes whether global and/or per-atom energy and virial should be calculated on this timestep, because some fix or variable or output will need it. These flags are passed to the various methods that compute particle interactions, so that they either compute and tally the corresponding data or can skip the extra calculations if the energy and virial are not needed. See the comments for the `Integrate::ev_set()` method which document the flag values.

At various points of the timestep, fixes are invoked, e.g. `fix->initial_integrate()`. In the code, this is actually done via the `Modify` class which stores all the `Fix` objects and lists of which should be invoked at what point in the timestep. Fixes are the LAMMPS mechanism for tailoring the operations of a timestep for a particular simulation. As described elsewhere, each fix has one or more methods, each of which is invoked at a specific stage of the timestep, as show in the timestep pseudo-code. All the active fixes defined in an input script, that are flagged to have an `initial_integrate()` method are invoked at the beginning of each timestep. Examples are *fix nve* or *fix nvt* or *fix npt* which perform the start-of-timestep velocity-Verlet integration operations to update velocities by a half-step, and coordinates by a full step. The `post_integrate()` method is next for operations that need to happen immediately after those updates. Only a few fixes use this, e.g. to reflect particles off box boundaries in the *FixWallReflect* class.

The `decide()` method in the `Neighbor` class determines whether neighbor lists need to be rebuilt on the current timestep (conditions can be changed using the *neigh\_modify every/delay/check* command. If not, coordinates of ghost atoms are acquired by each processor via the `forward_comm()` method of the `Comm` class. If neighbor lists need to be built, several operations within the inner if clause of the pseudo-code are first invoked. The `pre_exchange()` method of any defined fixes is invoked first. Typically this inserts or deletes particles from the system.

Periodic boundary conditions are then applied by the `Domain` class via its `pbcc()` method to remap particles that have moved outside the simulation box back into the box. Note that this is not done every timestep, but only when neighbor lists are rebuilt. This is so that each processor's sub-domain will have consistent (nearby) atom coordinates for its owned and ghost atoms. It is also why dumped atom coordinates may be slightly outside the simulation box if not

dumped on a step where the neighbor lists are rebuilt.

The box boundaries are then reset (if needed) via the `reset_box()` method of the `Domain` class, e.g. if box boundaries are shrink-wrapped to current particle coordinates. A change in the box size or shape requires internal information for communicating ghost atoms (`Comm` class) and neighbor list bins (`Neighbor` class) be updated. The `setup()` method of the `Comm` class and `setup_bins()` method of the `Neighbor` class perform the update.

The code is now ready to migrate atoms that have left a processor's geometric sub-domain to new processors. The `exchange()` method of the `Comm` class performs this operation. The `borders()` method of the `Comm` class then identifies ghost atoms surrounding each processor's sub-domain and communicates ghost atom information to neighboring processors. It does this by looping over all the atoms owned by a processor to make lists of those to send to each neighbor processor. On subsequent timesteps, the lists are used by the `Comm::forward_comm()` method.

Fixes with a `pre_neighbor()` method are then called. These typically re-build some data structure stored by the fix that depends on the current atoms owned by each processor.

Now that each processor has a current list of its owned and ghost atoms, LAMMPS is ready to rebuild neighbor lists via the `build()` method of the `Neighbor` class. This is typically done by binning all owned and ghost atoms, and scanning a stencil of bins around each owned atom's bin to make a Verlet list of neighboring atoms within the force cutoff plus neighbor skin distance.

In the next portion of the timestep, all interaction forces between particles are computed, after zeroing the per-atom force vector via the `force_clear()` method. If the `newton` flag is set to *on* by the `newton` command, forces are added to both owned and ghost atoms, otherwise only to owned (aka local) atoms.

Pairwise forces are calculated first, which enables the global virial (if requested) to be calculated cheaply (at  $O(N)$  cost instead of  $O(N^2)$  at the end of the `Pair::compute()` method), by a dot product of atom coordinates and forces. By including owned and ghost atoms in the dot product, the effect of periodic boundary conditions is correctly accounted for. Molecular topology interactions (bonds, angles, dihedrals, impropers) are calculated next (if supported by the current atom style). The final contribution is from long-range Coulombic interactions, invoked by the `KSpace` class.

The `pre_reverse()` method in fixes is used for operations that have to be done *before* the upcoming reverse communication (e.g. to perform additional data transfers or reductions for data computed during the force computation and stored with ghost atoms).

If the `newton` flag is on, forces on ghost atoms are communicated and summed back to their corresponding owned atoms. The `reverse_comm()` method of the `Comm` class performs this operation, which is essentially the inverse operation of sending copies of owned atom coordinates to other processor's ghost atoms.

At this point in the timestep, the total force on each (local) atom is known. Additional force constraints (external forces, SHAKE, etc) are applied by Fixes that have a `post_force()` method. The second half of the velocity-Verlet integration, `final_integrate()` is then performed (another half-step update of the velocities) via fixes like `nve`, `nvt`, `npt`.

At the end of the timestep, fixes that contain an `end_of_step()` method are invoked. These typically perform a diagnostic calculation, e.g. the `ave/time` and `ave/spatial` fixes. The final operation of the timestep is to perform any requested output, via the `write()` method of the `Output` class. There are 3 kinds of LAMMPS output: thermodynamic output to the screen and log file, snapshots of atom data to a dump file, and restart files. See the `thermo_style`, `dump`, and `restart` commands for more details.

The flow of control during energy minimization iterations is similar to that of a molecular dynamics timestep. Forces are computed, neighbor lists are built as needed, atoms migrate to new processors, and atom coordinates and forces are communicated to neighboring processors. The only difference is what `Fix` class operations are invoked when. Only a subset of LAMMPS fixes are useful during energy minimization, as explained in their individual doc pages. The relevant `Fix` class methods are `min_pre_exchange()`, `min_pre_force()`, and `min_post_force()`. Each fix is invoked at the appropriate place within the minimization iteration. For example, the `min_post_force()` method is analogous to the `post_force()` method for dynamics; it is used to alter or constrain forces on each atom, which affects the minimization procedure.

After all iterations are completed there is a `cleanup` step which calls the `post_run()` method of fixes to perform operations only required at the end of a calculations (like freeing temporary storage or creating final outputs).

## 15.7 Writing new styles

The *Modifying & extending LAMMPS* section of the manual gives an overview of how LAMMPS can be extended by writing new classes that derive from existing parent classes in LAMMPS. Here, some specific coding details are provided for writing code for LAMMPS.

### 15.7.1 Writing a new fix style

Writing fixes is a flexible way of extending LAMMPS. Users can implement many things using fixes:

- changing particles attributes (positions, velocities, forces, etc.). Examples: `FixNVE`, `FixFreeze`.
- reading/writing data. Example: `FixRestart`.
- adding or modifying properties due to geometry. Example: `FixWall`.
- interacting with other subsystems or external code: Examples: `FixTTM`, `FixExternal`, `FixLATTE`
- saving information for analysis or future use (previous positions, for instance). Examples: `Fix AveTime`, `FixStoreState`.

All fixes are derived from the `Fix` base class and must have a constructor with the signature: `FixPrintVel(class LAMMPS *, int, char **)`.

Every fix must be registered in LAMMPS by writing the following lines of code in the header before include guards:

```
#ifndef FIX_CLASS
// clang-format off
FixStyle(print/vel,FixPrintVel);
// clang-format on
#else
/* the definition of the FixPrintVel class comes here */
...
#endif
```

Where `print/vel` is the style name of your fix in the input script and `FixPrintVel` is the name of the class. The header file would be called `fix_print_vel.h` and the implementation file `fix_print_vel.cpp`. These conventions allow LAMMPS to automatically integrate it into the executable when compiling and associate your new fix class with the designated keyword when it parses the input script.

Let's write a simple fix which will print the average velocity at the end of each timestep. First of all, implement a constructor:

```
FixPrintVel::FixPrintVel(LAMMPS *lmp, int nargs, char **arg)
: Fix(lmp, nargs, arg)
{
  if (nargs < 4)
    error->all(FLERR,"Illegal fix print/vel command");

  nevery = utils::inumeric(FLERR,arg[3],false,lmp);
  if (nevery <= 0)
```

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```

error->all(FLERR,"Illegal fix print/vel command");
}

```

In the constructor you should parse your fix arguments which are specified in the script. All fixes have pretty much the same syntax: `fix <fix-ID> <fix group> <fix name> <fix arguments ...>`. The first 3 parameters are parsed by Fix base class constructor, while `<fix arguments>` should be parsed by you. In our case, we need to specify how often we want to print an average velocity. For instance, once in 50 timesteps: `fix 1 print/vel 50`. There is a special variable in the Fix class called `nevery` which specifies how often the method `end_of_step()` is called. Thus all we need to do is just set it up.

The next method we need to implement is `setmask()`:

```

int FixPrintVel::setmask()
{
    int mask = 0;
    mask |= FixConst::END_OF_STEP;
    return mask;
}

```

Here the user specifies which methods of your fix should be called during execution. The constant `END_OF_STEP` corresponds to the `end_of_step()` method. The most important available methods that are called during a timestep and the order in which they are called are shown in the previous section.

```

void FixPrintVel::end_of_step()
{
    // for add3, scale3
    using namespace MathExtra;

    double** v = atom->v;
    int nlocal = atom->nlocal;
    double localAvgVel[4]; // 4th element for particles count
    memset(localAvgVel, 0, 4 * sizeof(double));
    for (int particleInd = 0; particleInd < nlocal; ++particleInd) {
        add3(localAvgVel, v[particleInd], localAvgVel);
    }
    localAvgVel[3] = nlocal;
    double globalAvgVel[4];
    memset(globalAvgVel, 0, 4 * sizeof(double));
    MPI_Allreduce(localAvgVel, globalAvgVel, 4, MPI_DOUBLE, MPI_SUM, world);
    scale3(1.0 / globalAvgVel[3], globalAvgVel);
    if ((comm->me == 0) && screen) {
        fmt::print(screen, "{}, {}, {}\n",
                  globalAvgVel[0], globalAvgVel[1], globalAvgVel[2]);
    }
}

```

In the code above, we use `MathExtra` routines defined in `math_extra.h`. There are bunch of math functions to work with arrays of doubles as with math vectors. It is also important to note that LAMMPS code should always assume to be run in parallel and that atom data is thus distributed across the MPI ranks. Thus you can only process data from local atoms directly and need to use MPI library calls to combine or exchange data. For serial execution, LAMMPS comes bundled with the MPI STUBS library that contains the MPI library function calls in dummy versions that only work for a single MPI rank.

In this code we use an instance of `Atom` class. This object is stored in the `Pointers` class (see `pointers.h`) which is

the base class of the Fix base class. This object contains references to various class instances (the original instances are created and held by the LAMMPS class) with all global information about the simulation system. Data from the Pointers class is available to all classes inherited from it using protected inheritance. Hence when you write your own class, which is going to use LAMMPS data, don't forget to inherit from Pointers or pass an Pointer to it to all functions that need access. When writing fixes we inherit from class Fix which is inherited from Pointers so there is no need to inherit from it directly.

The code above computes average velocity for all particles in the simulation. Yet you have one unused parameter in fix call from the script: `group_name`. This parameter specifies the group of atoms used in the fix. So we should compute average for all particles in the simulation only if `group_name == "all"`, but it can be any group. The group membership information of an atom is contained in the `mask` property of an atom and the bit corresponding to a given group is stored in the `groupbit` variable which is defined in Fix base class:

```
for (int i = 0; i < nlocal; ++i) {
  if (atom->mask[i] & groupbit) {
    // Do all job here
  }
}
```

Class Atom encapsulates atoms positions, velocities, forces, etc. User can access them using particle index. Note, that particle indexes are usually changed every few timesteps because of neighbor list rebuilds and spatial sorting (to improve cache efficiency).

Let us consider another Fix example: We want to have a fix which stores atoms position from previous time step in your fix. The local atoms indexes may not be valid on the next iteration. In order to handle this situation there are several methods which should be implemented:

- `double memory_usage()`: return how much memory the fix uses (optional)
- `void grow_arrays(int)`: do reallocation of the per particle arrays in your fix
- `void copy_arrays(int i, int j, int delflag)`: copy i-th per-particle information to j-th. Used when atom sorting is performed. if `delflag` is set and atom `j` owns a body, move the body information to atom `i`.
- `void set_arrays(int i)`: sets i-th particle related information to zero

Note, that if your class implements these methods, it must call `add_callback` and `delete_callback` to constructor and destructor. Since we want to store positions of atoms from previous timestep, we need to add `double** xold` to the header file. Then add allocation code to the constructor:

```
FixSavePos::FixSavePos(LAMMPS *lmp, int narg, char **arg), xold(nullptr)
{
  //...
  memory->create(xold, atom->nmax, 3, "FixSavePos:x");
  atom->add_callback(0);
}

FixSavePos::~FixSavePos() {
  atom->delete_callback(id, 0);
  memory->destroy(xold);
}
```

Implement the aforementioned methods:

```
double FixSavePos::memory_usage()
{
  int nmax = atom->nmax;
```

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```

    double bytes = 0.0;
    bytes += nmax * 3 * sizeof(double);
    return bytes;
}

void FixSavePos::grow_arrays(int nmax)
{
    memory->grow(xold, nmax, 3, "FixSavePos:xold");
}

void FixSavePos::copy_arrays(int i, int j, int delflag)
{
    memcpy(xold[j], xold[i], sizeof(double) * 3);
}

void FixSavePos::set_arrays(int i)
{
    memset(xold[i], 0, sizeof(double) * 3);
}

int FixSavePos::pack_exchange(int i, double *buf)
{
    int m = 0;
    buf[m++] = xold[i][0];
    buf[m++] = xold[i][1];
    buf[m++] = xold[i][2];

    return m;
}

int FixSavePos::unpack_exchange(int nlocal, double *buf)
{
    int m = 0;
    xold[nlocal][0] = buf[m++];
    xold[nlocal][1] = buf[m++];
    xold[nlocal][2] = buf[m++];

    return m;
}

```

Now, a little bit about memory allocation. We use the Memory class which is just a bunch of template functions for allocating 1D and 2D arrays. So you need to add include `memory.h` to have access to them.

Finally, if you need to write/read some global information used in your fix to the restart file, you might do it by setting flag `restart_global = 1` in the constructor and implementing methods `void write_restart(FILE *fp)` and `void restart(char *buf)`. If, in addition, you want to write the per-atom property to restart files additional settings and functions are needed:

- a fix flag indicating this needs to be set `restart_peratom = 1`;
- `atom->add_callback()` and `atom->delete_callback()` must be called a second time with the final argument set to 1 instead of 0 (indicating restart processing instead of per-atom data memory management).
- the functions `void pack_restart(int i, double *buf)` and `void unpack_restart(int nlocal, int nth)` need to be implemented

## 15.8 Notes for developers and code maintainers

This section documents how some of the code functionality within LAMMPS works at a conceptual level. Comments on code in source files typically document what a variable stores, what a small section of code does, or what a function does and its input/outputs. The topics on this page are intended to document code functionality at a higher level.

Available topics are:

- *Reading and parsing of text and text files*
  - *Requesting and accessing neighbor lists*
  - *Fix contributions to instantaneous energy, virial, and cumulative energy*
  - *KSpace PPPM FFT grids*
- 

### 15.8.1 Reading and parsing of text and text files

It is frequently required for a class in LAMMPS to read in additional data from a file, e.g. potential parameters from a potential file for manybody potentials. LAMMPS provides several custom classes and convenience functions to simplify the process. They offer the following benefits:

- better code reuse and fewer lines of code needed to implement reading and parsing data from a file
- better detection of format errors, incompatible data, and better error messages
- exit with an error message instead of silently converting only part of the text to a number or returning a 0 on unrecognized text and thus reading incorrect values
- re-entrant code through avoiding global static variables (as used by `strtok()`)
- transparent support for translating unsupported UTF-8 characters to their ASCII equivalents (the text-to-value conversion functions **only** accept ASCII characters)

In most cases (e.g. potential files) the same data is needed on all MPI ranks. Then it is best to do the reading and parsing only on MPI rank 0, and communicate the data later with one or more `MPI_Bcast()` calls. For reading generic text and potential parameter files the custom classes *TextFileReader* and *PotentialFileReader* are available. They allow reading the file as individual lines for which they can return a tokenizer class (see below) for parsing the line. Or they can return blocks of numbers as a vector directly. The documentation on *File reader classes* contains an example for a typical case.

When reading per-atom data, the data on each line of the file usually needs to include an atom ID so it can be associated with a particular atom. In that case the data can be read in multi-line chunks and broadcast to all MPI ranks with `utils::read_lines_from_file()`. Those chunks are then split into lines, parsed, and applied only to atoms the MPI rank “owns”.

For splitting a string (incrementally) into words and optionally converting those to numbers, the *Tokenizer* and *ValueTokenizer* can be used. Those provide a superset of the functionality of `strtok()` from the C-library and the latter also includes conversion to different types. Any errors while processing the string in those classes will result in an exception, which can be caught and the error processed as needed. Unlike the C-library functions `atoi()`, `atof()`, `strtol()`, or `strtod()` the conversion will check if the converted text is a valid integer or floating point number and will not silently return an unexpected or incorrect value. For example, `atoi()` will return 12 when converting “12.5”, while the *ValueTokenizer* class will throw an *InvalidIntegerException* if `ValueTokenizer::next_int()` is called on the same string.

## 15.8.2 Requesting and accessing neighbor lists

LAMMPS uses Verlet-style neighbor lists to avoid having to loop over *all* pairs of *all* atoms when computing pairwise properties with a cutoff (e.g. pairwise forces or radial distribution functions). There are three main algorithms that can be selected by the *neighbor command*: *bin* (the default, uses binning to achieve linear scaling with system size), *nsq* (without binning, quadratic scaling), *multi* (with binning, optimized for varying cutoffs or polydisperse granular particles). In addition to how the neighbor lists are constructed a number of different variants of neighbor lists need to be created (e.g. “full” or “half”) for different purposes and styles and those may be required in every time step (“perpetual”) or on some steps (“occasional”).

The neighbor list creation is managed by the `Neighbor` class. Individual classes can obtain a neighbor list by creating an instance of a `NeighRequest` class which is stored in a list inside the `Neighbor` class. The `Neighbor` class will then analyze the various requests and apply optimizations where neighbor lists that have the same settings will be created only once and then copied, or a list may be constructed by processing a neighbor list from a different request that is a superset of the requested list. The neighbor list build is then *processed in parallel*.

The most commonly required neighbor list is a so-called “half” neighbor list, where each pair of atoms is listed only once (except when the *newton command setting* for pair is off; in that case pairs straddling sub-domains or periodic boundaries will be listed twice). Thus these are the default settings when a neighbor list request is created in:

```
void Pair::init_style()
{
  neighbor->add_request(this);
}

void Pair::init_list(int /*id*/, NeighList *ptr)
{
  list = ptr;
}
```

The `this` pointer argument is required so the neighbor list code can access the requesting class instance to store the assembled neighbor list with that instance by calling its `init_list()` member function. The optional second argument (omitted here) contains a bitmask of flags that determines the kind of neighbor list requested. The default value used here asks for a perpetual “half” neighbor list.

Non-default values of the second argument need to be used to adjust a neighbor list request to the specific needs of a style an additional request flag is needed. The *tersoff* pair style, for example, needs a “full” neighbor list:

```
void PairTersoff::init_style()
{
  // [...]
  neighbor->add_request(this, NeighConst::REQ_FULL);
}
```

When a pair style supports r-RESPA time integration with different cutoff regions, the request flag may depend on the corresponding r-RESPA settings. Here an example from pair style `lj/cut`:

```
void PairLJCut::init_style()
{
  int list_style = NeighConst::REQ_DEFAULT;

  if (update->whichflag == 1 && utils::strmatch(update->integrate_style, "^respa")) {
    auto respa = (Respa *) update->integrate;
    if (respa->level_inner >= 0) list_style = NeighConst::REQ_RESPA_INOUT;
    if (respa->level_middle >= 0) list_style = NeighConst::REQ_RESPA_ALL;
  }
}
```

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```

}
neighbor->add_request(this, list_style);
// [...]
}

```

Granular pair styles need neighbor lists based on particle sizes and not cutoff and also may require to have the list of previous neighbors available (“history”). For example with:

```

if (use_history) neighbor->add_request(this, NeighConst::REQ_SIZE | NeighConst::REQ_
↪HISTORY);
else neighbor->add_request(this, NeighConst::REQ_SIZE);

```

In case a class would need to make multiple neighbor list requests with different settings each request can set an id which is then used in the corresponding `init_list()` function to assign it to the suitable pointer variable. This is done for example by the *pair style meam*:

```

void PairMEAM::init_style()
{
// [...]
neighbor->add_request(this, NeighConst::REQ_FULL)->set_id(1);
neighbor->add_request(this)->set_id(2);
}
void PairMEAM::init_list(int id, NeighList *ptr)
{
if (id == 1) listfull = ptr;
else if (id == 2) listhalf = ptr;
}

```

Fixes may require a neighbor list that is only build occasionally (or just once) and this can also be indicated by a flag. As an example here is the request from the `FixPeriNeigh` class which is created internally by *Peridynamics pair styles*:

```

neighbor->add_request(this, NeighConst::REQ_FULL | NeighConst::REQ_OCCASIONAL);

```

It is also possible to request a neighbor list that uses a different cutoff than what is usually inferred from the pair style settings (largest cutoff of all pair styles plus neighbor list skin). The following is used in the *compute rdf* command implementation:

```

if (cutflag)
neighbor->add_request(this, NeighConst::REQ_OCCASIONAL)->set_cutoff(mycutneigh);
else
neighbor->add_request(this, NeighConst::REQ_OCCASIONAL);

```

The neighbor list request function has a slightly different set of arguments when created by a command style. In this case the neighbor list is *always* an occasional neighbor list, so that flag is not needed. However for printing the neighbor list summary the name of the requesting command should be set. Below is the request from the *delete atoms* command:

```

neighbor->add_request(this, "delete_atoms", NeighConst::REQ_FULL);

```

### 15.8.3 Fix contributions to instantaneous energy, virial, and cumulative energy

Fixes can calculate contributions to the instantaneous energy and/or virial of the system, both in a global and peratom sense. Fixes that perform thermostating or barostating can calculate the cumulative energy they add to or subtract from the system, which is accessed by the *ecouple* and *econserve* thermodynamic keywords. This subsection explains how both work and what flags to set in a new fix to enable this functionality.

Let's start with thermostating and barostating fixes. Examples are the *fix langevin* and *fix npt* commands. Here is what the fix needs to do:

- Set the variable *ecouple\_flag* = 1 in the constructor. Also set *scalar\_flag* = 1, *extscalar* = 1, and *global\_freq* to a timestep increment which matches how often the fix is invoked.
- Implement a *compute\_scalar()* method that returns the cumulative energy added or subtracted by the fix, e.g. by rescaling the velocity of atoms. The sign convention is that subtracted energy is positive, added energy is negative. This must be the total energy added to the entire system, i.e. an “extensive” quantity, not a per-atom energy. Cumulative means the summed energy since the fix was instantiated, even across multiple runs. This is because the energy is used by the *econserve* thermodynamic keyword to check that the fix is conserving the total energy of the system, i.e. potential energy + kinetic energy + coupling energy = a constant.

And here is how the code operates:

- The *Modify* class makes a list of all fixes that set *ecouple\_flag* = 1.
- The *thermo\_style custom* command defines *ecouple* and *econserve* keywords.
- These keywords sum the energy contributions from all the *ecouple\_flag* = 1 fixes by invoking the *energy\_couple()* method in the *Modify* class, which calls the *compute\_scalar()* method of each fix in the list.

Next, here is how a fix contributes to the instantaneous energy and virial of the system. First, it sets any or all of these flags to a value of 1 in their constructor:

- *energy\_global\_flag* to contribute to global energy, example: *fix indent*
- *energy\_peratom\_flag* to contribute to peratom energy, *fix cmap*
- *virial\_global\_flag* to contribute to global virial, example: *fix wall*
- *virial\_peratom\_flag* to contribute to peratom virial, example: *fix wall*

The fix must also do the following:

- For global energy, implement a *compute\_scalar()* method that returns the energy added or subtracted on this timestep. Here the sign convention is that added energy is positive, subtracted energy is negative.
- For peratom energy, invoke the *ev\_init(eflag,vflag)* function each time the fix is invoked, which initializes peratom energy storage. The value of *eflag* may need to be stored from an earlier call to the fix during the same timestep. See how the *fix cmap* command does this in *src/MOLECULE/fix\_cmap.cpp*. When an energy for one or more atoms is calculated, invoke the *ev\_tally()* function to tally the contribution to each atom. Both the *ev\_init()* and *ev\_tally()* methods are in the parent *Fix* class.
- For global and/or peratom virial, invoke the *v\_init(vflag)* function each time the fix is invoked, which initializes virial storage. When forces on one or more atoms are calculated, invoke the *v\_tally()* function to tally the contribution. Both the *v\_init()* and *v\_tally()* methods are in the parent *Fix* class. Note that there are several variants of *v\_tally()*; choose the one appropriate to your fix.

**Note:** The *ev\_init()* and *ev\_tally()* methods also account for global and peratom virial contributions. Thus you do not need to invoke the *v\_init()* and *v\_tally()* methods, if the fix also calculates peratom energies.

The fix must also specify whether (by default) to include or exclude these contributions to the global/peratom energy/virial of the system. For the fix to include the contributions, set either of both of these variables in the constructor:

- *thermo\_energy* = 1, for global and peratom energy
- *thermo\_virial* = 1, for global and peratom virial

Note that these variables are zeroed in `fix.cpp`. Thus if you don't set the variables, the contributions will be excluded (by default)

However, the user has ultimate control over whether to include or exclude the contributions of the fix via the *fix modify* command:

- `fix modify energy yes` to include global and peratom energy contributions
- `fix modify virial yes` to include global and peratom virial contributions

If the fix contributes to any of the global/peratom energy/virial values for the system, it should be explained on the fix doc page, along with the default values for the *energy yes/no* and *virial yes/no* settings of the *fix modify* command.

Finally, these 4 contributions are included in the output of 4 computes:

- global energy in *compute pe*
- peratom energy in *compute pe/atom*
- global virial in *compute pressure*
- peratom virial in *compute stress/atom*

These computes invoke a method of the `Modify` class to include contributions from fixes that have the corresponding flags set, e.g. *energy\_peratom\_flag* and *thermo\_energy* for *compute pe/atom*.

Note that each compute has an optional keyword to either include or exclude all contributions from fixes. Also note that *compute pe* and *compute pressure* are what is used (by default) by *thermodynamic output* to calculate values for its *pe* and *press* keywords.

## 15.8.4 KSpace PPPM FFT grids

The various *KSpace PPPM* styles in LAMMPS use FFTs to solve Poisson's equation. This subsection describes:

- how FFT grids are defined
- how they are decomposed across processors
- how they are indexed by each processor
- how particle charge and electric field values are mapped to/from the grid

An FFT grid cell is a 3d volume; grid points are corners of a grid cell and the code stores values assigned to grid points in vectors or 3d arrays. A global 3d FFT grid has points indexed 0 to N-1 inclusive in each dimension.

Each processor owns two subsets of the grid, each subset is brick-shaped. Depending on how it is used, these subsets are allocated as a 1d vector or 3d array. Either way, the ordering of values within contiguous memory is fastest, then y, z slowest.

For the 3d decomposition of the grid, the global grid is partitioned into bricks that correspond to the sub-domains of the simulation box that each processor owns. Often, this is a regular 3d array ( $P_x$  by  $P_y$  by  $P_z$ ) of bricks, where  $P$  = number of processors =  $P_x * P_y * P_z$ . More generally it can be a tiled decomposition, where each processor owns a brick and the union of all the bricks is the global grid. Tiled decompositions are produced by load balancing with the RCB algorithm; see the *balance rcb* command.

For the FFT decomposition of the grid, each processor owns a brick that spans the entire x dimension of the grid while the y and z dimensions are partitioned as a regular 2d array ( $P_1$  by  $P_2$ ), where  $P = P_1 * P_2$ .



The following indices store the inclusive bounds of the brick a processor owns, within the global grid:

```
nxlo_in,nxhi_in,nylo_in,nyhi_in,nzlo_in,nzhi_in = 3d decomposition brick
nxlo_fft,nxhi_fft,nylo_fft,nyhi_fft,nzlo_fft,nzhi_fft = FFT decomposition brick
nxlo_out,nxhi_out,nylo_out,nyhi_out,nzlo_out,nzhi_out = 3d decomposition brick + ghost
↪cells
```

The `in` and `fft` indices are from 0 to N-1 inclusive in each dimension, where N is the grid size.

The `out` indices index an array which stores the `in` subset of the grid plus ghost cells that surround it. These indices can thus be  $< 0$  or  $\geq N$ .

The number of ghost cells a processor owns in each of the 6 directions is a function of:

```
neighbor skin distance (since atoms can move outside a proc subdomain)
qdist = offset or charge from atom due to TIP4P fictitious charge
order = mapping stencil size
shift = factor used when order is an even number (see below)
```

Here is an explanation of how the PPPM variables `order`, `nlower` / `nupper`, `shift`, and `OFFSET` work. They are the relevant variables that determine how atom charge is mapped to grid points and how field values are mapped from grid points to atoms:

```
order = # of nearby grid points in each dim that atom charge/field are mapped to/from
nlower,nupper = extent of stencil around the grid point an atom is assigned to
OFFSET = large integer added/subtracted when mapping to avoid  $\text{int}(-0.75) = 0$  when  $-1$  is
↪the desired result
```

The `particle_map()` method assigns each atom to a grid point.

If `order` is even, say 4:

```
atom is assigned to grid point to its left (in each dim)
shift = OFFSET
nlower = -1, nupper = 2, which are offsets from assigned grid point
window of mapping grid pts is thus 2 grid points to left of atom, 2 to right
```

If `order` is odd, say 5:

```
atom is assigned to left/right grid pt it is closest to (in each dim)
shift = OFFSET + 0.5
nlower = 2, nupper = 2
if point is in left half of cell, then window of affected grid pts is 3 grid points to
↪left of atom, 2 to right
if point is in right half of cell, then window of affected grid pts is 2 grid points to
↪left of atom, 3 to right
```

These settings apply to each dimension, so that if `order` = 5, an atom's charge is mapped to 125 grid points that surround the atom.

## 15.9 Notes for updating code written for older LAMMPS versions

This section documents how C++ source files that are available *outside of the LAMMPS source distribution* (e.g. in external USER packages or as source files provided as a supplement to a publication) that are written for an older version of LAMMPS and thus need to be updated to be compatible with the current version of LAMMPS. Due to the active development of LAMMPS it is likely to always be incomplete. Please contact [developer@lammps.org](mailto:developer@lammps.org) in case you run across an issue that is not (yet) listed here. Please also review the latest information about the LAMMPS *programming style conventions*, especially if you are considering to submit the updated version for inclusion into the LAMMPS distribution.

Available topics in mostly chronological order are:

- *Setting flags in the constructor*
  - *Rename of `pack/unpack_comm()` to `pack/unpack_forward_comm()`*
  - *Use `ev_init()` to initialize variables derived from `eflag` and `vflag`*
  - *Use `utils::numeric()` functions instead of `force->numeric()`*
  - *Use `utils::open_potential()` function to open potential files*
  - *Simplify customized error messages*
  - *Use of “override” instead of “virtual”*
  - *Simplified and more compact neighbor list requests*
  - *Split of fix STORE into fix STORE/GLOBAL and fix STORE/PERATOM*
  - *Use `Output::get_dump_by_id()` instead of `Output::find_dump()`*
- 

### 15.9.1 Setting flags in the constructor

As LAMMPS gains additional functionality, new flags may need to be set in the constructor or a class to signal compatibility with such features. Most of the time the defaults are chosen conservatively, but sometimes the conservative choice is the uncommon choice, and then those settings need to be made when updating code.

Pair styles:

- `manybody_flag`: set to 1 if your pair style is not pair-wise additive
- `restartinfo`: set to 0 if your pair style does not store data in restart files

### 15.9.2 Rename of `pack/unpack_comm()` to `pack/unpack_forward_comm()`

Changed in version 8Aug2014.

In this change set the functions to pack data into communication buffers and to unpack data from communication buffers for *forward communications* were renamed from `pack_comm()` and `unpack_comm()` to `pack_forward_comm()` and `unpack_forward_comm()`, respectively. Also the meaning of the return value of these functions was changed: rather than returning the number of items per atom stored in the buffer, now the total number of items added (or unpacked) needs to be returned. Here is an example from the *PairEAM* class. Of course the member function declaration in corresponding header file needs to be updated accordingly.

Old:

```

int PairEAM::pack_comm(int n, int *list, double *buf, int pbc_flag, int *pbc)
{
  int m = 0;
  for (int i = 0; i < n; i++) {
    int j = list[i];
    buf[m++] = fp[j];
  }
  return 1;
}

```

New:

```

int PairEAM::pack_forward_comm(int n, int *list, double *buf, int pbc_flag, int *pbc)
{
  int m = 0;
  for (int i = 0; i < n; i++) {
    int j = list[i];
    buf[m++] = fp[j];
  }
  return m;
}

```

**Note:** Because the various “pack” and “unpack” functions are defined in the respective base classes as dummy functions doing nothing, and because of the the name mismatch the custom versions in the derived class will no longer be called, there will be no compilation error when this change is not applied. Only calculations will suddenly produce incorrect results because the required forward communication calls will cease to function correctly.

### 15.9.3 Use `ev_init()` to initialize variables derived from `eflag` and `vflag`

Changed in version 29Mar2019.

There are several variables that need to be initialized based on the values of the “eflag” and “vflag” variables and since sometimes there are new bits added and new variables need to be set to 1 or 0. To make this consistent, across all styles, there is now an inline function `ev_init(eflag, vflag)` that makes those settings consistently and calls either `ev_setup()` or `ev_unset()`. Example from a pair style:

Old:

```

if (eflag || vflag) ev_setup(eflag, vflag);
else evflag = vflag_fdotr = eflag_global = eflag_atom = 0;

```

New:

```

ev_init(eflag, vflag);

```

Not applying this change will not cause a compilation error, but can lead to inconsistent behavior and incorrect tallying of energy or virial.

## 15.9.4 Use `utils::numeric()` functions instead of `force->numeric()`

Changed in version 18Sep2020.

The “`numeric()`” conversion functions (including “`inumeric()`”, “`bnumeric()`”, and “`tnumeric()`”) have been moved from the `Force` class to the `utils` namespace. Also they take an additional argument that selects whether the `Error::all()` or `Error::one()` function should be called in case of an error. The former should be used when *all* MPI processes call the conversion function and the latter *must* be used when they are called from only one or a subset of the MPI processes.

Old:

```
val = force->numeric(FLERR, arg[1]);
num = force->inumeric(FLERR, arg[2]);
```

New:

```
val = utils::numeric(FLERR, true, arg[1], lmp);
num = utils::inumeric(FLERR, false, arg[2], lmp);
```

**See also:**

`utils::numeric()`, `utils::inumeric()`, `utils::bnumeric()`, `utils::tnumeric()`

## 15.9.5 Use `utils::open_potential()` function to open potential files

Changed in version 18Sep2020.

The `utils::open_potential()` function must be used to replace calls to `force->open_potential()` and should be used to replace `fopen()` for opening potential files for reading. The custom function does three additional steps compared to `fopen()`: 1) it will try to parse the `UNITS:` and `DATE:` metadata will stop with an error on a units mismatch and will print the date info, if present, in the log file; 2) for pair styles that support it, it will set up possible automatic unit conversions based on the embedded unit information and LAMMPS’ current units setting; 3) it will not only try to open a potential file at the given path, but will also search in the folders listed in the `LAMMPS_POTENTIALS` environment variable. This allows to keep potential files in a common location instead of having to copy them around for simulations.

Old:

```
fp = force->open_potential(filename);
fp = fopen(filename, "r");
```

New:

```
fp = utils::open_potential(filename, lmp);
```

## 15.9.6 Simplify customized error messages

Changed in version 14May2021.

Aided by features of the bundled `{fmt}` library, error messages now can have a variable number of arguments and the string will be interpreted as a `{fmt}` style format string so that custom error messages can be easily customized without having to use temporary buffers and `sprintf()`. Example:

Old:

```

if (fptr == NULL) {
    char str[128];
    sprintf(str, "Cannot open AEAM potential file %s", filename);
    error->one(FLERR, str);
}

```

New:

```

if (fptr == nullptr)
    error->one(FLERR, "Cannot open AEAM potential file {}: {}", filename,
↳utils::getsyserror());

```

### 15.9.7 Use of “override” instead of “virtual”

Changed in version 17Feb2022.

Since LAMMPS requires C++11 we switched to use the “override” keyword instead of “virtual” to indicate polymorphism in derived classes. This allows the C++ compiler to better detect inconsistencies when an override is intended or not. Please note that “override” has to be added to **all** polymorph functions in derived classes and “virtual” *only* to the function in the base class (or the destructor). Here is an example from the `FixWallReflect` class:

Old:

```

FixWallReflect(class LAMMPS *, int, char **);
virtual ~FixWallReflect();
int setmask();
void init();
void post_integrate();

```

New:

```

FixWallReflect(class LAMMPS *, int, char **);
~FixWallReflect() override;
int setmask() override;
void init() override;
void post_integrate() override;

```

This change set will neither cause a compilation failure, nor will it change functionality, but if you plan to submit the updated code for inclusion into the LAMMPS distribution, it will be requested for achieve a consistent *programming style*.

### 15.9.8 Simplified function names for forward and reverse communication

Changed in version 24Mar2022.

Rather than using the function name to distinguish between the different forward and reverse communication functions for styles, LAMMPS now uses the type of the “this” pointer argument.

Old:

```

comm->forward_comm_pair(this);
comm->forward_comm_fix(this);
comm->forward_comm_compute(this);

```

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```
comm->forward_comm_dump(this);  
comm->reverse_comm_pair(this);  
comm->reverse_comm_fix(this);  
comm->reverse_comm_compute(this);  
comm->reverse_comm_dump(this);
```

New:

```
comm->forward_comm(this);  
comm->reverse_comm(this);
```

This change is **required** or else the code will not compile.

## 15.9.9 Simplified and more compact neighbor list requests

Changed in version 24Mar2022.

This change set reduces the amount of code required to request a neighbor list. It enforces consistency and no longer requires to change internal data of the request. More information on neighbor list requests can be [found here](#). Example from the ComputeRDF class:

Old:

```
int irequest = neighbor->request(this,instance_me);  
neighbor->requests[irequest]->pair = 0;  
neighbor->requests[irequest]->compute = 1;  
neighbor->requests[irequest]->occasional = 1;  
if (cutflag) {  
    neighbor->requests[irequest]->cut = 1;  
    neighbor->requests[irequest]->cutoff = mycutneigh;  
}
```

New:

```
auto req = neighbor->add_request(this, NeighConst::REQ_OCCASIONAL);  
if (cutflag) req->set_cutoff(mycutneigh);
```

Public access to the NeighRequest class data members has been removed so this update is **required** to avoid compilation failure.

## 15.9.10 Split of fix STORE into fix STORE/GLOBAL and fix STORE/PERATOM

Changed in version 15Sep2022.

This change splits the GLOBAL and PERATOM modes of fix STORE into two separate fixes STORE/GLOBAL and STORE/PERATOM. There was very little shared code between the two fix STORE modes and the two different code paths had to be prefixed with if statements. Furthermore, some flags were used differently in the two modes leading to confusion. Splitting the code into two fix styles, makes it more easily maintainable. Since these are internal fixes, there is no user visible change.

Old:

```
#include "fix_store.h"

FixStore *fix = dynamic_cast<FixStore *>(
  modify->add_fix(fmt::format("{} {} STORE peratom 1 13",id_pole,group->names[0]));

FixStore *fix = dynamic_cast<FixStore *>(modify->get_fix_by_id(id_pole));
```

New:

```
#include "fix_store_peratom.h"

FixStorePeratom *fix = dynamic_cast<FixStorePeratom *>(
  modify->add_fix(fmt::format("{} {} STORE/PERATOM 1 13",id_pole,group->names[0]));

FixStorePeratom *fix = dynamic_cast<FixStorePeratom *>(modify->get_fix_by_id(id_pole));
```

Old:

```
#include "fix_store.h"

FixStore *fix = dynamic_cast<FixStore *>(
  modify->add_fix(fmt::format("{} {} STORE global 1 1",id_fix,group->names[igroup]));

FixStore *fix = dynamic_cast<FixStore *>(modify->get_fix_by_id(id_fix));
```

New:

```
#include "fix_store_global.h"

FixStoreGlobal *fix = dynamic_cast<FixStoreGlobal *>(
  modify->add_fix(fmt::format("{} {} STORE/GLOBAL 1 1",id_fix,group->names[igroup]));

FixStoreGlobal *fix = dynamic_cast<FixStoreGlobal *>(modify->get_fix_by_id(id_fix));
```

This change is **required** or else the code will not compile.

### 15.9.11 Use Output::get\_dump\_by\_id() instead of Output::find\_dump()

Changed in version 15Sep2022.

The accessor function to individual dump style instances has been changed from Output::find\_dump() returning the index of the dump instance in the list of dumps to Output::get\_dump\_by\_id() returning a pointer to the dump directly. Example:

Old:

```
int idump = output->find_dump(arg[iarg+1]);
if (idump < 0)
  error->all(FLERR, "Dump ID in hyper command does not exist");
memory->grow(dumplist,ndump+1,"hyper:dumplist");
dumplist[ndump++] = idump;

[...]
```

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```

if (dumpflag)
  for (int idump = 0; idump < ndump; idump++)
    output->dump[dumplist[idump]]->write();

```

New:

```

auto idump = output->get_dump_by_id(arg[iarg+1]);
if (!idump) error->all(FLERR, "Dump ID {} in hyper command does not exist", arg[iarg+1]);
dumplist.emplace_back(idump);

[...]

if (dumpflag) for (auto idump : dumplist) idump->write();

```

This change is **required** or else the code will not compile.

## 15.10 Writing plugins

Plugins provide a mechanism to add functionality to a LAMMPS executable without recompiling LAMMPS. The functionality for this and the *plugin command* are implemented in the *PLUGIN package* which must be installed to use plugins.

Plugins use the operating system's capability to load dynamic shared object (DSO) files in a way similar shared libraries and then reference specific functions in those DSOs. Any DSO file with plugins has to include an initialization function with a specific name, "lammplugin\_init", that has to follow specific rules described below. When loading the DSO with the "plugin" command, this function is looked up and called and will then register the contained plugin(s) with LAMMPS.

When the environment variable LAMMPS\_PLUGIN\_PATH is set, then LAMMPS will search the directory (or directories) listed in this path for files with names that end in *plugin.so* (e.g. *helloplugin.so*) and will try to load the contained plugins automatically at start-up. For plugins that are loaded this way, the behavior of LAMMPS should be identical to a binary where the corresponding code was compiled in statically as a package.

From the programmer perspective this can work because of the object oriented design of LAMMPS where all pair style commands are derived from the class *Pair*, all fix style commands from the class *Fix* and so on and usually only functions present in those base classes are called directly. When a *pair\_style command* or *fix command* is issued a new instance of such a derived class is created. This is done by a so-called factory function which is mapped to the style name. Thus when, for example, the LAMMPS processes the command `pair_style lj/cut 2.5`, LAMMPS will look up the factory function for creating the *PairLJCut* class and then execute it. The return value of that function is a *Pair \** pointer and the pointer will be assigned to the location for the currently active pair style.

A DSO file with a plugin thus has to implement such a factory function and register it with LAMMPS so that it gets added to the map of available styles of the given category. To register a plugin with LAMMPS an initialization function has to be present in the DSO file called `lammplugin_init` which is called with three `void *` arguments: a pointer to the current LAMMPS instance, a pointer to the opened DSO handle, and a pointer to the registration function. The registration function takes two arguments: a pointer to a `lammplugin_t` struct with information about the plugin and a pointer to the current LAMMPS instance. Please see below for an example of how the registration is done.



### 15.10.1 Members of `lammplugin_t`

Member	Description
<code>version</code>	LAMMPS Version string the plugin was compiled for
<code>style</code>	Style of the plugin (pair, bond, fix, command, etc.)
<code>name</code>	Name of the plugin style
<code>info</code>	String with information about the plugin
<code>author</code>	String with the name and email of the author
<code>creator.v1</code>	Pointer to factory function for pair, bond, angle, dihedral, improper, kspace, or command styles
<code>creator.v2</code>	Pointer to factory function for compute, fix, or region styles
<code>handle</code>	Pointer to the open DSO file handle

Only one of the two alternate creator entries can be used at a time and which of those is determined by the style of plugin. The “creator.v1” element is for factory functions of supported styles computing forces (i.e. pair, bond, angle, dihedral, or improper styles) or command styles and the function takes as single argument the pointer to the LAMMPS instance. The factory function is cast to the `lammplugin_factory1` type before assignment. The “creator.v2” element is for factory functions creating an instance of a fix, compute, or region style and takes three arguments: a pointer to the LAMMPS instance, an integer with the length of the argument list and a `char **` pointer to the list of arguments. The factory function pointer needs to be cast to the `lammplugin_factory2` type before assignment.

### 15.10.2 Pair style example

As an example, a hypothetical pair style plugin “morse2” implemented in a class `PairMorse2` in the files `pair_morse2.h` and `pair_morse2.cpp` with the factory function and initialization function would look like this:

```
#include "lammplugin.h"
#include "version.h"
#include "pair_morse2.h"

using namespace LAMMPS_NS;

static Pair *morse2creator(LAMMPS *lmp)
{
    return new PairMorse2(lmp);
}

extern "C" void lammplugin_init(void *lmp, void *handle, void *regfunc)
{
    lammplugin_regfunc register_plugin = (lammplugin_regfunc) regfunc;
    lammplugin_t plugin;

    plugin.version = LAMMPS_VERSION;
    plugin.style   = "pair";
    plugin.name    = "morse2";
    plugin.info    = "Morse2 variant pair style v1.0";
    plugin.author  = "Axel Kohlmeyer (akohlmey@gmail.com)";
    plugin.creator.v1 = (lammplugin_factory1 *) &morse2creator;
    plugin.handle  = handle;
    (*register_plugin)(amp;plugin, lmp);
}
```

The factory function in this example is called `morse2creator()`. It receives a pointer to the LAMMPS class

as only argument and thus has to be assigned to the *creator.v1* member of the plugin struct and cast to the `lammplugin_factory1` function pointer type. It returns a pointer to the allocated class instance derived from the `Pair` class. This function may be declared static to avoid clashes with other plugins. The name of the derived class, `PairMorse2`, however must be unique inside the entire LAMMPS executable.

### 15.10.3 Fix style example

If the factory function would be for a fix or compute, which take three arguments (a pointer to the LAMMPS class, the number of arguments and the list of argument strings), then the pointer type is `lammplugin_factory2` and it must be assigned to the *creator.v2* member of the plugin struct. Below is an example for that:

```
#include "lammplugin.h"
#include "version.h"
#include "fix_nve2.h"

using namespace LAMMPS_NS;

static Fix *nve2creator(LAMMPS *lmp, int argc, char **argv)
{
    return new FixNVE2(lmp,argc,argv);
}

extern "C" void lammplugin_init(void *lmp, void *handle, void *regfunc)
{
    lammplugin_regfunc register_plugin = (lammplugin_regfunc) regfunc;
    lammplugin_t plugin;

    plugin.version = LAMMPS_VERSION;
    plugin.style = "fix";
    plugin.name = "nve2";
    plugin.info = "NVE2 variant fix style v1.0";
    plugin.author = "Axel Kohlmeyer (akohlmey@gmail.com)";
    plugin.creator.v2 = (lammplugin_factory2 *) &nve2creator;
    plugin.handle = handle;
    (*register_plugin)(&plugin,lmp);
}
```

### 15.10.4 Command style example

Command styles also use the first variant of factory function as demonstrated in the following example, which also shows that the implementation of the plugin class may be within the same source file as the plugin interface code:

```
#include "lammplugin.h"

#include "comm.h"
#include "error.h"
#include "command.h"
#include "version.h"

#include <cstring>
```

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```

namespace LAMMPS_NS {
  class Hello : public Command {
  public:
    Hello(class LAMMPS *lmp) : Command(lmp) {};
    void command(int, char **);
  };
}

using namespace LAMMPS_NS;

void Hello::command(int argc, char **argv)
{
  if (argc != 1) error->all(FLERR, "Illegal hello command");
  if (comm->me == 0)
    utils::logmesg(lmp, fmt::format("Hello, {}!\n", argv[0]));
}

static void hellocreator(LAMMPS *lmp)
{
  return new Hello(lmp);
}

extern "C" void lammpsplugin_init(void *lmp, void *handle, void *regfunc)
{
  lammpsplugin_t plugin;
  lammpsplugin_regfunc register_plugin = (lammpsplugin_regfunc) regfunc;

  plugin.version = LAMMPS_VERSION;
  plugin.style   = "command";
  plugin.name    = "hello";
  plugin.info    = "Hello world command v1.1";
  plugin.author  = "Axel Kohlmeyer (akohlmey@gmail.com)";
  plugin.creator.v1 = (lammpsplugin_factory1 *) &hellocreator;
  plugin.handle  = handle;
  (*register_plugin)(amp;plugin, lmp);
}

```

### 15.10.5 Additional Details

The initialization function **must** be called `lammpsplugin_init`, it **must** have C bindings and it takes three void pointers as arguments. The first is a pointer to the LAMMPS class that calls it and it needs to be passed to the registration function. The second argument is a pointer to the internal handle of the DSO file, this needs to be added to the plugin info struct, so that the DSO can be closed and unloaded when all its contained plugins are unloaded. The third argument is a function pointer to the registration function and needs to be stored in a variable of `lammpsplugin_regfunc` type and then called with a pointer to the `lammpsplugin_t` struct and the pointer to the LAMMPS instance as arguments to register a single plugin. There may be multiple calls to multiple plugins in the same initialization function.

To register a plugin a struct of the `lammpsplugin_t` needs to be filled with relevant info: current LAMMPS version string, kind of style, name of style, info string, author string, pointer to factory function, and the DSO handle. The registration function is called with a pointer to the address of this struct and the pointer of the LAMMPS class. The registration function will then add the factory function of the plugin style to the respective style map under the provided name. It will also make a copy of the struct in a list of all loaded plugins and update the reference counter for loaded

plugins from this specific DSO file.

The pair style itself (i.e. the PairMorse2 class in this example) can be written just like any other pair style that is included in LAMMPS. For a plugin, the use of the PairStyle macro in the section encapsulated by `#ifdef PAIR_CLASS` is not needed, since the mapping of the class name to the style name is done by the plugin registration function with the information from the `lammpsplugin_t` struct. It may be included in case the new code is intended to be later included in LAMMPS directly.

A plugin may be registered under an existing style name. In that case the plugin will override the existing code. This can be used to modify the behavior of existing styles or to debug new versions of them without having to re-compile or re-install all of LAMMPS.

### 15.10.6 Compiling plugins

Plugins need to be compiled with the same compilers and libraries (e.g. MPI) and compilation settings (MPI on/off, OpenMP, integer sizes) as the LAMMPS executable and library. Otherwise the plugin will likely not load due to mismatches in the function signatures (LAMMPS is C++ so scope, type, and number of arguments are encoded into the symbol names and thus differences in them will lead to failed plugin load commands. Compilation of the plugin can be managed via both, CMake or traditional GNU makefiles. Some examples that can be used as a template are in the `examples/plugins` folder. The CMake script code has some small adjustments to allow building the plugins for running unit tests with them.

Another example that converts the KIM package into a plugin can be found in the `examples/kim/plugin` folder. No changes to the sources of the KIM package themselves are needed; only the plugin interface and loader code needs to be added. This example only supports building with CMake, but is probably a more typical example. To compile you need to run CMake with `-DLAMMPS_SOURCE_DIR=<path/to/lammps/src/folder>`. Other configuration setting are identical to those for compiling LAMMPS.

A second example for a plugin from a package is in the `examples/PACKAGES/pace/plugin` folder that will create a plugin from the ML-PACE package. In this case the bulk of the code is in a static external library that is being downloaded and compiled first and then combined with the pair style wrapper and the plugin loader. This example also contains a NSIS script that can be used to create an Installer package for Windows (the mutual licensing terms of the external library and LAMMPS conflict when distributing binaries, so the ML-PACE package cannot be linked statically, but the LAMMPS headers required to build the plugin are also available under a less restrictive license). This will automatically set the required environment variable and launching a (compatible) LAMMPS binary will load and register the plugin and the ML-PACE package can then be used as it was linked into LAMMPS.

## 15.11 Adding tests for unit testing

This section discusses adding or expanding tests for the unit test infrastructure included into the LAMMPS source code distribution. Unlike example inputs, unit tests focus on testing the “local” behavior of individual features, tend to run fast, and should be set up to cover as much of the added code as possible. When contributing code to the distribution, the LAMMPS developers will appreciate if additions to the integrated unit test facility are included.

Given the complex nature of MD simulations where many operations can only be performed when suitable “real” simulation environment has been set up, not all tests will be unit tests in the strict definition of the term. They are rather executed on a more abstract level by issuing LAMMPS script commands and then inspecting the changes to the internal data. For some classes of tests, generic test programs have been written that can be applied to parts of LAMMPS that use the same interface (via polymorphism) and those are driven by input files, so tests can be added by simply adding more of those input files. Those tests should be seen more as a hybrid between unit and regression tests.

When adding tests it is recommended to also *enable support for code coverage reporting*, and study the coverage reports so that it is possible to monitor which parts of the code of a given file are executed during the tests and which tests would need to be added to increase the coverage.

The tests are grouped into categories and corresponding folders. The following sections describe how the tests are implemented and executed in those categories with increasing complexity of tests and implementation.

### 15.11.1 Tests for utility functions

These tests are driven by programs in the `unittest/utils` folder and most closely resemble conventional unit tests. There is one test program for each namespace or group of classes or file. The naming convention for the sources and executables is that they start with `test_`. The following sources and groups of tests are currently available:

File name:	Test name:	Description:
<code>test_argutils.cpp</code>	ArgInfo	Tests for ArgInfo class used by LAMMPS
<code>test_fmtlib.cpp</code>	FmtLib	Tests for <code>fmtlib::</code> functions used by LAMMPS
<code>test_math_eigen_impl.cpp</code>	MathEigen	Tests for <code>MathEigen::</code> classes and functions
<code>test_mempool.cpp</code>	MemPool	Tests for <i>MyPage</i> and <i>MyPoolChunk</i>
<code>test_tokenizer.cpp</code>	Tokenizer	Tests for <i>Tokenizer</i> and <i>ValueTokenizer</i>
<code>test_utils.cpp</code>	Utils	Tests for <code>utils::</code> <i>functions</i>

To add tests either an existing source file needs to be modified or a new source file needs to be added to the distribution and enabled for testing. To add a new file suitable CMake script code needs to be added to the `CMakeLists.txt` file in the `unittest/utils` folder. Example:

```
add_executable(test_tokenizer test_tokenizer.cpp)
target_link_libraries(test_tokenizer PRIVATE lammops GTest::GMockMain GTest::GMock_
↳GTest::GTest)
add_test(Tokenizer test_tokenizer)
```

This adds instructions to build the `test_tokenizer` executable from `test_tokenizer.cpp` and links it with the GoogleTest libraries and the LAMMPS library as well as it uses the `main()` function from the GoogleMock library of GoogleTest. The third line registers the executable as a test program to be run from `ctest` under the name `Tokenizer`.

The test executable itself will execute multiple individual tests through the GoogleTest framework. In this case each test consists of creating a tokenizer class instance with a given string and explicit or default separator choice, and then executing member functions of the class and comparing their results with expected values. A few examples:

```
TEST(Tokenizer, empty_string)
{
    Tokenizer t("", " ");
    ASSERT_EQ(t.count(), 0);
}

TEST(Tokenizer, two_words)
{
    Tokenizer t("test word", " ");
    ASSERT_EQ(t.count(), 2);
}

TEST(Tokenizer, default_separators)
{
    Tokenizer t(" \r\n test \t word \f");
    ASSERT_THAT(t.next(), Eq("test"));
    ASSERT_THAT(t.next(), Eq("word"));
    ASSERT_EQ(t.count(), 2);
}
```

Each of these TEST functions will become an individual test run by the test program. When using the `ctest` command as a front end to run the tests, their output will be suppressed and only a summary printed, but adding the `'-V'` option will then produce output from the tests above like the following:

```
[...]
1: [ RUN      ] Tokenizer.empty_string
1: [         OK ] Tokenizer.empty_string (0 ms)
1: [ RUN      ] Tokenizer.two_words
1: [         OK ] Tokenizer.two_words (0 ms)
1: [ RUN      ] Tokenizer.default_separators
1: [         OK ] Tokenizer.default_separators (0 ms)
[...]
```

The MathEigen test collection has been adapted from a standalone test and does not use the GoogleTest framework and thus not representative. The other test sources, however, can serve as guiding examples for additional tests.

### 15.11.2 Tests for individual LAMMPS commands

The tests `unittest/commands` are a bit more complex as they require to first create a `LAMMPS` class instance and then use the `C++ API` to pass individual commands to that LAMMPS instance. For that reason these tests use a GoogleTest “test fixture”, i.e. a class derived from `testing::Test` that will create (and delete) the required LAMMPS class instance for each set of tests in a `TEST_F()` function. Please see the individual source files for different examples of setting up suitable test fixtures. Here is an example for implementing a test using a fixture by first checking the default value and then issuing LAMMPS commands and checking whether they have the desired effect:

```
TEST_F(SimpleCommandsTest, ResetTimestep)
{
    ASSERT_EQ(lmp->update->ntimestep, 0);

    BEGIN_HIDE_OUTPUT();
    command("reset_timestep 10");
    END_HIDE_OUTPUT();
    ASSERT_EQ(lmp->update->ntimestep, 10);

    BEGIN_HIDE_OUTPUT();
    command("reset_timestep 0");
    END_HIDE_OUTPUT();
    ASSERT_EQ(lmp->update->ntimestep, 0);

    TEST_FAILURE(".*ERROR: Timestep must be >= 0.*", command("reset_timestep -10"));
    TEST_FAILURE(".*ERROR: Illegal reset_timestep .*", command("reset_timestep"));
    TEST_FAILURE(".*ERROR: Illegal reset_timestep .*", command("reset_timestep 10 10"));
    TEST_FAILURE(".*ERROR: Expected integer .*", command("reset_timestep xxx"));
}
```

Please note the use of the `BEGIN_HIDE_OUTPUT` and `END_HIDE_OUTPUT` functions that will capture output from running LAMMPS. This is normally discarded but by setting the verbose flag (via setting the `TEST_ARGS` environment variable, `TEST_ARGS=-v`) it can be printed and used to understand why tests fail unexpectedly.

Another complexity of these tests stems from the need to capture situations where LAMMPS will stop with an error, i.e. handle so-called “death tests”. Here the LAMMPS code will operate differently depending on whether it was configured to throw C++ exceptions on errors or call either `exit()` or `MPI_Abort()`. In the latter case, the test code also needs to detect whether LAMMPS was compiled with the OpenMPI library, as OpenMPI is **only** compatible the death test options of the GoogleTest library when C++ exceptions are enabled; otherwise those “death tests” must be

skipped to avoid reporting bogus failures. The specifics of this step are implemented in the `TEST_FAILURE()` macro. These tests operate by capturing the screen output when executing the failing command and then comparing that with a provided regular expression string pattern. Example:

```
TEST_F(SimpleCommandsTest, UnknownCommand)
{
    TEST_FAILURE(".*ERROR: Unknown command.*", lmp->input->one("XXX one two"));
}
```

The following test programs are currently available:

File name:	Test name:	Description:
test_simple_commands.cpp	SimpleCommands	Tests for LAMMPS commands that do not require a box
test_lattice_region.cpp	LatticeRegion	Tests to validate the <i>lattice</i> and <i>region</i> commands
test_groups.cpp	GroupTest	Tests to validate the <i>group</i> command
test_variables.cpp	VariableTest	Tests to validate the <i>variable</i> command
test_kim_commands.cpp	KimCommands	Tests for several commands from the <i>KIM package</i>
test_reset_ids.cpp	ResetIDs	Tests to validate the <i>reset_atom_ids</i> and <i>reset_mol_ids</i> commands

### 15.11.3 Tests for the C-style library interface

Tests for validating the LAMMPS C-style library interface are in the `unittest/c-library` folder. They are implemented in either way used for utility functions and for LAMMPS commands, but use the functions implemented in the `src/library.cpp` file as much as possible. There may be some overlap with other tests, but only in as much as is required to test the C-style library API. The tests are distributed over multiple test programs which tries to match the grouping of the functions in the source code and *in the manual*.

This group of tests also includes tests invoking LAMMPS in parallel through the library interface, provided that LAMMPS was compiled with MPI support. These include tests where LAMMPS is run in multi-partition mode or only on a subset of the MPI world communicator. The CMake script code for adding this kind of test looks like this:

```
if (BUILD_MPI)
    add_executable(test_library_mpi test_library_mpi.cpp)
    target_link_libraries(test_library_mpi PRIVATE lammps GTest::GTest GTest::GMock)
    target_compile_definitions(test_library_mpi PRIVATE ${TEST_CONFIG_DEFS})
    add_mpi_test(NAME LibraryMPI NUM_PROCS 4 COMMAND $<TARGET_FILE:test_library_mpi>)
endif()
```

Note the custom function `add_mpi_test()` which adapts how `ctest` will execute the test so it is launched in parallel (with 4 MPI ranks).

### 15.11.4 Tests for the Python module and package

The `unittest/python` folder contains primarily tests for classes and functions in the LAMMPS python module but also for commands in the PYTHON package. These tests are only enabled, if the necessary prerequisites are detected or enabled during configuration and compilation of LAMMPS (shared library build enabled, Python interpreter found, Python development files found).

The Python tests are implemented using the `unittest` standard Python module and split into multiple files with similar categories as the tests for the C-style library interface.

### 15.11.5 Tests for the Fortran interface

Tests for using the Fortran module are in the `unittest/fortran` folder. Since they are also using the GoogleTest library, they require to also implement test wrappers in C++ that will call fortran functions which provide a C function interface through `ISO_C_BINDINGS` that will in turn call the functions in the LAMMPS Fortran module. This part of the unit tests is incomplete since the Fortran module it is based on is incomplete as well.

### 15.11.6 Tests for the C++-style library interface

The tests in the `unittest/cplusplus` folder are somewhat similar to the tests for the C-style library interface, but do not need to test the several convenience and utility functions that are only available through the C-style interface. Instead it can focus on the more generic features that are used internally. This part of the unit tests is currently still mostly in the planning stage.

### 15.11.7 Tests for reading and writing file formats

The `unittest/formats` folder contains test programs for reading and writing files like data files, restart files, potential files or dump files. This covers simple things like the file i/o convenience functions in the `utils::` namespace to complex tests of atom styles where creating and deleting of atoms with different properties is tested in different ways and through script commands or reading and writing of data or restart files.

### 15.11.8 Tests for styles computing or modifying forces

These are tests common configurations for pair styles, bond styles, angle styles, kspace styles and certain fix styles. Those are tests driven by some test executables build from sources in the `unittest/force-styles` folder and use LAMMPS input template and data files as well as input files in YAML format from the `unittest/force-styles/tests` folder. The YAML file names have to follow some naming conventions so they get associated with the test programs and categorized and listed with canonical names in the list of tests as displayed by `ctest -N`. If you add a new YAML file, you need to re-run CMake to update the corresponding list of tests.

A minimal YAML file for a (molecular) pair style test will look something like the following (see `mol-pair-zero.yaml`):

```
---
lammps_version: 24 Aug 2020
date_generated: Tue Sep 15 09:44:21 202
epsilon: 1e-14
prerequisites: ! |
  atom full
  pair zero
pre_commands: ! ""
```

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```

post_commands: ! ""
input_file: in.fourmol
pair_style: zero 8.0
pair_coeff: ! |
  * *
extract: ! ""
natoms: 29
init_vdwl: 0
init_coul: 0

[...]

```

The following table describes the available keys and their purpose for testing pair styles:

Key:	Description:
lammmps_version	LAMMPS version used to last update the reference data
date_generated	date when the file was last updated
epsilon	base value for the relative precision required for tests to pass
prerequisites	list of style kind / style name pairs required to run the test
pre_commands	LAMMPS commands to be executed before the input template file is read
post_commands	LAMMPS commands to be executed right before the actual tests
input_file	LAMMPS input file template based on pair style zero
pair_style	arguments to the pair_style command to be tested
pair_coeff	list of pair_coeff arguments to set parameters for the input template
extract	list of keywords supported by Pair::extract() and their dimension
natoms	number of atoms in the input file template
init_vdwl	non-Coulomb pair energy after “run 0”
init_coul	Coulomb pair energy after “run 0”
init_stress	stress tensor after “run 0”
init_forces	forces on atoms after “run 0”
run_vdwl	non-Coulomb pair energy after “run 4”
run_coul	Coulomb pair energy after “run 4”
run_stress	stress tensor after “run 4”
run_forces	forces on atoms after “run 4”

The test program will read all this data from the YAML file and then create a LAMMPS instance, apply the settings/commands from the YAML file as needed and then issue a “run 0” command, write out a restart file, a data file and a coeff file. The actual test will then compare computed energies, stresses, and forces with the reference data, issue a “run 4” command and compare to the second set of reference data. This will be run with both the newton\_pair setting enabled and disabled and is expected to generate the same results (allowing for some numerical noise). Then it will restart from the previously generated restart and compare with the reference and also start from the data file. A final check will use multi-cutoff r-RESPA (if supported by the pair style) at a 1:1 split and compare to the Verlet results. These sets of tests are run with multiple test fixtures for accelerated styles (OPT, OPENMP, INTEL) and for the latter two with 4 OpenMP threads enabled. For these tests the relative error (epsilon) is lowered by a common factor due to the additional numerical noise, but the tests are still comparing to the same reference data.

Additional tests will check whether all listed extract keywords are supported and have the correct dimensionality and the final set of tests will set up a few pairs of atoms explicitly and in such a fashion that the forces on the atoms computed from Pair::compute() will match individually with the results from Pair::single(), if the pair style does support that functionality.

With this scheme a large fraction of the code of any tested pair style will be executed and consistent results are required for different settings and between different accelerated pair style variants and the base class, as well as for computing

individual pairs through the `Pair::single()` where supported.

The `test_pair_style` tester is used with 4 categories of test inputs:

- pair styles compatible with molecular systems using bonded interactions and exclusions. For pair styles requiring a KSpace style the KSpace computations are disabled. The YAML files match the pattern “mol-pair-\*.yaml” and the tests are correspondingly labeled with “MolPairStyle:\*”
- pair styles not compatible with the previous input template. The YAML files match the pattern “atomic-pair-\*.yaml” and the tests are correspondingly labeled with “AtomicPairStyle:\*”
- manybody pair styles. The YAML files match the pattern “atomic-pair-\*.yaml” and the tests are correspondingly labeled with “AtomicPairStyle:\*”
- kspace styles. The YAML files match the pattern “kspace-\*.yaml” and the tests are correspondingly labeled with “KSpaceStyle:\*”. In these cases a compatible pair style is defined, but the computation of the pair style contributions is disabled.

The `test_bond_style` and `test_angle_style` are set up in a similar fashion and share support functions with the pair style tester. The final group of tests in this section is for fix styles that add/manipulate forces and velocities, e.g. for time integration, thermostats and more.

Adding a new test is easiest done by copying and modifying an existing test for a style that is similar to one to be tested. The file name should follow the naming conventions described above and after copying the file, the first step is to replace the style names where needed. The coefficient values do not have to be meaningful, just in a reasonable range for the given system. It does not matter if some forces are large, for as long as they do not diverge.

The template input files define a large number of index variables at the top that can be modified inside the YAML file to control the behavior. For example, if a pair style requires a “newton on” setting, the following can be used in as the “pre\_commands” section:

```
pre_commands: ! |
  variable newton_pair delete
  variable newton_pair index on
```

And for a pair style requiring a kspace solver the following would be used as the “post\_commands” section:

```
post_commands: ! |
  pair_modify table 0
  kspace_style pppm/tip4p 1.0e-6
  kspace_modify gewald 0.3
  kspace_modify compute no
```

Note that this disables computing the kspace contribution, but still will run the setup. The “gewald” parameter should be set explicitly to speed up the run. For styles with long-range electrostatics, typically two tests are added one using the (slower) analytic approximation of the `erfc()` function and the other using the tabulated coulomb, to test both code paths. The reference results in the YAML files then should be compared manually, if they agree well enough within the limits of those two approximations.

The `test_pair_style` and equivalent programs have special command line options to update the YAML files. Running a command like

```
$ test_pair_style mol-pair-lennard_mdf.yaml -g new.yaml
```

will read the settings from the `mol-pair-lennard_mdf.yaml` file and then compute the reference data and write a new file with to `new.yaml`. If this step fails, there are likely some (LAMMPS or YAML) syntax issues in the YAML file that need to be resolved and then one can compare the two files to see if the output is as expected.

It is also possible to do an update in place with:

```
$ test_pair_style mol-pair-lennard_mdf.yaml -u
```

And one can finally run the full set of tests with:

```
$ test_pair_style mol-pair-lennard_mdf.yaml
```

This will just print a summary of the groups of tests. When using the “-v” flag the test will also keep any LAMMPS output and when using the “-s” flag, there will be some statistics reported on the relative errors for the individual checks which can help to figure out what would be a good choice of the epsilon parameter. It should be as small as possible to catch any unintended side effects from changes elsewhere, but large enough to accommodate the numerical noise due to the implementation of the potentials and differences in compilers.

---

**Note:** These kinds of tests can be very sensitive to compiler optimization and thus the expectation is that they pass with compiler optimization turned off. When compiler optimization is enabled, there may be some failures, but one has to carefully check whether those are acceptable due to the enhanced numerical noise from reordering floating-point math operations or due to the compiler mis-compiling the code. That is not always obvious.

---

### 15.11.9 Tests for programs in the tools folder

The `unittest/tools` folder contains tests for programs in the `tools` folder. This currently only contains tests for the LAMMPS shell, which are implemented as a python scripts using the `unittest` Python module and launching the tool commands through the `subprocess` Python module.

## 15.12 C++ base classes

LAMMPS is designed to be used as a C++ class library where one can set up and drive a simulation through creating a class instance and then calling some abstract operations or commands on that class or its member class instances. These are interfaced to the *C library API*, which providing an additional level of abstraction simplification for common operations. The C API is also the basis for calling LAMMPS from Python or Fortran.

When used from a C++ program, most of the symbols and functions in LAMMPS are wrapped into the `LAMMPS_NS` namespace so they will not collide with your own classes or other libraries. This, however, does not extend to the additional libraries bundled with LAMMPS in the `lib` folder and some of the low-level code of some packages.

Behind the scenes this is implemented through inheritance and polymorphism where base classes define the abstract interface and derived classes provide the specialized implementation for specific models or optimizations or ports to accelerator platforms. This document will provide an outline of the fundamental class hierarchy and some selected examples for derived classes of specific models.

---

**Note:** Please see the *note about thread-safety* in the library Howto doc page.

---

### 15.12.1 LAMMPS Class

The LAMMPS class is encapsulating an MD simulation state and thus it is the class that needs to be created when starting a new simulation system state. The LAMMPS executable essentially creates one instance of this class and passes the command line flags and tells it to process the provided input (a file or `stdin`). It shuts the class down when control is returned to it and then exits. When using LAMMPS as a library from another code it is required to create an instance of this class, either directly from C++ with `new LAMMPS()` or through one of the library interface functions like `lammps_open()` of the C-library interface, or the `lammps.lammps` class constructor of the Python module, or the `lammps()` constructor of the Fortran module.

In order to avoid clashes of function names, all of the core code in LAMMPS is placed into the `LAMMPS_NS` namespace. Functions or variables outside of that namespace must be “static”, i.e. visible only to the scope of the file/object they are defined in. Code in packages or the libraries in the `lib` folder may not adhere to this as some of them are adapted from legacy code or consist of external libraries with their own requirements and policies.

---

class **LAMMPS**

*LAMMPS* simulation instance.

The *LAMMPS* class contains pointers of all constituent class instances and global variables that are used by a *LAMMPS* simulation. Its contents represent the entire state of the simulation.

The *LAMMPS* class manages the components of an MD simulation by creating, deleting, and initializing instances of the classes it is composed of, processing command line flags, and providing access to some global properties. The specifics of setting up and running a simulation are handled by the individual component class instances.

#### Public Functions

const char \***match\_style**(const char \*style, const char \*name)

Return name of package that a specific style belongs to.

This function checks the given name against all list of styles for all type of styles and if the name and the style match, it returns which package this style belongs to.

##### Parameters

- **style** – Type of style (e.g. atom, pair, fix, etc.)
- **name** – Name of style

##### Returns

Name of the package this style is part of

**LAMMPS**(int, char\*\*, MPI\_Comm)

Create a *LAMMPS* simulation instance

The *LAMMPS* constructor starts up a simulation by allocating all fundamental classes in the necessary order, parses input switches and their arguments, initializes communicators, screen and logfile output FILE pointers.

##### Parameters

- **narg** – number of arguments
- **arg** – list of arguments
- **communicator** – MPI communicator used by this *LAMMPS* instance

**~LAMMPS()**

Shut down a *LAMMPS* simulation instance

The *LAMMPS* destructor shuts down the simulation by deleting top-level class instances, closing screen and log files for the global instance (aka “world”) and files and MPI communicators in sub-partitions (“universes”). Then it deletes the fundamental class instances and copies of data inside the class.

**Public Static Functions**

static bool **is\_installed\_pkg**(const char \*pkg)

Return true if a *LAMMPS* package is enabled in this binary

**Parameters**

**pkg** – name of package

**Returns**

true if yes, else false

class **Pointers**

Base class for *LAMMPS* features.

The *Pointers* class contains references to many of the pointers and members of the *LAMMPS\_NS::LAMMPS* class. Derived classes thus gain access to the constituent class instances in the *LAMMPS* composite class and thus to the core functionality of *LAMMPS*.

This kind of construct is needed, since the *LAMMPS* constructor should only be run once per *LAMMPS* instance and thus classes cannot be derived from *LAMMPS* itself. The *Pointers* class constructor instead only initializes C++ references to component pointer in the *LAMMPS* class.

Subclassed by *Atom*, *Input*, *PotentialFileReader*

**15.12.2 LAMMPS Atom and AtomVec Base Classes**

class **Atom** : protected *Pointers*

Class to provide access to atom data.

The *Atom* class provides access to atom style related global settings and per-atom data that is stored with atoms and migrates with them from sub-domain to sub-domain as atoms move around. This includes topology data, which is stored with either one specific atom or all atoms involved depending on the settings of the *newton command*.

The actual per-atom data is allocated and managed by one of the various classes derived from the *AtomVec* class as determined by the *atom\_style command*. The pointers in the *Atom* class are updated by the *AtomVec* class as needed.

## Public Functions

**Atom**(class *LAMMPS*\*)

*Atom* class constructor

This resets and initialized all kinds of settings, parameters, and pointer variables for per-atom arrays. This also initializes the factory for creating instances of classes derived from the AtomVec base class, which correspond to the selected atom style.

### Parameters

**\_lmp** – pointer to the base *LAMMPS* class

virtual int **add\_custom**(const char\*, int, int)

Add a custom per-atom property with the given name and type and size.

This function will add a custom per-atom property with one or more values with the name “name” to the list of custom properties. This function is called, e.g. from *fix property/atom*.

### Parameters

- **name** – Name of the property (w/o a “i\_” or “d\_” or “i2\_” or “d2\_” prefix)
- **flag** – Data type of property: 0 for int, 1 for double
- **cols** – Number of values: 0 for a single value, 1 or more for a vector of values

### Returns

index of property in the respective list of properties

virtual void **remove\_custom**(int, int, int)

Remove a custom per-atom property of a given type and size.

This will remove a property that was requested, e.g. by the *fix property/atom* command. It frees the allocated memory and sets the pointer to `nullptr` for the entry in the list so it can be reused. The lists of these pointers are never compacted or shrink, so that indices to name mappings remain valid.

### Parameters

- **index** – Index of property in the respective list of properties
- **flag** – Data type of property: 0 for int, 1 for double
- **cols** – Number of values: 0 for a single value, 1 or more for a vector of values

void **\*extract**(const char\*)

Provide access to internal data of the *Atom* class by keyword

This function is a way to access internal per-atom data. This data is distributed across MPI ranks and thus only the data for “local” atoms are expected to be available. Whether also data for “ghost” atoms is stored and up-to-date depends on various simulation settings.

This table lists a large part of the supported names, their data types, length of the data area, and a short description.

Name	Type	Items per atom	Description
mass	double	1	per-type mass. This array is <b>NOT</b> a per-atom array but of length <code>nTYPES+1</code> , element 0 is ignored.
id	tagint	1	atom ID of the particles
type	int	1	atom type of the particles
mask	int	1	bitmask for mapping to groups. Individual bits are set to 0 or 1 for each group.
image	imageint	1	3 image flags encoded into a single integer. See <a href="#">lammers_encode_image_flags()</a> .
x	double	3	x-, y-, and z-coordinate of the particles
v	double	3	x-, y-, and z-component of the velocity of the particles
f	double	3	x-, y-, and z-component of the force on the particles
molecule	int	1	molecule ID of the particles
q	double	1	charge of the particles
mu	double	3	dipole moment of the particles
omega	double	3	x-, y-, and z-component of rotational velocity of the particles
angmom	double	3	x-, y-, and z-component of angular momentum of the particles
torque	double	3	x-, y-, and z-component of the torque on the particles
radius	double	1	radius of the (extended) particles
rmass	double	1	per-atom mass of the particles. <code>nullptr</code> if per-type masses are used. See the <a href="#">rmass_flag</a> setting.
ellipsoid	int	1	1 if the particle is an ellipsoidal particle, 0 if not
line	int	1	1 if the particle is a line particle, 0 if not
tri	int	1	1 if the particle is a triangulated particle, 0 if not
body	int	1	1 if the particle is a body particle, 0 if not
quat	double	4	four quaternion components of the particles
i_name	int	1	single integer value defined by fix property/atom vector name
d_name	double	1	single double value defined by fix property/atom vector name
i2_name	int	n	N integer values defined by fix property/atom array name
d2_name	double	n	N double values defined by fix property/atom array name

*See also*

[lammers\\_extract\\_atom\(\)](#)

**See also:**

[extract\\_datatype](#)

**Parameters**

**name** – string with the keyword of the desired property. Typically the name of the pointer variable returned

**Returns**

pointer to the requested data cast to `void *` or `nullptr`

int **extract\_datatype**(const char\*)

Provide data type info about internal data of the *Atom* class

New in version 18Sep2020.

**See also:**

*extract*

**Parameters**

**name** – string with the keyword of the desired property.

**Returns**

data type constant for desired property or -1

struct **PerAtom**

### 15.12.3 LAMMPS Input Base Class

class **Input** : protected *Pointers*

Class for processing commands and input files.

The Input class contains methods for reading, pre-processing and parsing LAMMPS commands and input files and will dispatch commands to the respective class instances or contains the code to execute the commands directly. It also contains the instance of the Variable class which performs computations and text substitutions.

**Public Functions**

**Input**(class *LAMMPS\**, int, char\*\*)

*Input* class constructor

This sets up the input processing, processes the *-var* and *-echo* command line flags, holds the factory of commands and creates and initializes an instance of the Variable class.

To execute a command, a specific class instance, derived from *Command*, is created, then its *command()* member function executed, and finally the class instance is deleted.

**Parameters**

- **lmp** – pointer to the base *LAMMPS* class
- **argc** – number of entries in *argv*



- **argv** – argument vector

void **file()**

Process all input from the FILE \* pointer *infile*

This will read lines from *infile*, parse and execute them until the end of the file is reached. The *infile* pointer will usually point to `stdin` or the input file given with the `-in` command line flag.

void **file(const char\*)**

Process all input from the file *filename*

This function opens the file at the path *filename*, put the current file pointer stored in *infile* on a stack and instead assign *infile* with the newly opened file pointer. Then it will call the `Input::file()` function to read, parse and execute the contents of that file. When the end of the file is reached, it is closed and the previous file pointer from the *infile* file pointer stack restored to *infile*.

#### Parameters

**filename** – name of file with *LAMMPS* commands

char \***one(const std::string&)**

Process a single command from a string in *single*

This function takes the text in *single*, makes a copy, parses that, executes the command and returns the name of the command (without the arguments). If there was no command in *single* it will return `nullptr`.

#### Parameters

**single** – string with *LAMMPS* command

#### Returns

string with name of the parsed command w/o arguments

## 15.13 Platform abstraction functions

The `platform` sub-namespace inside the `LAMMPS_NS` namespace provides a collection of wrapper and convenience functions and utilities that perform common tasks for which platform specific code would be required or for which a more high-level abstraction would be convenient and reduce duplicated code. This reduces redundant implementations and encourages consistent behavior and thus has some overlap with the “*utils*” sub-namespace.

### 15.13.1 Time functions

double `LAMMPS_NS::platform::cputime()`

Return the consumed CPU time for the current process in seconds

This is a wrapper around the POSIX function `getrusage()` and its Windows equivalent. It is to be used in a similar fashion than `MPI_Wtime()`. Its resolution may be rather low so it can only be trusted when observing processes consuming at seconds or more of CPU time.

#### Returns

used CPU time in seconds

double `LAMMPS_NS::platform::walltime()`

Return the wall clock state for the current process in seconds

This clock is counting continuous time is initialized during Load of the executable/library. Its absolute value must be considered arbitrary and thus elapsed wall times are measured in taking differences. It is therefore to be used in a similar fashion as `MPI_Wtime()` but has a different offset, usually leading to better resolution.

**Returns**

wall clock time in seconds

`void LAMMPS_NS::platform::usleep(int usec)`

Suspend execution for a microsecond interval

This emulates the `usleep(3)` BSD function call also mentioned in POSIX.1-2001. This is not a precise delay; it may be longer, but not shorter.

**Parameters**

`usec` – length of delay in microseconds

### 15.13.2 Platform information functions

`std::string LAMMPS_NS::platform::os_info()`

Return string with the operating system version and architecture info

**Returns**

string with info about the OS and the platform is is running on

`std::string LAMMPS_NS::platform::compiler_info()`

Return string with compiler version info

This function uses predefined compiler macros to identify Compilers and their version and configuration info.

**Returns**

string with the compiler information text

`std::string LAMMPS_NS::platform::cxx_standard()`

Return string with C++ standard version used to compile *LAMMPS*.

This function uses predefined compiler macros to identify the C++ standard version used to compile *LAMMPS* with.

**Returns**

string with the C++ standard version or “unknown”

`std::string LAMMPS_NS::platform::openmp_standard()`

Return string with OpenMP standard version info

This function uses predefined compiler macros to identify OpenMP support and the supported version of the standard.

**Returns**

string with the openmp information text

`std::string LAMMPS_NS::platform::mpi_vendor()`

Return string with MPI vendor info

This function uses predefined macros to identify the vendor of the MPI library used.

**Returns**

string with the MPI vendor information text

```
std::string LAMMPS_NS::platform::mpi_info(int &major, int &minor)
```

Return string with MPI version info

This function uses predefined macros and MPI function calls to identify the version of the MPI library used.

#### Parameters

- **major** – major version of the MPI standard (set on exit)
- **minor** – minor version of the MPI standard (set on exit)

#### Returns

string with the MPI version information text

```
std::string LAMMPS_NS::platform::compress_info()
```

Return string with list of available compression types and executables

This function tests which of the supported compression executables are available for reading or writing compressed files where supported.

#### Returns

string with list of available compression tools

### 15.13.3 File and path functions and global constants

```
constexpr char LAMMPS_NS::platform::filepathsep[] = "/"
```

Platform specific file path component separator

This is a string with the character that separates directories and filename in paths on a platform. If multiple are characters are provided, the first is the preferred one.

```
constexpr char LAMMPS_NS::platform::pathvarsep = ':'
```

Platform specific path environment variable component separator

This is the character that separates entries in “PATH”-style environment variables.

```
const char *LAMMPS_NS::platform::guesspath(FILE *fp, char *buf, int len)
```

Try to detect pathname from FILE pointer

Currently only supported on Linux, MacOS, and Windows. Otherwise will report “(unknown)”.

On Linux the folder `/proc/self/fd` holds symbolic links to the actual pathnames associated with each open file descriptor of the current process. On MacOS the same kind of information can be obtained using `fcntl(fd, F_GETPATH, buf)`. On Windows we use `GetFinalPathNameByHandleA()` which is available with Windows Vista and later. If the buffer is too small (< 16 bytes) a null pointer is returned.

This function is used to provide a filename with error messages in functions where the filename is not passed as an argument, but the FILE \* pointer.

#### Parameters

- **fp** – FILE pointer struct from STDIO library for which we want to detect the name
- **buf** – storage buffer for pathname. output will be truncated if not large enough
- **len** – size of storage buffer. output will be truncated to this length - 1

**Returns**

pointer to the storage buffer with path or a NULL pointer if buf is invalid or the buffer size is too small

std::string LAMMPS\_NS::platform::path\_basename(const std::string &path)

Strip off leading part of path, return just the filename

**Parameters**

**path** – file path

**Returns**

file name

std::string LAMMPS\_NS::platform::path\_join(const std::string &a, const std::string &b)

Join two pathname segments

This uses the forward slash ‘/’ character unless *LAMMPS* is compiled for Windows where it uses the backward slash ‘\’

**Parameters**

- **a** – first path
- **b** – second path

**Returns**

combined path

bool LAMMPS\_NS::platform::file\_is\_readable(const std::string &path)

Check if file exists and is readable

**Parameters**

**path** – file path

**Returns**

true if file exists and is readable

bool LAMMPS\_NS::platform::is\_console(FILE \*fp)

Check if a file pointer may be connected to a console

**Parameters**

**fp** – file pointer

**Returns**

true if the file pointer is flagged as a TTY

bool LAMMPS\_NS::platform::path\_is\_directory(const std::string &path)

Check if a path is a directory

**Parameters**

**path** – directory path

**Returns**

true if the directory exists

std::string LAMMPS\_NS::platform::current\_directory()

Get string with path to the current directory

**Returns**

path to the current directory or empty string

`std::vector<std::string> LAMMPS_NS::platform::list_directory(const std::string &dir)`

Get list of entries in a directory

This provides a list of strings of the entries in the directory without the leading path name while also skipping over “..” and “.”.

**Parameters**

**dir** – path to directory

**Returns**

vector with strings of all directory entries

`int LAMMPS_NS::platform::chdir(const std::string &path)`

Change current directory

**Parameters**

**path** – new current working directory path

**Returns**

-1 if unsuccessful, otherwise  $\geq 0$

`int LAMMPS_NS::platform::mkdir(const std::string &path)`

Create a directory

Unlike the the `mkdir()` or `_mkdir()` functions of the C library, this function will also try to create non-existing sub-directories in case they don't exist, and thus behave like the `mkdir -p` command rather than plain `mkdir` or ``md`.

**Parameters**

**path** – directory path

**Returns**

-1 if unsuccessful, otherwise  $\geq 0$

`int LAMMPS_NS::platform::rmdir(const std::string &path)`

Delete a directory

Unlike the the `rmdir()` or `_rmdir()` functions of the C library, this function will check for the contents of the folder and recurse into any sub-folders, if necessary and delete all contained folders and their contents before deleting the folder *path*.

**Parameters**

**path** – directory path

**Returns**

-1 if unsuccessful, otherwise  $\geq 0$

`int LAMMPS_NS::platform::unlink(const std::string &path)`

Delete a file

**Parameters**

**path** – path to file to be deleted

**Returns**

0 on success, -1 on error

### 15.13.4 Standard I/O function wrappers

constexpr bigint LAMMPS\_NS::platform::END\_OF\_FILE = -1  
constant to seek to the end of the file

bigint LAMMPS\_NS::platform::ftell(FILE \*fp)

Get current file position

**Parameters**

**fp** – FILE pointer of the given file

**Returns**

current FILE pointer position cast to a bigint

int LAMMPS\_NS::platform::fseek(FILE \*fp, bigint pos)

Set absolute file position

If the absolute position is END\_OF\_FILE, then position at the end of the file.

**Parameters**

- **fp** – FILE pointer of the given file
- **pos** – new position of the FILE pointer

**Returns**

0 if successful, otherwise -1

int LAMMPS\_NS::platform::ftruncate(FILE \*fp, bigint length)

Truncate file to a given length and re-position file pointer

**Parameters**

- **fp** – FILE pointer of the given file
- **length** – length to which the file is being truncated to

**Returns**

0 if successful, otherwise -1

FILE \*LAMMPS\_NS::platform::popen(const std::string &cmd, const std::string &mode)

Open a pipe to a command for reading or writing

**Parameters**

- **cmd** – command for the pipe
- **mode** – “r” for reading from *cmd* or “w” for writing to *cmd*

**Returns**

file pointer to the pipe if successful or null

int LAMMPS\_NS::platform::pclose(FILE \*fp)

Close a previously opened pipe

**Parameters**

**fp** – FILE pointer for the pipe

**Returns**

exit status of the pipe command or -1 in case of errors

### 15.13.5 Environment variable functions

int LAMMPS\_NS::platform::putenv(const std::string &vardef)

Add variable to the environment

**Parameters**

**vardef** – variable name or variable definition (NAME=value)

**Returns**

-1 if failure otherwise 0

int LAMMPS\_NS::platform::unsetenv(const std::string &variable)

Delete variable from the environment

**Parameters**

**variable** – variable name

**Returns**

-1 if failure otherwise 0

std::vector<std::string> LAMMPS\_NS::platform::list\_pathenv(const std::string &var)

Get list of entries in a path environment variable

This provides a list of strings of the entries in an environment variable that is containing a “path” like “PATH” or “LD\_LIBRARY\_PATH”.

**Parameters**

**var** – name of the environment variable

**Returns**

vector with strings of all entries in that path variable

std::string LAMMPS\_NS::platform::find\_exe\_path(const std::string &cmd)

Find pathname of an executable in the standard search path

This function will traverse the list of directories in the PATH environment variable and look for the executable *cmd*. If the file exists and is executable the full path is returned as string, otherwise an empty string is returned.

On Windows the *cmd* string must not include an extension as this function will automatically append the extensions “.exe”, “.com” and “.bat” and look for those paths. On Windows also the current directory is checked (and first), while otherwise not unless “.” exists in the PATH environment variable.

Because of the nature of the check, this will not detect shell functions built-in command or aliases.

**Parameters**

**cmd** – name of command

**Returns**

vector with strings of all directory entries

### 15.13.6 Dynamically loaded object or library functions

void \*LAMMPS\_NS::platform::**dlopen**(const std::string &fname)

Open a shared object file or library

**Parameters**

**fname** – name or path of the shared object

**Returns**

handle to the shared object or null

int LAMMPS\_NS::platform::**dlclose**(void \*handle)

Close a shared object

This releases the object corresponding to the provided handle. Resolved symbols associated with this handle may not be used after this call

**Parameters**

**handle** – handle to an opened shared object

**Returns**

0 if successful, non-zero of not

void \*LAMMPS\_NS::platform::**dlsym**(void \*handle, const std::string &symbol)

Resolve a symbol in shared object

**Parameters**

- **handle** – handle to an opened shared object
- **symbol** – name of the symbol to extract

**Returns**

pointer to the resolved symbol or null

std::string LAMMPS\_NS::platform::**dlerror**()

Obtain error diagnostic info after dynamic linking function calls

Return a human-readable string describing the most recent error that occurred when using one of the functions for dynamic loading objects the last call to this function. The string is empty, if there was no error.

**Returns**

string with error message or empty

### 15.13.7 Compressed file I/O functions

bool LAMMPS\_NS::platform::**has\_compress\_extension**(const std::string &file)

Check if a file name ends in a known extension for a compressed file format

Currently supported file extensions are: .gz, .bz2, .zst, .xz, .lzma, lz4

**Parameters**

**file** – name of the file to check

**Returns**

true if the file has a known extension, otherwise false

FILE \*LAMMPS\_NS::platform::**compressed\_read**(const std::string &file)

Open pipe to compressed text file for reading



**Parameters****file** – name of the file to open**Returns**

FILE pointer to pipe using for reading the compressed file.

FILE \*LAMMPS\_NS::platform::compressed\_write(const std::string &amp;file)

Open pipe to compressed text file for writing

**Parameters****file** – name of the file to open**Returns**

FILE pointer to pipe using for reading the compressed file.

## 15.14 Utility functions

The `utils` sub-namespace inside the `LAMMPS_NS` namespace provides a collection of convenience functions and utilities that perform common tasks that are required repeatedly throughout the LAMMPS code like reading or writing to files with error checking or translation of strings into specific types of numbers with checking for validity. This reduces redundant implementations and encourages consistent behavior and thus has some overlap with the “*platform*” sub-namespace.

### 15.14.1 I/O with status check and similar functions

The the first two functions are wrappers around the corresponding C library calls `fgets()` or `fread()`. They will check if there were errors on reading or an unexpected end-of-file state was reached. In that case, the functions will stop with an error message, indicating the name of the problematic file, if possible unless the `error` argument is a NULL pointer.

The `utils::fgets_trunc()` function will work similar for `fgets()` but it will read in a whole line (i.e. until the end of line or end of file), but store only as many characters as will fit into the buffer including a final newline character and the terminating NULL byte. If the line in the file is longer it will thus be truncated in the buffer. This function is used by `utils::read_lines_from_file()` to read individual lines but make certain they follow the size constraints.

The `utils::read_lines_from_file()` function will read the requested number of lines of a maximum length into a buffer and will return 0 if successful or 1 if not. It also guarantees that all lines are terminated with a newline character and the entire buffer with a NULL character.

---

```
void LAMMPS_NS::utils::sfgets(const char *srcname, int srcline, char *s, int size, FILE *fp, const char
                             *filename, Error *error)
```

Safe wrapper around `fgets()` which aborts on errors or EOF and prints a suitable error message to help debugging.

Use `nullptr` as the error parameter to avoid the abort on EOF or error.

**Parameters**

- **srcname** – name of the calling source file (from `FLERR` macro)
- **srcline** – line in the calling source file (from `FLERR` macro)
- **s** – buffer for storing the result of `fgets()`
- **size** – size of buffer `s` (max number of bytes read by `fgets()`)
- **fp** – file pointer used by `fgets()`

- **filename** – file name associated with fp (may be a null pointer; then *LAMMPS* will try to detect)
- **error** – pointer to Error class instance (for abort) or nullptr

void LAMMPS\_NS::utils::sfread(const char \*srcname, int srcline, void \*s, size\_t size, size\_t num, FILE \*fp, const char \*filename, Error \*error)

Safe wrapper around fread() which aborts on errors or EOF and prints a suitable error message to help debugging.

Use nullptr as the error parameter to avoid the abort on EOF or error.

#### Parameters

- **srcname** – name of the calling source file (from FLERR macro)
- **srcline** – line in the calling source file (from FLERR macro)
- **s** – buffer for storing the result of fread()
- **size** – size of data elements read by fread()
- **num** – number of data elements read by fread()
- **fp** – file pointer used by fread()
- **filename** – file name associated with fp (may be a null pointer; then *LAMMPS* will try to detect)
- **error** – pointer to Error class instance (for abort) or nullptr

char \*LAMMPS\_NS::utils::fgets\_trunc(char \*s, int size, FILE \*fp)

Wrapper around fgets() which reads whole lines but truncates the data to the buffer size and ensures a newline char at the end.

This function is useful for reading line based text files with possible comments that should be parsed later. This applies to data files, potential files, atomfile variable files and so on. It is used instead of fgets() by utils::read\_lines\_from\_file().

#### Parameters

- **s** – buffer for storing the result of fgets()
- **size** – size of buffer s (max number of bytes returned)
- **fp** – file pointer used by fgets()

int LAMMPS\_NS::utils::read\_lines\_from\_file(FILE \*fp, int nlines, int nmax, char \*buffer, int me, MPI\_Comm comm)

Read N lines of text from file into buffer and broadcast them

This function uses repeated calls to fread() to fill a buffer with newline terminated text. If a line does not end in a newline (e.g. at the end of a file), it is added. The caller has to allocate an nlines by nmax sized buffer for storing the text data. Reading is done by MPI rank 0 of the given communicator only, and thus only MPI rank 0 needs to provide a valid file pointer.

#### Parameters

- **fp** – file pointer used by fread
- **nlines** – number of lines to be read
- **nmax** – maximum length of a single line
- **buffer** – buffer for storing the data.
- **me** – MPI rank of calling process in MPI communicator

- **comm** – MPI communicator for broadcast

**Returns**

1 if the read was short, 0 if read was successful

## 15.14.2 String to number conversions with validity check

These functions should be used to convert strings to numbers. They are strongly preferred over C library calls like `atoi()` or `atof()` since they check if the **entire** string is a valid (floating-point or integer) number, and will error out instead of silently returning the result of a partial conversion or zero in cases where the string is not a valid number. This behavior improves detecting typos or issues when processing input files.

Similarly the `utils::logical()` function will convert a string into a boolean and will only accept certain words.

The `do_abort` flag should be set to `true` in case this function is called only on a single MPI rank, as that will then trigger the a call to `Error::one()` for errors instead of `Error::all()` and avoids a “hanging” calculation when run in parallel.

Please also see `utils::is_integer()` and `utils::is_double()` for testing strings for compliance without conversion.

```
double LAMMPS_NS::utils::numeric(const char *file, int line, const std::string &str, bool do_abort, LAMMPS
                               *Imp)
```

Convert a string to a floating point number while checking if it is a valid floating point or integer number

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to number
- **do\_abort** – determines whether to call `Error::one()` or `Error::all()`
- **Imp** – pointer to top-level *LAMMPS* class instance

**Returns**

double precision floating point number

```
double LAMMPS_NS::utils::numeric(const char *file, int line, const char *str, bool do_abort, LAMMPS *Imp)
```

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to number
- **do\_abort** – determines whether to call `Error::one()` or `Error::all()`
- **Imp** – pointer to top-level *LAMMPS* class instance

**Returns**

double precision floating point number

int LAMMPS\_NS::utils::inumeric(const char \*file, int line, const std::string &str, bool do\_abort, LAMMPS \*Imp)

Convert a string to an integer number while checking if it is a valid integer number (regular int)

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to number
- **do\_abort** – determines whether to call Error::one() or Error::all()
- **Imp** – pointer to top-level LAMMPS class instance

**Returns**

integer number (regular int)

int LAMMPS\_NS::utils::inumeric(const char \*file, int line, const char \*str, bool do\_abort, LAMMPS \*Imp)

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to number
- **do\_abort** – determines whether to call Error::one() or Error::all()
- **Imp** – pointer to top-level LAMMPS class instance

**Returns**

double precision floating point number

bigint LAMMPS\_NS::utils::bnumeric(const char \*file, int line, const std::string &str, bool do\_abort, LAMMPS \*Imp)

Convert a string to an integer number while checking if it is a valid integer number (bigint)

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to number
- **do\_abort** – determines whether to call Error::one() or Error::all()
- **Imp** – pointer to top-level LAMMPS class instance

**Returns**

integer number (bigint)

bigint LAMMPS\_NS::utils::bnumeric(const char \*file, int line, const char \*str, bool do\_abort, LAMMPS \*Imp)

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to number

- **do\_abort** – determines whether to call `Error::one()` or `Error::all()`
- **lmp** – pointer to top-level *LAMMPS* class instance

**Returns**

double precision floating point number

```
tagint LAMMPS_NS::utils::tnumeric(const char *file, int line, const std::string &str, bool do_abort, LAMMPS
                               *lmp)
```

Convert a string to an integer number while checking if it is a valid integer number (tagint)

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to number
- **do\_abort** – determines whether to call `Error::one()` or `Error::all()`
- **lmp** – pointer to top-level *LAMMPS* class instance

**Returns**

integer number (tagint)

```
tagint LAMMPS_NS::utils::tnumeric(const char *file, int line, const char *str, bool do_abort, LAMMPS *lmp)
```

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to number
- **do\_abort** – determines whether to call `Error::one()` or `Error::all()`
- **lmp** – pointer to top-level *LAMMPS* class instance

**Returns**

double precision floating point number

```
int LAMMPS_NS::utils::logical(const char *file, int line, const std::string &str, bool do_abort, LAMMPS *lmp)
```

Convert a string to a boolean while checking whether it is a valid boolean term. Valid terms are ‘yes’, ‘no’, ‘true’, ‘false’, ‘on’, ‘off’, and ‘1’, ‘0’. Only lower case is accepted.

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to logical
- **do\_abort** – determines whether to call `Error::one()` or `Error::all()`
- **lmp** – pointer to top-level *LAMMPS* class instance

**Returns**

1 if string resolves to “true”, otherwise 0

int LAMMPS\_NS::utils::logical(const char \*file, int line, const char \*str, bool do\_abort, LAMMPS \*Imp)

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

**Parameters**

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be converted to logical
- **do\_abort** – determines whether to call Error::one() or Error::all()
- **Imp** – pointer to top-level LAMMPS class instance

**Returns**

1 if string resolves to “true”, otherwise 0

### 15.14.3 String processing

The following are functions to help with processing strings and parsing files or arguments.

---

char \*LAMMPS\_NS::utils::strdup(const std::string &text)

Make C-style copy of string in new storage

This allocates a storage buffer and copies the C-style or C++ style string into it. The buffer is allocated with “new” and thus needs to be deallocated with “delete[]”.

**Parameters**

**text** – string that should be copied

**Returns**

new buffer with copy of string

std::string LAMMPS\_NS::utils::lowercase(const std::string &line)

Convert string to lowercase

**Parameters**

**line** – string that should be converted

**Returns**

new string with all lowercase characters

std::string LAMMPS\_NS::utils::uppercase(const std::string &line)

Convert string to uppercase

**Parameters**

**line** – string that should be converted

**Returns**

new string with all uppercase characters

std::string LAMMPS\_NS::utils::trim(const std::string &line)

Trim leading and trailing whitespace. Like TRIM() in Fortran.

**Parameters**

**line** – string that should be trimmed

**Returns**

new string without whitespace (string)

`std::string LAMMPS_NS::utils::trim_comment` (const std::string &line)

Return string with anything from the first '#' character onward removed

**Parameters**

**line** – string that should be trimmed

**Returns**

new string without comment (string)

`std::string LAMMPS_NS::utils::star_subst` (const std::string &name, bigint step, int pad)

Replace first '\*' character in a string with a number, optionally zero-padded

If there is no '\*' character in the string, return the original string. If the number requires more characters than the value of the *pad* argument, do not add zeros; otherwise add as many zeroes as needed to the left to make the number representation *pad* characters wide.

**Parameters**

- **name** – string with file containing a '\*' (or not)
- **step** – step number to replace the (first) '\*'
- **pad** – zero-padding (may be zero)

**Returns**

processed string

`inline bool LAMMPS_NS::utils::has_utf8` (const std::string &line)

Check if a string will likely have UTF-8 encoded characters

UTF-8 uses the 7-bit standard ASCII table for the first 127 characters and all other characters are encoded as multiple bytes. For the multi-byte characters the first byte has either the highest two, three, or four bits set followed by a zero bit and followed by one, two, or three more bytes, respectively, where the highest bit is set and the second highest bit set to 0. The remaining bits combined are the character code, which is thus limited to 21-bits.

For the sake of efficiency this test only checks if a character in the string has the highest bit set and thus is very likely an UTF-8 character. It will not be able to tell this this is a valid UTF-8 character or whether it is a 2-byte, 3-byte, or 4-byte character.

*See also*

[\*utils::utf8\\_subst\(\)\*](#)

**Parameters**

**line** – string that should be checked

**Returns**

true if string contains UTF-8 encoded characters (bool)

`std::string LAMMPS_NS::utils::utf8_subst` (const std::string &line)

Replace known UTF-8 characters with ASCII equivalents

*See also*

`utils::has_utf8()`

**Parameters**

**line** – string that should be converted

**Returns**

new string with ascii replacements (string)

`size_t LAMMPS_NS::utils::count_words(const char *text)`

Count words in C-string, ignore any whitespace matching `"\t\r\n\f"`

**Parameters**

**text** – string that should be searched

**Returns**

number of words found

`size_t LAMMPS_NS::utils::count_words(const std::string &text)`

Count words in string, ignore any whitespace matching `"\t\r\n\f"`

**Parameters**

**text** – string that should be searched

**Returns**

number of words found

`size_t LAMMPS_NS::utils::count_words(const std::string &text, const std::string &separators)`

Count words in string with custom choice of separating characters

**Parameters**

- **text** – string that should be searched
- **separators** – string containing characters that will be treated as whitespace

**Returns**

number of words found

`size_t LAMMPS_NS::utils::trim_and_count_words(const std::string &text, const std::string &separators = "\t\r\n\f")`

Count words in a single line, trim anything from `#` onward

**Parameters**

- **text** – string that should be trimmed and searched
- **separators** – string containing characters that will be treated as whitespace

**Returns**

number of words found

`std::string LAMMPS_NS::utils::join_words(const std::vector<std::string> &words, const std::string &sep)`

Take list of words and join them with a given separator text.

This is the inverse operation of what the `split_words()` function Tokenizer classes do.

**Parameters**

- **words** – STL vector with strings
- **sep** – separator string (may be empty)



**Returns**

string with the concatenated words and separators

```
std::vector<std::string> LAMMPS_NS::utils::split_words(const std::string &text)
```

Take text and split into non-whitespace words.

This can handle strings with single and double quotes, escaped quotes, and escaped codes within quotes, but due to using an STL container and STL strings is rather slow because of making copies. Designed for parsing command lines and similar text and not for time critical processing. Use a tokenizer class if performance matters.

*See also*

*Tokenizer, ValueTokenizer*

**Parameters**

**text** – string that should be split

**Returns**

STL vector with the words

```
std::vector<std::string> LAMMPS_NS::utils::split_lines(const std::string &text)
```

Take multi-line text and split into lines

**Parameters**

**text** – string that should be split

**Returns**

STL vector with the lines

```
bool LAMMPS_NS::utils::strmatch(const std::string &text, const std::string &pattern)
```

Match text against a simplified regex pattern

More flexible and specific matching of a string against a pattern. This function is supposed to be a more safe, more specific and simple to use API to find pattern matches. The purpose is to replace uses of either `strncmp()` or `strstr()` in the code base to find sub-strings safely. With `strncmp()` finding prefixes, the number of characters to match must be counted, which can lead to errors, while using “`^pattern`” will do the same with less problems. Matching for suffixes using `strstr()` is not as specific as ‘`pattern$`’, and complex matches, e.g. “`^rigid.*Vsmall.*`”, to match all small body optimized rigid fixes require only one test.

The use of `std::string` arguments allows for simple concatenation even with `char *` type variables. Example: `utils::strmatch(text, std::string("^") + charptr)`

**Parameters**

- **text** – the text to be matched against the pattern
- **pattern** – the search pattern, which may contain regexp markers

**Returns**

true if the pattern matches, false if not

```
std::string LAMMPS_NS::utils::strfind(const std::string &text, const std::string &pattern)
```

Find sub-string that matches a simplified regex pattern

This function is a companion function to `utils::strmatch()`. Arguments and logic is the same, but instead of a boolean, it returns the sub-string that matches the regex pattern. There can be only one match. This can be used as a more flexible alternative to `strstr()`.

**Parameters**

- **text** – the text to be matched against the pattern
- **pattern** – the search pattern, which may contain regexp markers

**Returns**

the string that matches the pattern or an empty one

`bool LAMMPS_NS::utils::is_integer(const std::string &str)`

Check if string can be converted to valid integer

**Parameters**

**str** – string that should be checked

**Returns**

true, if string contains valid a integer, false otherwise

`bool LAMMPS_NS::utils::is_double(const std::string &str)`

Check if string can be converted to valid floating-point number

**Parameters**

**str** – string that should be checked

**Returns**

true, if string contains valid number, false otherwise

`bool LAMMPS_NS::utils::is_id(const std::string &str)`

Check if string is a valid ID ID strings may contain only letters, numbers, and underscores.

**Parameters**

**str** – string that should be checked

**Returns**

true, if string contains valid id, false otherwise

`int LAMMPS_NS::utils::is_type(const std::string &str)`

Check if string is a valid type label, or numeric type, or numeric type range. Numeric type or type range may only contain digits or the '\*' character. Type label strings may not contain a digit, or a '\*', or a '#' character as the first character to distinguish them from comments and numeric types or type ranges. They also may not contain any whitespace. If the string is a valid numeric type or type range the function returns 0, if it is a valid type label the function returns 1, otherwise it returns -1.

**Parameters**

**str** – string that should be checked

**Returns**

0, 1, or -1, depending on whether the string is valid numeric type, valid type label or neither, respectively

### 15.14.4 Potential file functions

`std::string LAMMPS_NS::utils::get_potential_file_path(const std::string &path)`

Determine full path of potential file. If file is not found in current directory, search directories listed in LAMMPS\_POTENTIALS environment variable

**Parameters**

**path** – file path

**Returns**

full path to potential file

`std::string LAMMPS_NS::utils::get_potential_date(const std::string &path, const std::string &potential_name)`

Read potential file and return DATE field if it is present

**Parameters**

- **path** – file path
- **potential\_name** – name of potential that is being read

**Returns**

DATE field if present

`std::string LAMMPS_NS::utils::get_potential_units(const std::string &path, const std::string &potential_name)`

Read potential file and return UNITS field if it is present

**Parameters**

- **path** – file path
- **potential\_name** – name of potential that is being read

**Returns**

UNITS field if present

`int LAMMPS_NS::utils::get_supported_conversions(const int property)`

Return bitmask of available conversion factors for a given property

**Parameters**

**property** – property to be converted

**Returns**

bitmask indicating available conversions

`double LAMMPS_NS::utils::get_conversion_factor(const int property, const int conversion)`

Return unit conversion factor for given property and selected from/to units

**Parameters**

- **property** – property to be converted
- **conversion** – constant indicating the conversion

**Returns**

conversion factor

FILE \*LAMMPS\_NS::utils::open\_potential(const std::string &name, LAMMPS \*Imp, int \*auto\_convert)

Open a potential file as specified by *name*

If opening the file directly fails, the function will search for it in the list of folder pointed to by the environment variable LAMMPS\_POTENTIALS (if it is set).

If the potential file has a UNITS tag in the first line, the tag's value is compared to the current unit style setting. The behavior of the function then depends on the value of the *auto\_convert* parameter. If it is a null pointer, then the unit values must match or else the open will fail with an error. Otherwise the bitmask that *auto\_convert* points to is used check for compatibility with possible automatic conversions by the calling function. If compatible, the bitmask is set to the required conversion or `utils::NOCONVERT`.

#### Parameters

- **name** – file- or pathname of the potential file
- **Imp** – pointer to top-level LAMMPS class instance
- **auto\_convert** – pointer to unit conversion bitmask or `nullptr`

#### Returns

FILE pointer of the opened potential file or `nullptr`

## 15.14.5 Argument processing

template<typename TYPE>

void LAMMPS\_NS::utils::bounds(const char \*file, int line, const std::string &str, bigint nmin, bigint nmax, TYPE &nlo, TYPE &nhi, Error \*error)

Compute index bounds derived from a string with a possible wildcard

This functions processes the string in *str* and set the values of *nlo* and *nhi* according to the following five cases:

- a single number, *i*: *nlo* = *i*; *nhi* = *i*;
- a single asterisk, *\**: *nlo* = *nmin*; *nhi* = *nmax*;
- a single number followed by an asterisk, *i\**: *nlo* = *i*; *nhi* = *nmax*;
- a single asterisk followed by a number, *\*i*: *nlo* = *nmin*; *nhi* = *i*;
- two numbers with an asterisk in between. *i\*j*: *nlo* = *i*; *nhi* = *j*;

#### Parameters

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – string to be processed
- **nmin** – smallest possible lower bound
- **nmax** – largest allowed upper bound
- **nlo** – lower bound
- **nhi** – upper bound
- **error** – pointer to Error class for out-of-bounds messages

```
int LAMMPS_NS::utils::expand_args(const char *file, int line, int nargs, char **arg, int mode, char **&earg,
                                LAMMPS *lmp)
```

Expand list of arguments when containing fix/compute wildcards

This function searches the list of arguments in *arg* for strings of the kind `c_ID[*]` or `f_ID[*]` referring to computes or fixes. Any such strings are replaced by one or more strings with the `*` character replaced by the corresponding possible numbers as determined from the fix or compute instance. Other strings are just copied. If the *mode* parameter is set to 0, expand global vectors, but not global arrays; if it is set to 1, expand global arrays (by column) but not global vectors.

If any expansion happens, the *earg* list and all its strings are new allocations and must be freed explicitly by the caller. Otherwise *arg* and *earg* will point to the same address and no explicit de-allocation is needed by the caller.

#### Parameters

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **nargs** – number of arguments in current list
- **arg** – argument list, possibly containing wildcards
- **mode** – select between global vectors(=0) and arrays (=1)
- **earg** – new argument list with wildcards expanded
- **lmp** – pointer to top-level *LAMMPS* class instance

#### Returns

number of arguments in expanded list

```
char *LAMMPS_NS::utils::expand_type(const char *file, int line, const std::string &str, int mode, LAMMPS
                                   *lmp)
```

Expand type label string into its equivalent numeric type

This function checks if a given string may be a type label and then searches the labelmap type indicated by the *mode* argument for the corresponding numeric type. If this is found a copy of the numeric type string is made and returned. Otherwise a null pointer is returned. If a string is returned, the calling code must free it with `delete[]`.

#### Parameters

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **str** – type string to be expanded
- **mode** – select labelmap using constants from *Atom* class
- **lmp** – pointer to top-level *LAMMPS* class instance

#### Returns

pointer to expanded string or null pointer

## 15.14.6 Convenience functions

template<typename ...**Args**>

void LAMMPS\_NS::utils::logmesg(LAMMPS \*Imp, const std::string &format, *Args*&&... args)

Send formatted message to screen and logfile, if available

This function simplifies the repetitive task of outputting some message to both the screen and/or the log file. The template wrapper with `fntlib` format and argument processing allows this function to work similar to `fmt::print()`.

### Parameters

- **Imp** – pointer to *LAMMPS* class instance
- **format** – format string of message to be printed
- **args** – arguments to format string

void LAMMPS\_NS::utils::logmesg(LAMMPS \*Imp, const std::string &mesg)

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

### Parameters

- **Imp** – pointer to *LAMMPS* class instance
- **mesg** – string with message to be printed

std::string LAMMPS\_NS::utils::errorurl(int errorcode)

Return text redirecting the user to a specific paragraph in the manual

The *LAMMPS* manual contains detailed detailed explanations for errors and warnings where a simple error message may not be sufficient. These can be reached through URLs with a numeric code. This function creates the corresponding text to be included into the error message that redirects the user to that URL.

### Parameters

**errorcode** – number pointing to a paragraph in the manual

void LAMMPS\_NS::utils::missing\_cmd\_args(const std::string &file, int line, const std::string &cmd, Error \*error)

Print error message about missing arguments for command

This function simplifies the repetitive reporting missing arguments to a command.

### Parameters

- **file** – name of source file for error message
- **line** – line number in source file for error message
- **cmd** – name of the failing command
- **error** – pointer to Error class instance (for abort) or nullptr

void LAMMPS\_NS::utils::flush\_buffers(LAMMPS \*Imp)

Flush output buffers

This function calls `fflush()` on screen and logfile FILE pointers if available and thus tells the operating system to output all currently buffered data. This is local operation and independent from buffering by a file system or an MPI library.

std::string LAMMPS\_NS::utils::getsyserror()

Return a string representing the current system error status

This is a wrapper around calling strerror(errno).

**Returns**

error string

std::string LAMMPS\_NS::utils::check\_packages\_for\_style(const std::string &style, const std::string &name, *LAMMPS* \*Imp)

Report if a requested style is in a package or may have a typo

**Parameters**

- **style** – type of style that is to be checked for
- **name** – name of style that was not found
- **Imp** – pointer to top-level *LAMMPS* class instance

**Returns**

string usable for error messages

double LAMMPS\_NS::utils::timespec2seconds(const std::string &timespec)

Convert a time string to seconds

The strings “off” and “unlimited” result in -1

**Parameters**

**timespec** – a string in the following format: ([[HH:]MM:]SS)

**Returns**

total in seconds

int LAMMPS\_NS::utils::date2num(const std::string &date)

Convert a *LAMMPS* version date to a number

This will generate a number YYYYMMDD from a date string (with or without blanks) that is suitable for numerical comparisons, i.e. later dates will generate a larger number.

The day may or may not have a leading zero, the month is identified by the first 3 letters (so there may be more) and the year may be 2 or 4 digits (the missing 2 digits will be assumed as 20. That is 04 corresponds to 2004).

No check is made whether the date is valid.

**Parameters**

**date** – string in the format (Day Month Year)

**Returns**

date code

std::string LAMMPS\_NS::utils::current\_date()

Return current date as string

This will generate a string containing the current date in YYYY-MM-DD format.

**Returns**

string with current date

### 15.14.7 Customized standard functions

int LAMMPS\_NS::utils::binary\_search(const double needle, const int n, const double \*haystack)

Binary search in a vector of ascending doubles of length N

If the value is smaller than the smallest value in the vector, 0 is returned. If the value is larger or equal than the largest value in the vector, N-1 is returned. Otherwise the index that satisfies the condition

haystack[index] <= value < haystack[index+1]

is returned, i.e. a value from 1 to N-2. Note that if there are tied values in the haystack, always the larger index is returned as only that satisfied the condition.

#### Parameters

- **needle** – search value for which are are looking for the closest index
- **n** – size of the haystack array
- **haystack** – array with data in ascending order.

#### Returns

index of value in the haystack array smaller or equal to needle

void LAMMPS\_NS::utils::merge\_sort(int \*index, int num, void \*ptr, int (\*comp)(int, int, void\*))

Custom merge sort implementation

This function provides a custom upward hybrid merge sort implementation with support to pass an opaque pointer to the comparison function, e.g. for access to class members. This avoids having to use global variables. For improved performance, it uses an in-place insertion sort on initial chunks of up to 64 elements and switches to merge sort from then on.

#### Parameters

- **index** – Array with indices to be sorted
- **num** – Length of the index array
- **ptr** – Pointer to opaque object passed to comparison function
- **comp** – Pointer to comparison function

---

## 15.15 Special Math functions

The `MathSpecial` namespace implements a selection of custom and optimized mathematical functions for a variety of applications.

double LAMMPS\_NS::MathSpecial::factorial(const int n)

Fast tabulated factorial function

This function looks up pre-computed factorial values for arguments of  $n = 0$  to a maximum of 167, which is the maximal value representable by a double precision floating point number. For other values of  $n$  a NaN value is returned.

#### Parameters

**n** – argument (valid:  $0 \leq n \leq 167$ )

#### Returns

value of  $n!$  as double precision number or NaN



```
double LAMMPS_NS::MathSpecial::exp2_x86(double x)
```

Fast implementation of  $2^x$  without argument checks for little endian CPUs

This function implements an optimized version of `pow(2.0, x)` that does not check for valid arguments and thus may only be used where arguments are well behaved. The implementation makes assumptions about the layout of double precision floating point numbers in memory and thus will only work on little endian CPUs. If little endian cannot be safely detected, the result of calling `pow(2.0, x)` will be returned. This function also is the basis for the fast exponential `fm_exp(x)`.

**Parameters**

**x** – argument

**Returns**

value of  $2^x$  as double precision number

```
double LAMMPS_NS::MathSpecial::fm_exp(double x)
```

Fast implementation of `exp(x)` for little endian CPUs

This function implements an optimized version of `exp(x)` for little endian CPUs. It calls the `exp2_x86(x)` function with a suitable prefactor to `x` to return `exp(x)`. The implementation makes assumptions about the layout of double precision floating point numbers in memory and thus will only work on little endian CPUs. If little endian cannot be safely detected, the result of calling the `exp(x)` implementation in the standard math library will be returned.

**Parameters**

**x** – argument

**Returns**

value of  $e^x$  as double precision number

```
static inline double LAMMPS_NS::MathSpecial::my_erfcx(const double x)
```

Fast scaled error function complement `exp(x*x)*erfc(x)` for coul/long styles

This is a portable fast implementation of `exp(x*x)*erfc(x)` that can be used in coul/long pair styles as a replacement for the polynomial expansion that is/was widely used. Unlike the polynomial expansion, that is only accurate at the level of single precision floating point it provides full double precision accuracy, but at comparable speed (unlike the `erfc()` implementation shipped with GNU standard math library).

**Parameters**

**x** – argument

**Returns**

value of  $e^{(x*x)*erfc(x)}$

```
static inline double LAMMPS_NS::MathSpecial::expmsq(double x)
```

Fast implementation of `exp(-x*x)` for little endian CPUs for coul/long styles

This function implements an optimized version of `exp(-x*x)` based on `exp2_x86()` for use with little endian CPUs. If little endian cannot be safely detected, the result of calling the `exp(-x*x)` implementation in the standard math library will be returned.

**Parameters**

**x** – argument

**Returns**

value of  $e^{(-x*x)}$  as double precision number

```
static inline double LAMMPS_NS::MathSpecial::square(const double &x)
```

Fast inline version of `pow(x, 2.0)`

**Parameters**

**x** – argument

**Returns** $x^*x$ 

```
static inline double LAMMPS_NS::MathSpecial::cube(const double &x)
```

Fast inline version of `pow(x, 3.0)`

**Parameters**

**x** – argument

**Returns** $x^*x$ 

```
static inline double LAMMPS_NS::MathSpecial::powsign(const int n)
```

```
static inline double LAMMPS_NS::MathSpecial::powint(const double &x, const int n)
```

```
static inline double LAMMPS_NS::MathSpecial::powsinxx(const double &x, int n)
```

---

## 15.16 Tokenizer classes

The purpose of the tokenizer classes is to simplify the recurring task of breaking lines of text down into words and/or numbers. Traditionally, LAMMPS code would be using the `strtok()` function from the C library for that purpose, but that function has two significant disadvantages: 1) it cannot be used concurrently from different LAMMPS instances since it stores its status in a global variable and 2) it modifies the string that it is processing. These classes were implemented to avoid both of these issues and also to reduce the amount of code that needs to be written.

The basic procedure is to create an instance of the tokenizer class with the string to be processed as an argument and then do a loop until all available tokens are read. The constructor has a default set of separator characters, but that can be overridden. The default separators are all “whitespace” characters, i.e. the space character, the tabulator character, the carriage return character, the linefeed character, and the form feed character.

Listing 1: Tokenizer class example listing entries of the PATH environment variable

```
#include "tokenizer.h"
#include <stdlib>
#include <string>
#include <iostream>

using namespace LAMMPS_NS;

int main(int, char **)
{
    const char *path = getenv("PATH");

    if (path != nullptr) {
        Tokenizer p(path, ".");
        while (p.has_next())
            std::cout << "Entry: " << p.next() << "\n";
    }
    return 0;
}
```

Most tokenizer operations cannot fail except for `LAMMPS_NS::Tokenizer::next()` (when used without first checking with `LAMMPS_NS::Tokenizer::has_next()`) and `LAMMPS_NS::Tokenizer::skip()`. In case of failure, the class will throw an exception, so you may need to wrap the code using the tokenizer into a try / catch block to handle errors. The `LAMMPS_NS::ValueTokenizer` class may also throw an exception when a (type of) number is requested as next token that is not compatible with the string representing the next word.

Listing 2: ValueTokenizer class example with exception handling

```
#include "tokenizer.h"
#include <cstdlib>
#include <string>
#include <iostream>

using namespace LAMMPS_NS;

int main(int, char **)
{
    const char *text = "1 2 3 4 5 20.0 21 twentytwo 2.3";
    double num1(0), num2(0), num3(0), num4(0);

    ValueTokenizer t(text);
    // read 4 doubles after skipping over 5 numbers
    try {
        t.skip(5);
        num1 = t.next_double();
        num2 = t.next_double();
        num3 = t.next_double();
        num4 = t.next_double();
    } catch (TokenizerException &e) {
        std::cout << "Reading numbers failed: " << e.what() << "\n";
    }
    std::cout << "Values: " << num1 << " " << num2 << " " << num3 << " " << num4 << "\n";
    return 0;
}
```

This code example should produce the following output:

```
Reading numbers failed: Not a valid floating-point number: 'twentytwo'
Values: 20 21 0 0
```

class **Tokenizer**

### Public Functions

**Tokenizer**(std::string str, std::string separators = TOKENIZER\_DEFAULT\_SEPARATORS)

Class for splitting text into words

This tokenizer will break down a string into sub-strings (i.e words) separated by the given separator characters. If the string contains certain known UTF-8 characters they will be replaced by their ASCII equivalents processing the string.

*See also*

*ValueTokenizer*, *utils::split\_words()*, *utils::utf8\_subst()*

**Parameters**

- **str** – string to be processed
- **\_separators** – string with separator characters (default: ”\t\r\n\f”)

void **reset()**

Re-position the tokenizer state to the first word, i.e. the first non-separator character

void **skip**(int n = 1)

Skip over a given number of tokens

**Parameters**

**n** – number of tokens to skip over

bool **has\_next()** const

Indicate whether more tokens are available

**Returns**

true if there are more tokens, false if not

bool **contains**(const std::string &str) const

Search the text to be processed for a sub-string.

**Parameters**

**str** – string to be searched for

**Returns**

true if string was found, false if not

std::string **next()**

Retrieve next token.

**Returns**

string with the next token

size\_t **count()**

Count number of tokens in text.

**Returns**

number of counted tokens

std::vector<std::string> **as\_vector()**

Retrieve the entire text converted to an STL vector of tokens.

**Returns**

The STL vector

class **TokenizerException** : public exception

General Tokenizer exception class

Subclassed by *InvalidFloatException*, *InvalidIntegerException*

## Public Functions

explicit **TokenizerException**(const std::string &msg, const std::string &token)

Thrown during retrieving or skipping tokens

### Parameters

- **msg** – String with error message
- **token** – String of the token/word that caused the error

inline const char \***what**() const noexcept override

Retrieve message describing the thrown exception

### Returns

string with error message

class **ValueTokenizer**

## Public Functions

**ValueTokenizer**(const std::string &str, const std::string &separators =  
TOKENIZER\_DEFAULT\_SEPARATORS)

Class for reading text with numbers

*See also*

*Tokenizer*

**See also:**

Tokenizer *InvalidIntegerException InvalidFloatException*

### Parameters

- **str** – String to be processed
- **separators** – String with separator characters (default: "\t\r\n\f")

std::string **next\_string**()

Retrieve next token

### Returns

string with next token

tagint **next\_tagint**()

Retrieve next token and convert to tagint

### Returns

value of next token

bigint **next\_bigint**()

Retrieve next token and convert to bigint

### Returns

value of next token

int **next\_int**()

Retrieve next token and convert to int

**Returns**

value of next token

double **next\_double**()

Retrieve next token and convert to double

**Returns**

value of next token

bool **has\_next**() const

Indicate whether more tokens are available

**Returns**

true if there are more tokens, false if not

bool **contains**(const std::string &value) const

Search the text to be processed for a sub-string.

**Parameters**

**value** – string with value to be searched for

**Returns**

true if string was found, false if not

void **skip**(int ntokens = 1)

Skip over a given number of tokens

**Parameters**

**n** – number of tokens to skip over

size\_t **count**()

Count number of tokens in text.

**Returns**

number of counted tokens

class **InvalidIntegerException** : public *TokenizerException*

Exception thrown by ValueTokenizer when trying to convert an invalid integer string

### Public Functions

inline explicit **InvalidIntegerException**(const std::string &token)

Thrown during converting string to integer number

**Parameters**

**token** – String of the token/word that caused the error

class **InvalidFloatException** : public *TokenizerException*

Exception thrown by ValueTokenizer when trying to convert an floating point string

## Public Functions

inline explicit **InvalidFloatException**(const std::string &token)

Thrown during converting string to floating point number

### Parameters

**token** – String of the token/word that caused the error

## 15.17 Argument parsing classes

The purpose of argument parsing classes is to simplify and unify how arguments of commands in LAMMPS are parsed and to make abstractions of repetitive tasks.

The `LAMMPS_NS::ArgInfo` class provides an abstraction for parsing references to compute or fix styles, variables or custom integer or double properties handled by *fix property/atom*. These would start with a “c\_”, “f\_”, “v\_”, “d\_”, “d2\_”, “i\_”, or “i2\_” followed by the ID or name of that instance and may be postfixed with one or two array indices “[<number>]” with numbers > 0.

A typical code segment would look like this:

Listing 3: Usage example for ArgInfo class

```
int nvalues = 0;
for (iarg = 0; iarg < nargnew; iarg++) {
    ArgInfo argi(arg[iarg]);

    which[nvalues] = argi.get_type();
    argindex[nvalues] = argi.get_index1();
    ids[nvalues] = argi.copy_name();

    if ((which[nvalues] == ArgInfo::UNKNOWN)
        || (which[nvalues] == ArgInfo::NONE)
        || (argi.get_dim() > 1))
        error->all(FLERR, "Illegal compute XXX command");

    nvalues++;
}
```

class **ArgInfo**

### Public Types

enum **ArgTypes**

constants for argument types

*Values:*

enumerator **ERROR**

enumerator **UNKNOWN**

enumerator **NONE**

enumerator **X**

enumerator **V**

enumerator **F**

enumerator **COMPUTE**

enumerator **FIX**

enumerator **VARIABLE**

enumerator **KEYWORD**

enumerator **TYPE**

enumerator **MOLECULE**

enumerator **DNAME**

enumerator **INAME**

enumerator **DENSITY\_NUMBER**

enumerator **DENSITY\_MASS**

enumerator **MASS**

enumerator **TEMPERATURE**

enumerator **BIN1D**

enumerator **BIN2D**

enumerator **BIN3D**

enumerator **BINSPHERE**

enumerator **BINCYLINDER**



## Public Functions

**ArgInfo**(const std::string &arg, int allowed = *COMPUTE* | *FIX* | *VARIABLE*)

Class for processing references to fixes, computes and variables

This class provides an abstraction for the repetitive task of parsing arguments that may contain references to fixes, computes, variables, or custom per-atom properties. It will identify the name and the index value in the first and second dimension, if present.

### Parameters

- **arg** – string with possible reference
- **allowed** – integer with bitmap of allowed types of references

inline int **get\_type**() const

get type of reference

Return a type constant for the reference. This may be either *COMPUTE*, *FIX*, *VARIABLE* (if not restricted to a subset of those by the “allowed” argument of the constructor) or *NONE*, if it is not a recognized or allowed reference, or *UNKNOWN*, in case some error happened identifying or parsing the values of the indices

### Returns

integer with a constant from ArgTypes enumerator

inline int **get\_dim**() const

get dimension of reference

This will return either 0, 1, 2 depending on whether the reference has no, one or two “[{number}]” postfixes.

### Returns

integer with the dimensionality of the reference

inline int **get\_index1**() const

get index of first dimension

This will return the number in the first “[{number}]” postfix or 0 if there is no postfix.

### Returns

integer with index or the postfix or 0

inline int **get\_index2**() const

get index of second dimension

This will return the number in the second “[{number}]” postfix or -1 if there is no second postfix.

### Returns

integer with index of the postfix or -1

inline const char \***get\_name**() const

return reference to the ID or name of the reference

This string is pointing to an internal storage element and is only valid to use while the ArgInfo class instance is in scope. If you need a long-lived string make a copy with `copy_name()`.

### Returns

C-style char \* string

char \*copy\_name()

make copy of the ID of the reference as C-style string

The ID is copied into a buffer allocated with “new” and thus must be later deleted with “delete []” to avoid a memory leak. Because it is a full copy in a newly allocated buffer, the lifetime of this string extends beyond the the time the ArgInfo class is in scope.

**Returns**

copy of string as char \*

---

## 15.18 File reader classes

The purpose of the file reader classes is to simplify the recurring task of reading and parsing files. They can use the *ValueTokenizer* class to process the read in text. The *TextFileReader* is a more general version while *PotentialFileReader* is specialized to implement the behavior expected for looking up and reading/parsing files with potential parameters in LAMMPS. The potential file reader class requires a LAMMPS instance, requires to be run on MPI rank 0 only, will use the *utils::get\_potential\_file\_path* function to look up and open the file, and will call the *LAMMPS\_NS::Error* class in case of failures to read or to convert numbers, so that LAMMPS will be aborted.

Listing 4: Use of PotentialFileReader class in pair style coul/streitz

```
PotentialFileReader reader(lmp, file, "coul/streitz");
char * line;

while((line = reader.next_line(NPARAMS_PER_LINE))) {
  try {
    ValueTokenizer values(line);
    std::string iname = values.next_string();

    int ielement;
    for (ielement = 0; ielement < nelements; ielement++)
      if (iname == elements[ielement]) break;

    if (nparams == maxparam) {
      maxparam += DELTA;
      params = (Param *) memory->srealloc(params,maxparam*sizeof(Param),
                                         "pair:params");
    }

    params[nparams].ielement = ielement;
    params[nparams].chi = values.next_double();
    params[nparams].eta = values.next_double();
    params[nparams].gamma = values.next_double();
    params[nparams].zeta = values.next_double();
    params[nparams].zcore = values.next_double();

  } catch (TokenizerException & e) {
    error->one(FLERR, e.what());
  }
  nparams++;
}
```

A file that would be parsed by the reader code fragment looks like this:

```
# DATE: 2015-02-19 UNITS: metal CONTRIBUTOR: Ray Shan CITATION: Streitzi and Mintmire,  
↪Phys Rev B, 50, 11996-12003 (1994)  
#  
# X (eV)          J (eV)          gamma (1/AA)    zeta (1/AA)    Z (e)  
  
Al      0.000000      10.328655      0.000000      0.968438      0.763905  
O       5.484763      14.035715      0.000000      2.143957      0.000000
```

class **TextFileReader**

### Public Functions

**TextFileReader**(const std::string &filename, const std::string &filetype)

Class for reading and parsing text files

The value of the class member variable *ignore\_comments* controls whether any text following the pound sign (#) should be ignored (true) or not (false). Default: true, i.e. ignore.

*See also*

[TextFileReader](#)

#### Parameters

- **filename** – Name of file to be read
- **filetype** – Description of file type for error messages

**TextFileReader**(FILE \*fp, std::string filetype)

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

This function is useful in combination with [utils::open\\_potential\(\)](#).

---

**Note:** The FILE pointer is not closed in the destructor, but will be advanced when reading from it.

---

#### Parameters

- **fp** – File descriptor of the already opened file
- **filetype** – Description of file type for error messages

virtual **~TextFileReader**()

Closes the file

void **set\_bufsize**(int)

adjust line buffer size

void **rewind**()

Reset file to the beginning

void **skip\_line**()

Read the next line and ignore it

char **\*next\_line**(int nparams = 0)

Read the next line(s) until *nparams* words have been read.

This reads a line and counts the words in it, if the number is less than the requested number, it will read the next line, as well. Output will be a string with all read lines combined. The purpose is to somewhat replicate the reading behavior of formatted files in Fortran.

If the *ignore\_comments* class member has the value *true*, then any text read in is truncated at the first '#' character.

**Parameters**

**nparams** – Number of words that must be read. Default: 0

**Returns**

String with the concatenated text

void **next\_dvector**(double \*list, int n)

Read lines until *n* doubles have been read and stored in array *list*

This reads lines from the file using the *next\_line()* function, and splits them into floating-point numbers using the *ValueTokenizer* class and stores the number in the provided list.

**Parameters**

- **list** – Pointer to array with suitable storage for *n* doubles
- **n** – Number of doubles to be read

*ValueTokenizer* **next\_values**(int nparams, const std::string &separators =  
TOKENIZER\_DEFAULT\_SEPARATORS)

Read text until *nparams* words are read and passed to a tokenizer object for custom parsing.

This reads lines from the file using the *next\_line()* function, and splits them into floating-point numbers using the *ValueTokenizer* class and stores the number in the provided list.

**Parameters**

- **nparams** – Number of words to be read
- **separators** – String with list of separators.

**Returns**

*ValueTokenizer* object for read in text

## Public Members

bool **ignore\_comments**

Controls whether comments are ignored.

class **PotentialFileReader** : protected *Pointers*

## Public Functions

**PotentialFileReader**(class *LAMMPS* \*Imp, const std::string &filename, const std::string &potential\_name, const std::string &name\_suffix, const int auto\_convert = 0)

Class for reading and parsing *LAMMPS* potential files

The value of the class member variable *ignore\_comments* controls whether any text following the pound sign (#) should be ignored (true) or not (false). Default: true, i.e. ignore.

*See also*

*TextFileReader*

### Parameters

- **Imp** – Pointer to *LAMMPS* instance
- **filename** – Name of file to be read
- **potential\_name** – Name of potential style for error messages
- **name\_suffix** – Suffix added to potential name in error messages
- **auto\_convert** – Bitmask of supported unit conversions

**~PotentialFileReader**() override

Closes the file

void **ignore\_comments**(bool value)

Set comment (= text after '#') handling preference for the file to be read

### Parameters

**value** – Comment text is ignored if true, or not if false

void **rewind**()

Reset file to the beginning

void **skip\_line**()

Read a line but ignore its content

char \***next\_line**(int nparams = 0)

Read the next line(s) until *nparams* words have been read.

This reads a line and counts the words in it, if the number is less than the requested number, it will read the next line, as well. Output will be a string with all read lines combined. The purpose is to somewhat replicate the reading behavior of formatted files in Fortran.

### Parameters

**nparams** – Number of words that must be read. Default: 0

### Returns

String with the concatenated text

void **next\_dvector**(double \*list, int n)

Read lines until *n* doubles have been read and stored in array *list*

This reads lines from the file using the *next\_line*() function, and splits them into floating-point numbers using the *ValueTokenizer* class and stores the number in the provided list.

### Parameters

- **list** – Pointer to array with suitable storage for *n* doubles

- **n** – Number of doubles to be read

*ValueTokenizer* **next\_values**(int nparams, const std::string &separators =  
TOKENIZER\_DEFAULT\_SEPARATORS)

Read text until *nparams* words are read and passed to a tokenizer object for custom parsing.

This reads lines from the file using the `next_line()` function, and splits them into floating-point numbers using the `ValueTokenizer` class and stores the number in the provided list.

#### Parameters

- **nparams** – Number of words to be read
- **separators** – String with list of separators.

#### Returns

`ValueTokenizer` object for read in text

double **next\_double**()

Read next line and convert first word to a double

#### Returns

Value of first word in line as double

int **next\_int**()

Read next line and convert first word to an int

#### Returns

Value of first word in line as int

tagint **next\_tagint**()

Read next line and convert first word to a tagint

#### Returns

Value of first word in line as tagint

bigint **next\_bigint**()

Read next line and convert first word to a bigint

#### Returns

Value of first word in line as bigint

std::string **next\_string**()

Read next line and return first word

#### Returns

First word of read in line

---

## 15.19 Memory pool classes

The memory pool classes are used for cases where otherwise many small memory allocations would be needed and where the data would be either all used or all freed. One example for that is the storage of neighbor lists. The memory management strategy is based on the assumption that allocations will be in chunks of similar sizes. The allocation is then not done per individual call for a reserved chunk of memory, but for a “page” that can hold multiple chunks of data. A parameter for the maximum chunk size must be provided, as that is used to determine whether a new page of memory must be used.

The `MyPage` class offers two ways to reserve a chunk: 1) with `MyPage::get()` the chunk size needs to be known in advance, 2) with `MyPage::vget()` a pointer to the next chunk is returned, but its size is registered later with `MyPage::vgot()`.

Listing 5: Example of using `MyPage`

```
#include "my_page.h"
using namespace LAMMPS_NS;

MyPage<double> *dpage = new MyPage<double>;
// max size of chunk: 256, size of page: 10240 doubles (=81920 bytes)
dpage->init(256,10240);

double **build_some_lists(int num)
{
    dpage->reset();
    double **dlist = new double*[num];
    for (int i=0; i < num; ++i) {
        double *dptr = dpage.vget();
        int jnum = 0;
        for (int j=0; j < jmax; ++j) {
            // compute some dvalue for eligible loop index j
            dptr[j] = dvalue;
            ++jnum;
        }
        if (dpage.status() != 0) {
            // handle out of memory or jnum too large errors
        }
        dpage.vgot(jnum);
        dlist[i] = dptr;
    }
    return dlist;
}
```

```
template<class T>
```

```
class MyPage
```

Templated class for storing chunks of datums in pages.

The size of the chunk may vary from call to call, but must be less or equal than the `maxchunk` setting. The chunks are not returnable like with `malloc()` (i.e. you cannot call `free()` on them individually). One can only reset and start over. The purpose of this class is to replace many small memory allocations via `malloc()` with a few large ones. Since the pages are never freed until the class is re-initialized, they can be re-used without having to re-allocate them by calling the `reset()` method.

The settings `maxchunk`, `pagesize`, and `pagedelta` control the memory allocation strategy. The `maxchunk` value represents the expected largest number of items per chunk. If there is less space left on the current page, a new page is allocated for the next chunk. The `pagesize` value represents how many items can fit on a single page. It should have space for multiple chunks of size `maxchunk`. The combination of these two parameters determines how much memory is wasted by either switching to the next page too soon or allocating too large pages that never get properly used. It is an error, if a requested chunk is larger than `maxchunk`. The `pagedelta` parameter determines how many pages are allocated in one go. In combination with the `pagesize` setting, this determines how often blocks of memory get allocated (fewer allocations will result in faster execution).

**Note:** This is a template class with explicit instantiation. If the class is used with a new data type a new explicit instantiation may need to be added at the end of the file `src/my_page.cpp` to avoid symbol lookup errors.

---

## Public Functions

### MyPage()

Create a class instance

Need to call `init()` before use to define allocation settings

int **init**(int user\_maxchunk = 1, int user\_pagesize = 1024, int user\_pagedelta = 1)

(Re-)initialize the set of pages and allocation parameters.

This also frees all previously allocated storage and allocates the first page(s).

#### Parameters

- **user\_maxchunk** – Expected maximum number of items for one chunk
- **user\_pagesize** – Number of items on a single memory page
- **user\_pagedelta** – Number of pages to allocate with one malloc

#### Returns

1 if there were invalid parameters, 2 if there was an allocation error or 0 if successful

*T* \***get**(int n = 1)

Pointer to location that can store *N* items.

This will allocate more pages as needed. If the parameter *N* is larger than the *maxchunk* setting an error is flagged.

#### Parameters

**n** – number of items for which storage is requested

#### Returns

memory location or null pointer, if error or allocation failed

inline *T* \***vget**()

Get pointer to location that can store *maxchunk* items.

This will return the same pointer as the previous call to this function unless `vgot()` is called afterwards to record how many items of the chunk were actually used.

#### Returns

pointer to chunk of memory or null pointer if run out of memory

inline void **vgot**(int n)

Mark *N* items as used of the chunk reserved with a preceding call to `vget()`.

This will advance the internal pointer inside the current memory page. It is not necessary to call this function for *N* = 0, that is the reserved storage was not used. A following call to `vget()` will then reserve the same location again. It is an error if *N* > *maxchunk*.

#### Parameters

**n** – Number of items used in previously reserved chunk

void **reset**()

Reset state of memory pool without freeing any memory



```
inline double size() const
```

Return total size of allocated pages

**Returns**

total storage used in bytes

```
inline int status() const
```

Return error status

**Returns**

0 if no error, 1 requested chunk size > maxchunk, 2 if malloc failed

```
template<class T>
```

```
class MyPoolChunk
```

Templated class for storing chunks of datums in pages.

The size of the chunk may vary from call to call between the *minchunk* and *maxchunk* setting. Chunks may be returned to the pool for re-use. Chunks can be reserved in *nbin* different sizes between *minchunk* and *maxchunk*. The *chunksperpage* setting specifies how many chunks are stored on any page and the *pagedelta* setting determines how many pages are allocated in one go. Pages are never freed, so they can be re-used without re-allocation.

---

**Note:** This is a template class with explicit instantiation. If the class is used with a new data type a new explicit instantiation may need to be added at the end of the file `src/my_pool_chunk.cpp` to avoid symbol lookup errors.

---

## Public Functions

```
MyPoolChunk(int user_minchunk = 1, int user_maxchunk = 1, int user_nbin = 1, int user_chunkperpage = 1024, int user_pagedelta = 1)
```

Create a class instance and set memory pool parameters

**Parameters**

- **user\_minchunk** – Minimal chunk size
- **user\_maxchunk** – Maximal chunk size
- **user\_nbin** – Number of bins of different chunk sizes
- **user\_chunkperpage** – Number of chunks per page
- **user\_pagedelta** – Number of pages to allocate in one go

```
~MyPoolChunk()
```

Destroy class instance and free all allocated memory

```
T *get(int &index)
```

Return pointer/index of unused chunk of size maxchunk

**Parameters**

**index** – Index of chunk in memory pool

**Returns**

Pointer to requested chunk of storage

*T* \***get**(int n, int &index)

Return pointer/index of unused chunk of size N

**Parameters**

- **n** – Size of chunk
- **index** – Index of chunk in memory pool

**Returns**

Pointer to requested chunk of storage

void **put**(int index)

Put indexed chunk back into memory pool via free list

**Parameters**

**index** – Memory chunk index returned by call to *get()*

double **size**() const

Return total size of allocated pages

**Returns**

total storage used in bytes

inline int **status**() const

Return error status

**Returns**

0 if no error, 1 if invalid input, 2 if malloc() failed, 3 if chunk > maxchunk

---

## 15.20 Eigensolver functions

The `MathEigen` sub-namespace of the `LAMMPS_NS` namespace contains functions and classes for eigensolvers. Currently only the *jacobi3 function* is used in various places in LAMMPS. That function is built on top of a group of more generic eigensolvers that are maintained in the `math_eigen_impl.h` header file. This header contains the implementation of three template classes:

1. “Jacobi” calculates all of the eigenvalues and eigenvectors of a dense, symmetric, real matrix.
2. The “PEigenDense” class only calculates the principal eigenvalue (ie. the largest or smallest eigenvalue), and its corresponding eigenvector. However it is much more efficient than “Jacobi” when applied to large matrices (larger than 13x13). PEigenDense also can understand complex-valued Hermitian matrices.
3. The “LambdaLanczos” class is a generalization of “PEigenDense” which can be applied to arbitrary sparse matrices.

The “`math_eigen_impl.h`” code is an amalgamation of `jacobi_pd` by Andrew Jewett at Scripps Research (under CC0-1.0 license) and `LambdaLanczos` by Yuya Kurebayashi at Tohoku University (under MIT license)

---

int `MathEigen::jacobi3`(double const \*const \*mat, double \*eval, double \*\*evec)

A specialized function which finds the eigenvalues and eigenvectors of a 3x3 matrix (in double \*\* format).

**Parameters**

- **mat** – the 3x3 matrix you wish to diagonalize
  - **eval** – store the eigenvalues here
-

- **evect** – store the eigenvectors here...

### Returns

0 if eigenvalue calculation converged, 1 if it failed

```
int MathEigen::jacobi3(double const mat[3][3], double *eval, double evect[3][3])
```

This is an overloaded member function, provided for convenience. It differs from the above function only in what argument(s) it accepts.

## 15.21 Communication buffer coding with *ubuf*

LAMMPS uses communication buffers where it collects data from various class instances and then exchanges the data with neighboring sub-domains. For simplicity those buffers are defined as `double` buffers and used for doubles and integer numbers. This presents a unique problem when 64-bit integers are used. While the storage needed for a `double` is also 64-bit, it cannot be used by a simple assignment. To get around that limitation, LAMMPS uses the *ubuf* union. It is used in the various “pack” and “unpack” functions in the LAMMPS classes to store and retrieve integers that may be 64-bit from the communication buffers.

union **ubuf**

*#include <lmptype.h>* Data structure for packing 32-bit and 64-bit integers into double (communication) buffers

Using this union avoids aliasing issues by having member types (`double`, `int`) referencing the same buffer memory location.

The explicit constructor for 32-bit integers prevents compilers from (incorrectly) calling the double constructor when storing an `int` into a double buffer.

**Usage:**

Listing 6: To copy an integer into a double buffer:

```
double buf[2];
int    foo = 1;
tagint bar = 2<<40;
buf[1] = ubuf(foo).d;
buf[2] = ubuf(bar).d;
```

Listing 7: To copy from a double buffer back to an int:

```
foo = (int)    ubuf(buf[1]).i;  
bar = (tagint) ubuf(buf[2]).i;
```

The typecasts prevent compiler warnings about possible truncation issues.

### Public Functions

inline **ubuf**(const double &arg)

inline **ubuf**(const int64\_t &arg)

inline **ubuf**(const int &arg)

### Public Members

double **d**

int64\_t **i**

## **Part III**

# **Command Reference**



## COMMANDS

### 16.1 angle\_coeff command

#### 16.1.1 Syntax

```
angle_coeff N args
```

- N = numeric angle type (see asterisk form below), or type label
- args = coefficients for one or more angle types

#### 16.1.2 Examples

```
angle_coeff 1 300.0 107.0
angle_coeff * 5.0
angle_coeff 2*10 5.0

labelmap angle 1 hydroxyl
angle_coeff hydroxyl 300.0 107.0
```

#### 16.1.3 Description

Specify the angle force field coefficients for one or more angle types. The number and meaning of the coefficients depends on the angle style. Angle coefficients can also be set in the data file read by the *read\_data* command or in a restart file.

*N* can be specified in one of two ways. An explicit numeric value can be used, as in the first example above. Or *N* can be a type label, which is an alphanumeric string defined by the *labelmap* command or in a section of a data file read by the *read\_data* command.

For numeric values only, a wild-card asterisk can be used to set the coefficients for multiple angle types. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If *N* is the number of angle types, then an asterisk with no numeric values means all types from 1 to *N*. A leading asterisk means all types from 1 to *n* (inclusive). A trailing asterisk means all types from *n* to *N* (inclusive). A middle asterisk means all types from *m* to *n* (inclusive).

Note that using an *angle\_coeff* command can override a previous setting for the same angle type. For example, these commands set the coeffs for all angle types, then overwrite the coeffs for just angle type 2:

```
angle_coeff * 200.0 107.0 1.2
angle_coeff 2 50.0 107.0
```

A line in a data file that specifies angle coefficients uses the exact same format as the arguments of the *angle\_coeff* command in an input script, except that wild-card asterisks should not be used since coefficients for all  $N$  types must be listed in the file. For example, under the “Angle Coeffs” section of a data file, the line that corresponds to the first example above would be listed as

```
1 300.0 107.0
```

The *angle\_style class2* is an exception to this rule, in that an additional argument is used in the input script to allow specification of the cross-term coefficients. See its doc page for details.

---

The list of all angle styles defined in LAMMPS is given on the *angle\_style* doc page. They are also listed in more compact form on the *Commands angle* doc page.

On either of those pages, click on the style to display the formula it computes and its coefficients as specified by the associated *angle\_coeff* command.

---

## 16.1.4 Restrictions

This command must come after the simulation box is defined by a *read\_data*, *read\_restart*, or *create\_box* command.

An angle style must be defined before any angle coefficients are set, either in the input script or in a data file.

## 16.1.5 Related commands

*angle\_style*

## 16.1.6 Default

none

# 16.2 angle\_style command

## 16.2.1 Syntax

```
angle_style style
```

- style = *none* or *zero* or *hybrid* or *amoeba* or *charmm* or *class2* or *class2/p6* or *cosine* or *cosine/buck6d* or *cosine/delta* or *cosine/periodic* or *cosine/shift* or *cosine/shift/exp* or *cosine/squared* or *cross* or *dipole* or *fourier* or *fourier/simple* or *gaussian* or *harmonic* or *mm3* or *quartic* or *spica* or *table*



## 16.2.2 Examples

```
angle_style harmonic
angle_style charmm
angle_style hybrid harmonic cosine
```

## 16.2.3 Description

Set the formula(s) LAMMPS uses to compute angle interactions between triplets of atoms, which remain in force for the duration of the simulation. The list of angle triplets is read in by a *read\_data* or *read\_restart* command from a data or restart file.

Hybrid models where angles are computed using different angle potentials can be setup using the *hybrid* angle style.

The coefficients associated with a angle style can be specified in a data or restart file or via the *angle\_coeff* command.

All angle potentials store their coefficient data in binary restart files which means *angle\_style* and *angle\_coeff* commands do not need to be re-specified in an input script that restarts a simulation. See the *read\_restart* command for details on how to do this. The one exception is that *angle\_style hybrid* only stores the list of sub-styles in the restart file; angle coefficients need to be re-specified.

---

**Note:** When both an angle and pair style is defined, the *special\_bonds* command often needs to be used to turn off (or weight) the pairwise interaction that would otherwise exist between 3 bonded atoms.

---

In the formulas listed for each angle style, *theta* is the angle between the 3 atoms in the angle.

---

Here is an alphabetic list of angle styles defined in LAMMPS. Click on the style to display the formula it computes and coefficients specified by the associated *angle\_coeff* command.

Click on the style to display the formula it computes, any additional arguments specified in the *angle\_style* command, and coefficients specified by the associated *angle\_coeff* command.

There are also additional accelerated pair styles included in the LAMMPS distribution for faster performance on CPUs, GPUs, and KNLs. The individual style names on the [Commands angle](#) page are followed by one or more of (g,i,k,o,t) to indicate which accelerated styles exist.

- *none* - turn off angle interactions
- *zero* - topology but no interactions
- *hybrid* - define multiple styles of angle interactions
- *amoeba* - AMOEBA angle
- *charmm* - CHARMM angle
- *class2* - COMPASS (class 2) angle
- *class2/p6* - COMPASS (class 2) angle expanded to 6th order
- *cosine* - angle with cosine term
- *cosine/buck6d* - same as cosine with Buckingham term between 1-3 atoms
- *cosine/delta* - angle with difference of cosines
- *cosine/periodic* - DREIDING angle
- *cosine/shift* - angle cosine with a shift

- *cosine/shift/exp* - cosine with shift and exponential term in spring constant
  - *cosine/squared* - angle with cosine squared term
  - *cross* - cross term coupling angle and bond lengths
  - *dipole* - angle that controls orientation of a point dipole
  - *fourier* - angle with multiple cosine terms
  - *fourier/simple* - angle with a single cosine term
  - *gaussian* - multi-centered Gaussian-based angle potential
  - *harmonic* - harmonic angle
  - *mesocnt* - piecewise harmonic and linear angle for bending-buckling of nanotubes
  - *mm3* - anharmonic angle
  - *quartic* - angle with cubic and quartic terms
  - *spica* - harmonic angle with repulsive SPICA pair style between 1-3 atoms
  - *table* - tabulated by angle
- 

## 16.2.4 Restrictions

Angle styles can only be set for atom\_styles that allow angles to be defined.

Most angle styles are part of the MOLECULE package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info. The doc pages for individual bond potentials tell if it is part of a package.

## 16.2.5 Related commands

*angle\_coeff*

## 16.2.6 Default

```
angle_style none
```

## 16.3 atom\_modify command

### 16.3.1 Syntax

```
atom_modify keyword values ...
```

- one or more keyword/value pairs may be appended
- keyword = *id* or *map* or *first* or *sort*

```

id value = yes or no
map value = yes or array or hash
first value = group-ID = group whose atoms will appear first in internal atom lists
sort values = Nfreq binsize
    Nfreq = sort atoms spatially every this many time steps
    binsize = bin size for spatial sorting (distance units)

```

### 16.3.2 Examples

```

atom_modify map yes
atom_modify map hash sort 10000 2.0
atom_modify first colloid

```

### 16.3.3 Description

Modify certain attributes of atoms defined and stored within LAMMPS, in addition to what is specified by the *atom\_style* command. The *id* and *map* keywords must be specified before a simulation box is defined; other keywords can be specified any time.

The *id* keyword determines whether non-zero atom IDs can be assigned to each atom. If the value is *yes*, which is the default, IDs are assigned, whether you use the *create\_atoms* or *read\_data* or *read\_restart* commands to initialize atoms. If the value is *no* the IDs for all atoms are assumed to be 0.

If atom IDs are used, they must all be positive integers. They should also be unique, though LAMMPS does not check for this. Typically they should also be consecutively numbered (from 1 to *Natoms*), though this is not required. Molecular *atom styles* are those that store bond topology information (styles *bond*, *angle*, *molecular*, *full*). These styles require atom IDs since the IDs are used to encode the topology. Some other LAMMPS commands also require the use of atom IDs. E.g. some many-body pair styles use them to avoid double computation of the I-J interaction between two atoms.

The only reason not to use atom IDs is if you are running an atomic simulation so large that IDs cannot be uniquely assigned. For a default LAMMPS build this limit is  $2^{31}$  or about 2 billion atoms. However, even in this case, you can use 64-bit atom IDs, allowing  $2^{63}$  or about  $9e18$  atoms, if you build LAMMPS with the `-DLAMMPS_BIGBIG` switch. This is described on the *Build\_settings* doc page. If atom IDs are not used, they must be specified as 0 for all atoms, e.g. in a data or restart file.

The *map* keyword determines how atoms with specific IDs are found when required. An example are the *bond* (*angle*, etc) methods which need to find the local index of an atom with a specific global ID which is a *bond* (*angle*, etc) partner. LAMMPS performs this operation efficiently by creating a “map”, which is either an *array* or *hash* table, as described below.

When the *map* keyword is not specified in your input script, LAMMPS only creates a map for *atom\_styles* for molecular systems which have permanent bonds (*angles*, etc). No map is created for atomic systems, since it is normally not needed. However some LAMMPS commands require a map, even for atomic systems, and will generate an error if one does not exist. The *map* keyword thus allows you to force the creation of a map. The *yes* value will create either an *array* or *hash* style map, as explained in the next paragraph. The *array* and *hash* values create an array-style or hash-style map respectively.

For an *array*-style map, each processor stores a lookup table of length *N*, where *N* is the largest atom ID in the system. This is a fast, simple method for many simulations, but requires too much memory for large simulations. For a *hash*-style map, a hash table is created on each processor, which finds an atom ID in constant time (independent of the global number of atom IDs). It can be slightly slower than the *array* map, but its memory cost is proportional to the number of atoms owned by a processor, i.e.  $N/P$  when *N* is the total number of atoms in the system and *P* is the number of processors.

The *first* keyword allows a *group* to be specified whose atoms will be maintained as the first atoms in each processor's list of owned atoms. This is only useful when the specified group is a small fraction of all the atoms, and there are other operations LAMMPS is performing that will be sped-up significantly by being able to loop over the smaller set of atoms. Otherwise the reordering required by this option will be a net slow-down. The *neigh\_modify include* and *comm\_modify group* commands are two examples of commands that require this setting to work efficiently. Several *fixes*, most notably time integration fixes like *fix nve*, also take advantage of this setting if the group they operate on is the group specified by this command. Note that specifying "all" as the group-ID effectively turns off the *first* option.

It is OK to use the *first* keyword with a group that has not yet been defined, e.g. to use the *atom\_modify first* command at the beginning of your input script. LAMMPS does not use the group until a simulation is run.

The *sort* keyword turns on a spatial sorting or reordering of atoms within each processor's sub-domain every *Nfreq* timesteps. If *Nfreq* is set to 0, then sorting is turned off. Sorting can improve cache performance and thus speed-up a LAMMPS simulation, as discussed in a paper by (*Meloni*). Its efficacy depends on the problem size (atoms/processor), how quickly the system becomes disordered, and various other factors. As a general rule, sorting is typically more effective at speeding up simulations of liquids as opposed to solids. In tests we have done, the speed-up can range from zero to 3-4x.

Reordering is performed every *Nfreq* timesteps during a dynamics run or iterations during a minimization. More precisely, reordering occurs at the first reneighboring that occurs after the target timestep. The reordering is performed locally by each processor, using bins of the specified *binsize*. If *binsize* is set to 0.0, then a binsize equal to half the *neighbor* cutoff distance (force cutoff plus skin distance) is used, which is a reasonable value. After the atoms have been binned, they are reordered so that atoms in the same bin are adjacent to each other in the processor's 1d list of atoms.

The goal of this procedure is for atoms to put atoms close to each other in the processor's one-dimensional list of atoms that are also near to each other spatially. This can improve cache performance when pairwise interactions and neighbor lists are computed. Note that if bins are too small, there will be few atoms/bin. Likewise if bins are too large, there will be many atoms/bin. In both cases, the goal of cache locality will be undermined.

---

**Note:** Running a simulation with sorting on versus off should not change the simulation results in a statistical sense. However, a different ordering will induce round-off differences, which will lead to diverging trajectories over time when comparing two simulations. Various commands, particularly those which use random numbers (e.g. *velocity create*, and *fix langevin*), may generate (statistically identical) results which depend on the order in which atoms are processed. The order of atoms in a *dump* file will also typically change if sorting is enabled.

---

### 16.3.4 Restrictions

The *first* and *sort* options cannot be used together. Since sorting is on by default, it will be turned off if the *first* keyword is used with a group-ID that is not "all".

### 16.3.5 Related commands

none

### 16.3.6 Default

By default, *id* is yes. By default, atomic systems (no bond topology info) do not use a map. For molecular systems (with bond topology info), a map is used. The default map style is array if no atom ID is larger than 1 million, otherwise the default is hash. By default, a “first” group is not defined. By default, sorting is enabled with a frequency of 1000 and a binsize of 0.0, which means the neighbor cutoff will be used to set the bin size. If no neighbor cutoff is defined, sorting will be turned off.

(Meloni) Meloni, Rosati and Colombo, J Chem Phys, 126, 121102 (2007).

## 16.4 atom\_style command

### 16.4.1 Syntax

```
atom_style style args
```

- style = *amoeba* or *angle* or *atomic* or *body* or *bond* or *charge* or *dielectric* or *dipole* or *dpd* or *edpd* or *electron* or *ellipsoid* or *full* or *line* or *mdpd* or *mesont* or *molecular* or *oxdna* or *peri* or *smd* or *sph* or *sphere* or *bpm/sphere* or *spin* or *tdpd* or *tri* or *template* or *wavepacket* or *hybrid*

args = none for any style except the following

*body* args = *bstyle* *bstyle-args*

*bstyle* = style of body particles

*bstyle-args* = additional arguments specific to the *bstyle*

see the [Howto body](#) doc

page for details

*sphere* arg = 0/1 (optional) for static/dynamic particle radii

*bpm/sphere* arg = 0/1 (optional) for static/dynamic particle radii

*tdpd* arg = *Nspecies*

*Nspecies* = # of chemical species

*template* arg = *template-ID*

*template-ID* = ID of molecule template specified in a separate [molecule](#) command

*hybrid* args = list of one or more sub-styles, each with their args

- accelerated styles (with same args) = *angle/kk* or *atomic/kk* or *bond/kk* or *charge/kk* or *full/kk* or *molecular/kk* or *spin/kk*

### 16.4.2 Examples

```
atom_style atomic
atom_style bond
atom_style full
atom_style body nparticle 2 10
atom_style hybrid charge bond
atom_style hybrid charge body nparticle 2 5
atom_style spin
atom_style template myMols
atom_style hybrid template twomols charge
atom_style tdpd 2
```

### 16.4.3 Description

Define what style of atoms to use in a simulation. This determines what attributes are associated with the atoms. This command must be used before a simulation is setup via a *read\_data*, *read\_restart*, or *create\_box* command.

**Note:** Many of the atom styles discussed here are only enabled if LAMMPS was built with a specific package, as listed below in the Restrictions section.

Once a style is assigned, it cannot be changed, so use a style general enough to encompass all attributes. E.g. with style *bond*, angular terms cannot be used or added later to the model. It is OK to use a style more general than needed, though it may be slightly inefficient.

The choice of style affects what quantities are stored by each atom, what quantities are communicated between processors to enable forces to be computed, and what quantities are listed in the data file read by the *read\_data* command.

These are the additional attributes of each style and the typical kinds of physical systems they are used to model. All styles store coordinates, velocities, atom IDs and types. See the *read\_data*, *create\_atoms*, and *set* commands for info on how to set these various quantities.

<i>amoeba</i>	molecular + charge + 1/5 neighbors	AMOEBHA/HIPPO polarized force fields
<i>angle</i>	bonds and angles	bead-spring polymers with stiffness
<i>atomic</i>	only the default values	coarse-grain liquids, solids, metals
<i>body</i>	mass, inertia moments, quaternion, angular momentum	arbitrary bodies
<i>bond</i>	bonds	bead-spring polymers
<i>charge</i>	charge	atomic system with charges
<i>dielectric</i>	dipole, area, curvature	system with surface polarization
<i>dipole</i>	charge and dipole moment	system with dipolar particles
<i>dpd</i>	internal temperature and internal energies	DPD particles
<i>edpd</i>	temperature and heat capacity	eDPD particles
<i>electron</i>	charge and spin and eradius	electronic force field
<i>ellipsoid</i>	shape, quaternion, angular momentum	aspherical particles
<i>full</i>	molecular + charge	bio-molecules
<i>line</i>	end points, angular velocity	rigid bodies
<i>mdpd</i>	density	mDPD particles
<i>mesont</i>	mass, radius, length, buckling, connections, tube id	mesoscopic nanotubes
<i>molecular</i>	bonds, angles, dihedrals, impropers	uncharged molecules
<i>oxdna</i>	nucleotide polarity	coarse-grained DNA and RNA models
<i>peri</i>	mass, volume	mesoscopic Peridynamic models
<i>smd</i>	volume, kernel diameter, contact radius, mass	solid and fluid SPH particles
<i>sph</i>	rho, esph, cv	SPH particles
<i>sphere</i>	diameter, mass, angular velocity	granular models
<i>bpm/sphere</i>	diameter, mass, angular velocity, quaternion	granular bonded particle models (BPM)
<i>spin</i>	magnetic moment	system with magnetic particles
<i>tdpd</i>	chemical concentration	tDPD particles
<i>template</i>	template index, template atom	small molecules with fixed topology
<i>tri</i>	corner points, angular momentum	rigid bodies
<i>wavepacket</i>	charge, spin, eradius, etag, cs_re, cs_im	AWPMD

**Note:** It is possible to add some attributes, such as a molecule ID, to atom styles that do not have them via the *fix property/atom* command. This command also allows new custom attributes consisting of extra integer or floating-point values to be added to atoms. See the *fix property/atom* page for examples of cases where this is useful and details on

how to initialize, access, and output the custom values.

All of the above styles define point particles, except the *sphere*, *bpm/sphere*, *ellipsoid*, *electron*, *peri*, *wavepacket*, *line*, *tri*, and *body* styles, which define finite-size particles. See the [Howto spherical](#) page for an overview of using finite-size particle models with LAMMPS.

All of the point-particle styles assign mass to particles on a per-type basis, using the *mass* command, The finite-size particle styles assign mass to individual particles on a per-particle basis.

For the *sphere* and *bpm/sphere* styles, the particles are spheres and each stores a per-particle diameter and mass. If the diameter > 0.0, the particle is a finite-size sphere. If the diameter = 0.0, it is a point particle. Note that by use of the *disc* keyword with the *fix nve/sphere*, *fix nvt/sphere*, *fix nph/sphere*, *fix npt/sphere* commands for the *sphere* style, spheres can be effectively treated as 2d discs for a 2d simulation if desired. See also the *set density/disc* command. The *sphere* and *bpm/sphere* styles take an optional 0 or 1 argument. A value of 0 means the radius of each sphere is constant for the duration of the simulation. A value of 1 means the radii may vary dynamically during the simulation, e.g. due to use of the *fix adapt* command.

For the *ellipsoid* style, the particles are ellipsoids and each stores a flag which indicates whether it is a finite-size ellipsoid or a point particle. If it is an ellipsoid, it also stores a shape vector with the 3 diameters of the ellipsoid and a quaternion 4-vector with its orientation.

For the *dielectric* style, each particle can be either a physical particle (e.g. an ion), or an interface particle representing a boundary element. For physical particles, the per-particle properties are the same as *atom\_style full*. For interface particles, in addition to these properties, each particle also has an area, a normal unit vector, a mean local curvature, the mean and difference of the dielectric constants of two sides of the interface, and the local dielectric constant at the boundary element. The distinction between the physical and interface particles is only meaningful when *fix polarize* commands are applied to the interface particles.

For the *dipole* style, a point dipole is defined for each point particle. Note that if you wish the particles to be finite-size spheres as in a Stockmayer potential for a dipolar fluid, so that the particles can rotate due to dipole-dipole interactions, then you need to use *atom\_style hybrid sphere dipole*, which will assign both a diameter and dipole moment to each particle.

For the *electron* style, the particles representing electrons are 3d Gaussians with a specified position and bandwidth or uncertainty in position, which is represented by the *eradius* = electron size.

For the *peri* style, the particles are spherical and each stores a per-particle mass and volume.

The *bpm/sphere* style is part of the BPM package.

The *oxdna* style is for coarse-grained nucleotides and stores the 3'-to-5' polarity of the nucleotide strand, which is set through the bond topology in the data file. The first (second) atom in a bond definition is understood to point towards the 3'-end (5'-end) of the strand. Note that this style is part of the CG-DNA package.

The *dpd* style is for dissipative particle dynamics (DPD) particles. Note that it is part of the DPD-REACT package, and is not for use with the *pair\_style dpd* or *dpd/stat* commands, which can simply use *atom\_style atomic*. *Atom\_style dpd* extends DPD particle properties with internal temperature (*dpdTheta*), internal conductive energy (*uCond*), internal mechanical energy (*uMech*), and internal chemical energy (*uChem*).

The *edpd* style is for energy-conserving dissipative particle dynamics (eDPD) particles which store a temperature (*edpd\_temp*), and heat capacity (*edpd\_cv*).

The *mdpd* style is for many-body dissipative particle dynamics (mDPD) particles which store a density (*rho*) for considering density-dependent many-body interactions.

The *tdpd* style is for transport dissipative particle dynamics (tDPD) particles which store a set of chemical concentration. An integer "cc\_species" is required to specify the number of chemical species involved in a tDPD system.

The *sph* style is for smoothed particle hydrodynamics (SPH) particles which store a density (*rho*), energy (*esph*), and heat capacity (*cv*).

The *smd* style is for a general formulation of Smooth Particle Hydrodynamics. Both fluids and solids can be modeled. Particles store the mass and volume of an integration point, a kernel diameter used for calculating the field variables (e.g. stress and deformation) and a contact radius for calculating repulsive forces which prevent individual physical bodies from penetrating each other.

For the *spin* style, a magnetic spin is associated to each atom. Those spins have a norm (their magnetic moment) and a direction.

The *wavepacket* style is similar to *electron*, but the electrons may consist of several Gaussian wave packets, summed up with coefficients  $cs = (cs\_re, cs\_im)$ . Each of the wave packets is treated as a separate particle in LAMMPS, wave packets belonging to the same electron must have identical *etag* values.

For the *line* style, the particles are idealized line segments and each stores a per-particle mass and length and orientation (i.e. the end points of the line segment).

For the *tri* style, the particles are planar triangles and each stores a per-particle mass and size and orientation (i.e. the corner points of the triangle).

The *template* style allows molecular topology (bonds, angles, etc) to be defined via a molecule template using the *molecule* command. The template stores one or more molecules with a single copy of the topology info (bonds, angles, etc) of each. Individual atoms only store a template index and template atom to identify which molecule and which atom-within-the-molecule they represent. Using the *template* style instead of the *bond*, *angle*, *molecular* styles can save memory for systems comprised of a large number of small molecules, all of a single type (or small number of types). See the paper by Grime and Voth, in (*Grime*), for examples of how this can be advantageous for large-scale coarse-grained systems. The `examples/template` directory has a few demo inputs and examples showing the use of the *template* atom style versus *molecular*.

---

**Note:** When using the *template* style with a *molecule template* that contains multiple molecules, you should insure the atom types, bond types, angle\_types, etc in all the molecules are consistent. E.g. if one molecule represents H2O and another CO2, then you probably do not want each molecule file to define 2 atom types and a single bond type, because they will conflict with each other when a mixture system of H2O and CO2 molecules is defined, e.g. by the *read\_data* command. Rather the H2O molecule should define atom types 1 and 2, and bond type 1. And the CO2 molecule should define atom types 3 and 4 (or atom types 3 and 2 if a single oxygen type is desired), and bond type 2.

---

For the *body* style, the particles are arbitrary bodies with internal attributes defined by the “style” of the bodies, which is specified by the *bstyle* argument. Body particles can represent complex entities, such as surface meshes of discrete points, collections of sub-particles, deformable objects, etc.

The *Howto body* page describes the body styles LAMMPS currently supports, and provides more details as to the kind of body particles they represent. For all styles, each body particle stores moments of inertia and a quaternion 4-vector, so that its orientation and position can be time integrated due to forces and torques.

Note that there may be additional arguments required along with the *bstyle* specification, in the `atom_style body` command. These arguments are described on the *Howto body* doc page.

---

Typically, simulations require only a single (non-hybrid) atom style. If some atoms in the simulation do not have all the properties defined by a particular style, use the simplest style that defines all the needed properties by any atom. For example, if some atoms in a simulation are charged, but others are not, use the *charge* style. If some atoms have bonds, but others do not, use the *bond* style.

The only scenario where the *hybrid* style is needed is if there is no single style which defines all needed properties of all atoms. For example, as mentioned above, if you want dipolar particles which will rotate due to torque, you need to use “`atom_style hybrid sphere dipole`”. When a hybrid style is used, atoms store and communicate the union of all quantities implied by the individual styles.

When using the *hybrid* style, you cannot combine the *template* style with another molecular style that stores bond, angle, etc info on a per-atom basis.



LAMMPS can be extended with new atom styles as well as new body styles; see the [Modify](#) doc page.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 16.4.4 Restrictions

This command cannot be used after the simulation box is defined by a *read\_data* or *create\_box* command.

Many of the styles listed above are only enabled if LAMMPS was built with a specific package, as listed below. See the [Build package](#) page for more info.

The *amoeba* style is part of the AMOEBA package.

The *angle*, *bond*, *full*, *molecular*, and *template* styles are part of the MOLECULE package.

The *line* and *tri* styles are part of the ASPHERE package.

The *body* style is part of the BODY package.

The *dipole* style is part of the DIPOLE package.

The *peri* style is part of the PERI package for Peridynamics.

The *oxdna* style is part of the CG-DNA package for coarse-grained simulation of DNA and RNA.

The *electron* style is part of the EFF package for *electronic force fields*.

The *dpd* style is part of the DPD-REACT package for dissipative particle dynamics (DPD).

The *edpd*, *mdpd*, and *tdpd* styles are part of the DPD-MESO package for energy-conserving dissipative particle dynamics (eDPD), many-body dissipative particle dynamics (mDPD), and transport dissipative particle dynamics (tDPD), respectively.

The *sph* style is part of the SPH package for smoothed particle hydrodynamics (SPH). See [this PDF guide](#) to using SPH in LAMMPS.

The *mesont* style is part of the MESONT package.

The *spin* style is part of the SPIN package.

The *wavepacket* style is part of the AWPMD package for the *antisymmetrized wave packet MD method*.

## 16.4.5 Related commands

*read\_data, pair\_style*

## 16.4.6 Default

The default atom style is atomic. If atom\_style sphere is used its default argument is 0.

---

(Grime) Grime and Voth, to appear in J Chem Theory & Computation (2014).

## 16.5 balance command

### 16.5.1 Syntax

```
balance thresh style args ... keyword args ...
```

- thresh = imbalance threshold that must be exceeded to perform a re-balance
- one style/arg pair can be used (or multiple for x,y,z)
- style = *x* or *y* or *z* or *shift* or *rcb*
  - x* args = *uniform* or Px-1 numbers between 0 and 1
    - uniform* = evenly spaced cuts between processors in x dimension
    - numbers = Px-1 ascending values between 0 and 1, Px - # of processors in x<sub>dimension</sub>  
→dimension
    - x* can be specified together with *y* or *z*
  - y* args = *uniform* or Py-1 numbers between 0 and 1
    - uniform* = evenly spaced cuts between processors in y dimension
    - numbers = Py-1 ascending values between 0 and 1, Py - # of processors in y<sub>dimension</sub>  
→dimension
    - y* can be specified together with *x* or *z*
  - z* args = *uniform* or Pz-1 numbers between 0 and 1
    - uniform* = evenly spaced cuts between processors in z dimension
    - numbers = Pz-1 ascending values between 0 and 1, Pz - # of processors in z<sub>dimension</sub>  
→dimension
    - z* can be specified together with *x* or *y*
  - shift* args = dimstr Niter stopthresh
    - dimstr = sequence of letters containing "x" or "y" or "z", each not more than once
    - Niter = # of times to iterate within each dimension of dimstr sequence
    - stopthresh = stop balancing when this imbalance threshold is reached
  - rcb* args = none
- zero or more keyword/arg pairs may be appended
- keyword = *weight* or *out*
  - weight* style args = use weighted particle counts for the balancing
    - style = *group* or *neigh* or *time* or *var* or *store*
    - group args = Ngroup group1 weight1 group2 weight2 ...
      - Ngroup = number of groups with assigned weights
      - group1, group2, ... = group IDs
      - weight1, weight2, ... = corresponding weight factors

```

    neigh factor = compute weight based on number of neighbors
                    factor = scaling factor (> 0)
    time factor = compute weight based on time spend computing
                    factor = scaling factor (> 0)
    var name = take weight from atom-style variable
                    name = name of the atom-style variable
    store name = store weight in custom atom property defined by fix property/atom
↪command
    name = atom property name (without d_ prefix)
out arg = filename
    filename = write each processor's sub-domain to a file

```

## 16.5.2 Examples

```

balance 0.9 x uniform y 0.4 0.5 0.6
balance 1.2 shift xz 5 1.1
balance 1.0 shift xz 5 1.1
balance 1.1 rcb
balance 1.0 shift x 10 1.1 weight group 2 fast 0.5 slow 2.0
balance 1.0 shift x 10 1.1 weight time 0.8 weight neigh 0.5 weight store balance
balance 1.0 shift x 20 1.0 out tmp.balance

```

## 16.5.3 Description

This command adjusts the size and shape of processor sub-domains within the simulation box, to attempt to balance the number of atoms or particles and thus indirectly the computational cost (load) more evenly across processors. The load balancing is “static” in the sense that this command performs the balancing once, before or between simulations. The processor sub-domains will then remain static during the subsequent run. To perform “dynamic” balancing, see the *fix balance* command, which can adjust processor sub-domain sizes and shapes on-the-fly during a *run*.

Load-balancing is typically most useful if the particles in the simulation box have a spatially-varying density distribution or when the computational cost varies significantly between different particles. E.g. a model of a vapor/liquid interface, or a solid with an irregular-shaped geometry containing void regions, or *hybrid pair style simulations* which combine pair styles with different computational cost. In these cases, the LAMMPS default of dividing the simulation box volume into a regular-spaced grid of 3d bricks, with one equal-volume sub-domain per processor, may assign numbers of particles per processor in a way that the computational effort varies significantly. This can lead to poor performance when the simulation is run in parallel.

The balancing can be performed with or without per-particle weighting. With no weighting, the balancing attempts to assign an equal number of particles to each processor. With weighting, the balancing attempts to assign an equal aggregate computational weight to each processor, which typically induces a different number of atoms assigned to each processor. Details on the various weighting options and examples for how they can be used are *given below*.

Note that the *processors* command allows some control over how the box volume is split across processors. Specifically, for a  $P_x$  by  $P_y$  by  $P_z$  grid of processors, it allows choice of  $P_x$ ,  $P_y$ , and  $P_z$ , subject to the constraint that  $P_x * P_y * P_z = P$ , the total number of processors. This is sufficient to achieve good load-balance for some problems on some processor counts. However, all the processor sub-domains will still have the same shape and same volume.

The requested load-balancing operation is only performed if the current “imbalance factor” in particles owned by each processor exceeds the specified *thresh* parameter. The imbalance factor is defined as the maximum number of particles (or weight) owned by any processor, divided by the average number of particles (or weight) per processor. Thus an imbalance factor of 1.0 is perfect balance.

As an example, for 10000 particles running on 10 processors, if the most heavily loaded processor has 1200 particles, then the factor is 1.2, meaning there is a 20% imbalance. Note that a re-balance can be forced even if the current balance is perfect (1.0) by specifying a *thresh* < 1.0.

---

**Note:** Balancing is performed even if the imbalance factor does not exceed the *thresh* parameter if a “grid” style is specified when the current partitioning is “tiled”. The meaning of “grid” vs “tiled” is explained below. This is to allow forcing of the partitioning to “grid” so that the *comm\_style brick* command can then be used to replace a current *comm\_style tiled* setting.

---

When the balance command completes, it prints statistics about the result, including the change in the imbalance factor and the change in the maximum number of particles on any processor. For “grid” methods (defined below) that create a logical 3d grid of processors, the positions of all cutting planes in each of the 3 dimensions (as fractions of the box length) are also printed.

---

**Note:** This command attempts to minimize the imbalance factor, as defined above. But depending on the method a perfect balance (1.0) may not be achieved. For example, “grid” methods (defined below) that create a logical 3d grid cannot achieve perfect balance for many irregular distributions of particles. Likewise, if a portion of the system is a perfect lattice, e.g. the initial system is generated by the *create\_atoms* command, then “grid” methods may be unable to achieve exact balance. This is because entire lattice planes will be owned or not owned by a single processor.

---

---

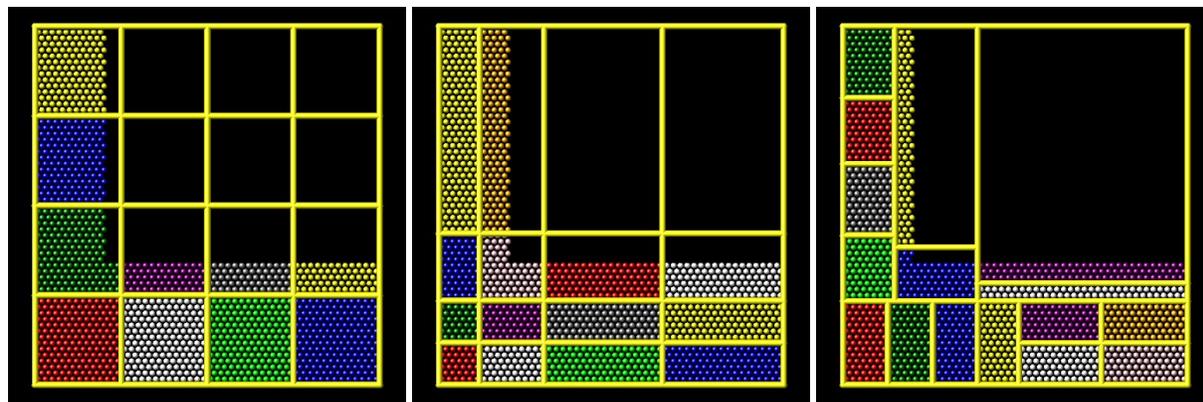
**Note:** The imbalance factor is also an estimate of the maximum speed-up you can hope to achieve by running a perfectly balanced simulation versus an imbalanced one. In the example above, the 10000 particle simulation could run up to 20% faster if it were perfectly balanced, versus when imbalanced. However, computational cost is not strictly proportional to particle count, and changing the relative size and shape of processor sub-domains may lead to additional computational and communication overheads, e.g. in the PPPM solver used via the *kpspace\_style* command. Thus you should benchmark the run times of a simulation before and after balancing.

---

---

The method used to perform a load balance is specified by one of the listed styles (or more in the case of *x,y,z*), which are described in detail below. There are 2 kinds of styles.

The *x*, *y*, *z*, and *shift* styles are “grid” methods which produce a logical 3d grid of processors. They operate by changing the cutting planes (or lines) between processors in 3d (or 2d), to adjust the volume (area in 2d) assigned to each processor, as in the following 2d diagram where processor sub-domains are shown and particles are colored by the processor that owns them.



The leftmost diagram is the default partitioning of the simulation box across processors (one sub-box for each of 16 processors); the middle diagram is after a “grid” method has been applied. The *rcb* style is a “tiling” method which

does not produce a logical 3d grid of processors. Rather it tiles the simulation domain with rectangular sub-boxes of varying size and shape in an irregular fashion so as to have equal numbers of particles (or weight) in each sub-box, as in the rightmost diagram above.

The “grid” methods can be used with either of the *comm\_style* command options, *brick* or *tiled*. The “tiling” methods can only be used with *comm\_style tiled*. Note that it can be useful to use a “grid” method with *comm\_style tiled* to return the domain partitioning to a logical 3d grid of processors so that “comm\_style brick” can afterwards be specified for subsequent *run* commands.

When a “grid” method is specified, the current domain partitioning can be either a logical 3d grid or a tiled partitioning. In the former case, the current logical 3d grid is used as a starting point and changes are made to improve the imbalance factor. In the latter case, the tiled partitioning is discarded and a logical 3d grid is created with uniform spacing in all dimensions. This becomes the starting point for the balancing operation.

When a “tiling” method is specified, the current domain partitioning (“grid” or “tiled”) is ignored, and a new partitioning is computed from scratch.

---

The *x*, *y*, and *z* styles invoke a “grid” method for balancing, as described above. Note that any or all of these 3 styles can be specified together, one after the other, but they cannot be used with any other style. This style adjusts the position of cutting planes between processor sub-domains in specific dimensions. Only the specified dimensions are altered.

The *uniform* argument spaces the planes evenly, as in the left diagrams above. The *numeric* argument requires listing Ps-1 numbers that specify the position of the cutting planes. This requires knowing  $P_s = P_x$  or  $P_y$  or  $P_z =$  the number of processors assigned by LAMMPS to the relevant dimension. This assignment is made (and the  $P_x$ ,  $P_y$ ,  $P_z$  values printed out) when the simulation box is created by the “create\_box” or “read\_data” or “read\_restart” command and is influenced by the settings of the *processors* command.

Each of the numeric values must be between 0 and 1, and they must be listed in ascending order. They represent the fractional position of the cutting place. The left (or lower) edge of the box is 0.0, and the right (or upper) edge is 1.0. Neither of these values is specified. Only the interior Ps-1 positions are specified. Thus if there are 2 processors in the x dimension, you specify a single value such as 0.75, which would make the left processor’s sub-domain 3x larger than the right processor’s sub-domain.

---

The *shift* style invokes a “grid” method for balancing, as described above. It changes the positions of cutting planes between processors in an iterative fashion, seeking to reduce the imbalance factor, similar to how the *fix balance shift* command operates.

The *dimstr* argument is a string of characters, each of which must be an “x” or “y” or “z”. Each character can appear zero or one time, since there is no advantage to balancing on a dimension more than once. You should normally only list dimensions where you expect there to be a density variation in the particles.

Balancing proceeds by adjusting the cutting planes in each of the dimensions listed in *dimstr*, one dimension at a time. For a single dimension, the balancing operation (described below) is iterated on up to *Niter* times. After each dimension finishes, the imbalance factor is re-computed, and the balancing operation halts if the *stoptresh* criterion is met.

A re-balance operation in a single dimension is performed using a recursive multisectioning algorithm, where the position of each cutting plane (line in 2d) in the dimension is adjusted independently. This is similar to a recursive bisectioning for a single value, except that the bounds used for each bisectioning take advantage of information from neighboring cuts if possible. At each iteration, the count of particles on either side of each plane is tallied. If the counts do not match the target value for the plane, the position of the cut is adjusted to be halfway between a low and high bound. The low and high bounds are adjusted on each iteration, using new count information, so that they become closer together over time. Thus as the recursion progresses, the count of particles on either side of the plane gets closer to the target value.

After the balanced plane positions are determined, if any pair of adjacent planes are closer together than the neighbor skin distance (as specified by the *neigh\_modify* command), then the plane positions are shifted to separate them by at

least this amount. This is to prevent particles being lost when dynamics are run with processor sub-domains that are too narrow in one or more dimensions.

Once the re-balancing is complete and final processor sub-domains assigned, particles are migrated to their new owning processor, and the balance procedure ends.

---

**Note:** At each re-balance operation, the bisectioning for each cutting plane (line in 2d) typically starts with low and high bounds separated by the extent of a processor's sub-domain in one dimension. The size of this bracketing region shrinks by 1/2 every iteration. Thus if *Niter* is specified as 10, the cutting plane will typically be positioned to 1 part in 1000 accuracy (relative to the perfect target position). For *Niter* = 20, it will be accurate to 1 part in a million. Thus there is no need to set *Niter* to a large value. LAMMPS will check if the threshold accuracy is reached (in a dimension) is less iterations than *Niter* and exit early. However, *Niter* should also not be set too small, since it will take roughly the same number of iterations to converge even if the cutting plane is initially close to the target value.

---

The *rcb* style invokes a “tiled” method for balancing, as described above. It performs a recursive coordinate bisectioning (RCB) of the simulation domain. The basic idea is as follows.

The simulation domain is cut into 2 boxes by an axis-aligned cut in one of the dimensions, leaving one new sub-box on either side of the cut. Which dimension is chosen for the cut depends on the particle (weight) distribution within the parent box. Normally the longest dimension of the box is cut, but if all (or most) of the particles are at one end of the box, a cut may be performed in another dimension to induce sub-boxes that are more cube-ish (3d) or square-ish (2d) in shape.

After the cut is made, all the processors are also partitioned into 2 groups, half assigned to the box on the lower side of the cut, and half to the box on the upper side. (If the processor count is odd, one side gets an extra processor.) The cut is positioned so that the number of (weighted) particles in the lower box is exactly the number that the processors assigned to that box should own for load balance to be perfect. This also makes load balance for the upper box perfect. The positioning of the cut is done iteratively, by a bisectioning method (median search). Note that counting particles on either side of the cut requires communication between all processors at each iteration.

That is the procedure for the first cut. Subsequent cuts are made recursively, in exactly the same manner. The subset of processors assigned to each box make a new cut in one dimension of that box, splitting the box, the subset of processors, and the particles in the box in two. The recursion continues until every processor is assigned a sub-box of the entire simulation domain, and owns the (weighted) particles in that sub-box.

---

This sub-section describes how to perform weighted load balancing using the *weight* keyword.

By default, all particles have a weight of 1.0, which means each particle is assumed to require the same amount of computation during a timestep. There are, however, scenarios where this is not a good assumption. Measuring the computational cost for each particle accurately would be impractical and slow down the computation. Instead the *weight* keyword implements several ways to influence the per-particle weights empirically by properties readily available or using the user's knowledge of the system. Note that the absolute value of the weights are not important; only their relative ratios affect which particle is assigned to which processor. A particle with a weight of 2.5 is assumed to require 5x more computational than a particle with a weight of 0.5. For all the options below the weight assigned to a particle must be a positive value; an error will be generated if a weight is  $\leq 0.0$ .

Below is a list of possible weight options with a short description of their usage and some example scenarios where they might be applicable. It is possible to apply multiple weight flags and the weightings they induce will be combined through multiplication. Most of the time, however, it is sufficient to use just one method.

The *group* weight style assigns weight factors to specified *groups* of particles. The *group* style keyword is followed by the number of groups, then pairs of group IDs and the corresponding weight factor. If a particle belongs to none of the specified groups, its weight is not changed. If it belongs to multiple groups, its weight is the product of the weight factors.

This weight style is useful in combination with pair style *hybrid*, e.g. when combining a more costly many-body potential with a fast pairwise potential. It is also useful when using *run\_style respa* where some portions of the system have many bonded interactions and others none. It assumes that the computational cost for each group remains constant over time. This is a purely empirical weighting, so a series test runs to tune the assigned weight factors for optimal performance is recommended.

The *neigh* weight style assigns the same weight to each particle owned by a processor based on the total count of neighbors in the neighbor list owned by that processor. The motivation is that more neighbors means a higher computational cost. The style does not use neighbors per atom to assign a unique weight to each atom, because that value can vary depending on how the neighbor list is built.

The *factor* setting is applied as an overall scale factor to the *neigh* weights which allows adjustment of their impact on the balancing operation. The specified *factor* value must be positive. A value  $> 1.0$  will increase the weights so that the ratio of max weight to min weight increases by *factor*. A value  $< 1.0$  will decrease the weights so that the ratio of max weight to min weight decreases by *factor*. In both cases the intermediate weight values increase/decrease proportionally as well. A value  $= 1.0$  has no effect on the *neigh* weights. As a rule of thumb, we have found a *factor* of about 0.8 often results in the best performance, since the number of neighbors is likely to overestimate the ideal weight.

This weight style is useful for systems where there are different cutoffs used for different pairs of interactions, or the density fluctuates, or a large number of particles are in the vicinity of a wall, or a combination of these effects. If a simulation uses multiple neighbor lists, this weight style will use the first suitable neighbor list it finds. It will not request or compute a new list. A warning will be issued if there is no suitable neighbor list available or if it is not current, e.g. if the *balance* command is used before a *run* or *minimize* command is used, in which case the neighbor list may not yet have been built. In this case no weights are computed. Inserting a *run 0 post no* command before issuing the *balance* command, may be a workaround for this case, as it will induce the neighbor list to be built.

The *time* weight style uses *timer data* to estimate weights. It assigns the same weight to each particle owned by a processor based on the total computational time spent by that processor. See details below on what time window is used. It uses the same timing information as is used for the *MPI task timing breakdown*, namely, for sections *Pair*, *Bond*, *Kspace*, and *Neigh*. The time spent in those portions of the timestep are measured for each MPI rank, summed, then divided by the number of particles owned by that processor. I.e. the weight is an effective CPU time/particle averaged over the particles on that processor.

The *factor* setting is applied as an overall scale factor to the *time* weights which allows adjustment of their impact on the balancing operation. The specified *factor* value must be positive. A value  $> 1.0$  will increase the weights so that the ratio of max weight to min weight increases by *factor*. A value  $< 1.0$  will decrease the weights so that the ratio of max weight to min weight decreases by *factor*. In both cases the intermediate weight values increase/decrease proportionally as well. A value  $= 1.0$  has no effect on the *time* weights. As a rule of thumb, effective values to use are typically between 0.5 and 1.2. Note that the timer quantities mentioned above can be affected by communication which occurs in the middle of the operations, e.g. pair styles with intermediate exchange of data within the force computation, and likewise for KSpace solves.

When using the *time* weight style with the *balance* command, the timing data is taken from the preceding run command, i.e. the timings are for the entire previous run. For the *fix balance* command the timing data is for only the timesteps since the last balancing operation was performed. If timing information for the required sections is not available, e.g. at the beginning of a run, or when the *timer* command is set to either *loop* or *off*, a warning is issued. In this case no weights are computed.

---

**Note:** The *time* weight style is the most generic option, and should be tried first, unless the *group* style is easily applicable. However, since the computed cost function is averaged over all particles on a processor, the weights may not be highly accurate. This style can also be effective as a secondary weight in combination with either *group* or *neigh* to offset some of inaccuracies in either of those heuristics.

---

The *var* weight style assigns per-particle weights by evaluating an *atom-style variable* specified by *name*. This is provided as a more flexible alternative to the *group* weight style, allowing definition of a more complex heuristics based

on information (global and per atom) available inside of LAMMPS. For example, atom-style variables can reference the position of a particle, its velocity, the volume of its Voronoi cell, etc.

The *store* weight style does not compute a weight factor. Instead it stores the current accumulated weights in a custom per-atom vector specified by *name*. This must be a vector defined as *d\_name* via the *fix property/atom* command. This means the values in the vector can be read as part of a data file with the *read\_data* command or specified with the *set* command. These weights can also be output in a *dump* file, so this is a way to examine, debug, or visualize the per-particle weights used during the load-balancing operation.

Note that the name of the custom per-atom vector is specified just as *name*, not as *d\_name* as it is for other commands that use different kinds of custom atom vectors or arrays as arguments.

---

The *out* keyword writes a text file to the specified *filename* with the results of the balancing operation. The file contains the bounds of the sub-domain for each processor after the balancing operation completes. The format of the file is compatible with the [Pizza.py mdump](#) tool which has support for manipulating and visualizing mesh files. An example is shown here for a balancing by 4 processors for a 2d problem:

```
ITEM: TIMESTEP
0
ITEM: NUMBER OF NODES
16
ITEM: BOX BOUNDS
0 10
0 10
0 10
ITEM: NODES
1 1 0 0 0
2 1 5 0 0
3 1 5 5 0
4 1 0 5 0
5 1 5 0 0
6 1 10 0 0
7 1 10 5 0
8 1 5 5 0
9 1 0 5 0
10 1 5 5 0
11 1 5 10 0
12 1 10 5 0
13 1 5 5 0
14 1 10 5 0
15 1 10 10 0
16 1 5 10 0
ITEM: TIMESTEP
0
ITEM: NUMBER OF SQUARES
4
ITEM: SQUARES
1 1 1 2 3 4
2 1 5 6 7 8
3 1 9 10 11 12
4 1 13 14 15 16
```

The coordinates of all the vertices are listed in the NODES section, 5 per processor. Note that the 4 sub-domains share vertices, so there will be duplicate nodes in the list.



The “SQUARES” section lists the node IDs of the 4 vertices in a rectangle for each processor (1 to 4).

For a 3d problem, the syntax is similar with 8 vertices listed for each processor, instead of 4, and “SQUARES” replaced by “CUBES”.

## 16.5.4 Restrictions

For 2d simulations, the *z* style cannot be used. Nor can a “z” appear in *dimstr* for the *shift* style.

Balancing through recursive bisectioning (*rcb* style) requires *comm\_style tiled*

## 16.5.5 Related commands

*group*, *processors*, *fix balance*, *comm\_style*

## 16.5.6 Default

none

# 16.6 bond\_coeff command

## 16.6.1 Syntax

```
bond_coeff N args
```

- N = numeric bond type (see asterisk form below), or type label
- args = coefficients for one or more bond types

## 16.6.2 Examples

```
bond_coeff 5 80.0 1.2
bond_coeff * 30.0 1.5 1.0 1.0
bond_coeff 1*4 30.0 1.5 1.0 1.0
bond_coeff 1 harmonic 200.0 1.0 (for bond_style hybrid)
```

```
labelmap bond 5 carbonyl
```

```
bond_coeff carbonyl 80.0 1.2
```

### 16.6.3 Description

Specify the bond force field coefficients for one or more bond types. The number and meaning of the coefficients depends on the bond style. Bond coefficients can also be set in the data file read by the *read\_data* command or in a restart file.

$N$  can be specified in one of several ways. An explicit numeric value can be used, as in the first example above. Or  $N$  can be a type label, which is an alphanumeric string defined by the *labelmap* command or in a section of a data file read by the *read\_data* command.

For numeric values only, a wild-card asterisk can be used to set the coefficients for multiple bond types. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If  $N$  is the number of bond types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $n$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive).

Note that using a *bond\_coeff* command can override a previous setting for the same bond type. For example, these commands set the coeffs for all bond types, then overwrite the coeffs for just bond type 2:

```
bond_coeff * 100.0 1.2
bond_coeff 2 200.0 1.2
```

A line in a data file that specifies bond coefficients uses the exact same format as the arguments of the *bond\_coeff* command in an input script, except that wild-card asterisks should not be used since coefficients for all  $N$  types must be listed in the file. For example, under the “Bond Coeffs” section of a data file, the line that corresponds to the first example above would be listed as

```
5 80.0 1.2
```

---

The list of all bond styles defined in LAMMPS is given on the *bond\_style* doc page. They are also listed in more compact form on the *Commands bond* doc page.

On either of those pages, click on the style to display the formula it computes and its coefficients as specified by the associated *bond\_coeff* command.

---

### 16.6.4 Restrictions

This command must come after the simulation box is defined by a *read\_data*, *read\_restart*, or *create\_box* command.

A bond style must be defined before any bond coefficients are set, either in the input script or in a data file.

### 16.6.5 Related commands

*bond\_style*

## 16.6.6 Default

none

## 16.7 bond\_style command

### 16.7.1 Syntax

```
bond_style style args
```

- style = *none* or *zero* or *hybrid* or *bpm/rotational* or *bpm/spring* or *class2* or *fene* or *fene/expand* or *fene/nm* or *gaussian* or *gromos* or *harmonic* or *harmonic/shift* or *harmonic/shift/cut* or *morse* or *nonlinear* or *oxdna/fene* or *oxdena2/fene* or *oxrna2/fene* or *quartic* or *special* or *table*
- args = none for any style except *hybrid*
  - *hybrid* args = list of one or more styles

### 16.7.2 Examples

```
bond_style harmonic
bond_style fene
bond_style hybrid harmonic fene
```

### 16.7.3 Description

Set the formula(s) LAMMPS uses to compute bond interactions between pairs of atoms. In LAMMPS, a bond differs from a pairwise interaction, which are set via the *pair\_style* command. Bonds are defined between specified pairs of atoms and remain in force for the duration of the simulation (unless the bond breaks which is possible in some bond potentials). The list of bonded atoms is read in by a *read\_data* or *read\_restart* command from a data or restart file. By contrast, pair potentials are typically defined between all pairs of atoms within a cutoff distance and the set of active interactions changes over time.

Hybrid models where bonds are computed using different bond potentials can be setup using the *hybrid* bond style.

The coefficients associated with a bond style can be specified in a data or restart file or via the *bond\_coeff* command.

All bond potentials store their coefficient data in binary restart files which means *bond\_style* and *bond\_coeff* commands do not need to be re-specified in an input script that restarts a simulation. See the *read\_restart* command for details on how to do this. The one exception is that *bond\_style hybrid* only stores the list of sub-styles in the restart file; bond coefficients need to be re-specified.

---

**Note:** When both a bond and pair style is defined, the *special\_bonds* command often needs to be used to turn off (or weight) the pairwise interaction that would otherwise exist between 2 bonded atoms.

---

In the formulas listed for each bond style,  $r$  is the distance between the 2 atoms in the bond.

---

Here is an alphabetic list of bond styles defined in LAMMPS. Click on the style to display the formula it computes and coefficients specified by the associated *bond\_coeff* command.

Click on the style to display the formula it computes, any additional arguments specified in the `bond_style` command, and coefficients specified by the associated `bond_coeff` command.

There are also additional accelerated pair styles included in the LAMMPS distribution for faster performance on CPUs, GPUs, and KNLs. The individual style names on the [Commands bond](#) doc page are followed by one or more of (g,i,k,o,t) to indicate which accelerated styles exist.

- *none* - turn off bonded interactions
- *zero* - topology but no interactions
- *hybrid* - define multiple styles of bond interactions
- *bpm/rotational* - breakable bond with forces and torques based on deviation from reference state
- *bpm/spring* - breakable bond with forces based on deviation from reference length
- *class2* - COMPASS (class 2) bond
- *fene* - FENE (finite-extensible non-linear elastic) bond
- *fene/expand* - FENE bonds with variable size particles
- *fene/nm* - FENE bonds with a generalized Lennard-Jones potential
- *gaussian* - multicentered Gaussian-based bond potential
- *gromos* - GROMOS force field bond
- *harmonic* - harmonic bond
- *harmonic/shift* - shifted harmonic bond
- *harmonic/shift/cut* - shifted harmonic bond with a cutoff
- *mesocnt* - Harmonic bond wrapper with parameterization presets for nanotubes
- *mm3* - MM3 anharmonic bond
- *morse* - Morse bond
- *nonlinear* - nonlinear bond
- *oxdna/fene* - modified FENE bond suitable for DNA modeling
- *oxdna2/fene* - same as oxdna but used with different pair styles
- *oxrna2/fene* - modified FENE bond suitable for RNA modeling
- *quartic* - breakable quartic bond
- *special* - enable special bond exclusions for 1-5 pairs and beyond
- *table* - tabulated by bond length

## 16.7.4 Restrictions

Bond styles can only be set for atom styles that allow bonds to be defined.

Most bond styles are part of the MOLECULE package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info. The doc pages for individual bond potentials tell if it is part of a package.

## 16.7.5 Related commands

*bond\_coeff*, *delete\_bonds*

## 16.7.6 Default

```
bond_style none
```

# 16.8 bond\_write command

## 16.8.1 Syntax

```
bond_write btype N inner outer file keyword itype jtype
```

- btype = bond types
- N = # of values
- inner,outer = inner and outer bond length (distance units)
- file = name of file to write values to
- keyword = section name in file for this set of tabulated values
- itype,jtype = 2 atom types (optional)

## 16.8.2 Examples

```
bond_write 1 500 0.5 3.5 table.txt Harmonic_1
bond_write 3 1000 0.1 6.0 table.txt Morse
```

## 16.8.3 Description

Write energy and force values to a file as a function of distance for the currently defined bond potential. This is useful for plotting the potential function or otherwise debugging its values. If the file already exists, the table of values is appended to the end of the file to allow multiple tables of energy and force to be included in one file.

The energy and force values are computed at distances from inner to outer for 2 interacting atoms forming a bond of type btype, using the appropriate *bond\_coeff* coefficients. N evenly spaced distances are used.

For example, for N = 7, inner = 1.0, and outer = 4.0, values are computed at r = 1.0, 1.5, 2.0, 2.5, 3.0, 3.5, 4.0.

The file is written in the format used as input for the *bond\_style table* option with *keyword* as the section name. Each line written to the file lists an index number (1-N), a distance (in distance units), an energy (in energy units), and a force

(in force units). In case a new file is created, the first line will be a comment with a “DATE:” and “UNITS:” tag with the current date and *units* settings. For subsequent invocations of the `bond_write` command for the same file, data will be appended and the current units settings will be compared to the data from the header, if present, and `bond_write` will refuse to add a table if the units are not the same.

## 16.8.4 Restrictions

All force field coefficients for bond and other kinds of interactions must be set before this command can be invoked.

Due to how the bond force is computed, an inner value  $> 0.0$  must be specified even if the potential has a finite value at  $r = 0.0$ .

## 16.8.5 Related commands

*bond\_style table*, *bond\_style*, *bond\_coeff*

## 16.8.6 Default

none

# 16.9 boundary command

## 16.9.1 Syntax

```
boundary x y z
```

- $x,y,z = p$  or  $s$  or  $f$  or  $m$ , one or two letters
  - $p$  is periodic
  - $f$  is non-periodic and fixed
  - $s$  is non-periodic and shrink-wrapped
  - $m$  is non-periodic and shrink-wrapped with a minimum value

## 16.9.2 Examples

```
boundary p p f
boundary p fs p
boundary s f fm
```

### 16.9.3 Description

Set the style of boundaries for the global simulation box in each dimension. A single letter assigns the same style to both the lower and upper face of the box. Two letters assigns the first style to the lower face and the second style to the upper face. The initial size of the simulation box is set by the *read\_data*, *read\_restart*, or *create\_box* commands.

The style *p* means the box is periodic, so that particles interact across the boundary, and they can exit one end of the box and re-enter the other end. A periodic dimension can change in size due to constant pressure boundary conditions or box deformation (see the *fix npt* and *fix deform* commands). The *p* style must be applied to both faces of a dimension.

The styles *f*, *s*, and *m* mean the box is non-periodic, so that particles do not interact across the boundary and do not move from one side of the box to the other.

For style *f*, the position of the face is fixed. If an atom moves outside the face it will be deleted on the next timestep that reneighboring occurs. This will typically generate an error unless you have set the *thermo\_modify lost* option to allow for lost atoms.

For style *s*, the position of the face is set so as to encompass the atoms in that dimension (shrink-wrapping), no matter how far they move. Note that when the difference between the current box dimensions and the shrink-wrap box dimensions is large, this can lead to lost atoms at the beginning of a run when running in parallel. This is due to the large change in the (global) box dimensions also causing significant changes in the individual sub-domain sizes. If these changes are farther than the communication cutoff, atoms will be lost. This is best addressed by setting initial box dimensions to match the shrink-wrapped dimensions more closely, by using *m* style boundaries (see below).

For style *m*, shrink-wrapping occurs, but is bounded by the value specified in the data or restart file or set by the *create\_box* command. For example, if the upper *z* face has a value of 50.0 in the data file, the face will always be positioned at 50.0 or above, even if the maximum *z*-extent of all the atoms becomes less than 50.0. This can be useful if you start a simulation with an empty box or if you wish to leave room on one side of the box, e.g. for atoms to evaporate from a surface.

For triclinic (non-orthogonal) simulation boxes, if the second dimension of a tilt factor (e.g. *y* for *xy*) is periodic, then the periodicity is enforced with the tilt factor offset. If the first dimension is shrink-wrapped, then the shrink wrapping is applied to the tilted box face, to encompass the atoms. E.g. for a positive *xy* tilt, the *xlo* and *xhi* faces of the box are planes tilting in the +*y* direction as *y* increases. These tilted planes are shrink-wrapped around the atoms to determine the *x* extent of the box.

See the [Howto triclinic](#) page for a geometric description of triclinic boxes, as defined by LAMMPS, and how to transform these parameters to and from other commonly used triclinic representations.

## 16.9.4 Restrictions

This command cannot be used after the simulation box is defined by a *read\_data* or *create\_box* command or *read\_restart* command. See the *change\_box* command for how to change the simulation box boundaries after it has been defined.

For 2d simulations, the z dimension must be periodic.

## 16.9.5 Related commands

See the *thermo\_modify* command for a discussion of lost atoms.

## 16.9.6 Default

```
boundary p p p
```

# 16.10 box command

## 16.10.1 Syntax

```
box keyword value ...
```

- one or more keyword/value pairs may be appended
- keyword = *tilt*  
*tilt* value = *small* or *large*

## 16.10.2 Examples

```
box tilt large  
box tilt small
```

## 16.10.3 Description

Set attributes of the simulation box.

For triclinic (non-orthogonal) simulation boxes, the *tilt* keyword allows simulation domains to be created with arbitrary tilt factors, e.g. via the *create\_box* or *read\_data* commands. Tilt factors determine how skewed the triclinic box is; see the *Howto triclinic* page for a discussion of triclinic boxes in LAMMPS.

LAMMPS normally requires that no tilt factor can skew the box more than half the distance of the parallel box length, which is the first dimension in the tilt factor (x for xz). If *tilt* is set to *small*, which is the default, then an error will be generated if a box is created which exceeds this limit. If *tilt* is set to *large*, then no limit is enforced. You can create a box with any tilt factors you wish.

Note that if a simulation box has a large tilt factor, LAMMPS will run less efficiently, due to the large volume of communication needed to acquire ghost atoms around a processor's irregular-shaped sub-domain. For extreme values of tilt, LAMMPS may also lose atoms and generate an error.



## 16.10.4 Restrictions

This command cannot be used after the simulation box is defined by a *read\_data* or *create\_box* command or *read\_restart* command.

## 16.10.5 Related commands

none

## 16.10.6 Default

The default value is *tilt = small*.

# 16.11 change\_box command

## 16.11.1 Syntax

```
change_box group-ID parameter args ... keyword args ...
```

- group-ID = ID of group of atoms to (optionally) displace
- one or more parameter/arg pairs may be appended

parameter = *x* or *y* or *z* or *xy* or *xz* or *yz* or *boundary* or *ortho* or *triclinic* or *set*  
 ↪ or *remap*

*x*, *y*, *z* args = style value(s)

style = *final* or *delta* or *scale* or *volume*

*final* values = lo hi

lo hi = box boundaries after displacement (distance units)

*delta* values = dlo dhi

dlo dhi = change in box boundaries after displacement (distance units)

*scale* values = factor

factor = multiplicative factor for change in box length after displacement

*volume* value = none = adjust this dim to preserve volume of system

*xy*, *xz*, *yz* args = style value

style = *final* or *delta*

*final* value = tilt

tilt = tilt factor after displacement (distance units)

*delta* value = dtilt

dtilt = change in tilt factor after displacement (distance units)

*boundary* args = *x y z*

*x,y,z* = *p* or *s* or *f* or *m*, one or two letters

*p* is periodic

*f* is non-periodic and fixed

*s* is non-periodic and shrink-wrapped

*m* is non-periodic and shrink-wrapped with a minimum value

*ortho* args = none = change box to orthogonal

*triclinic* args = none = change box to triclinic

*set* args = none = store state of current box

*remap* args = none = remap atom coords from last saved state to current box

- zero or more keyword/value pairs may be appended
- keyword = *units*

```
units value = lattice or box
  lattice = distances are defined in lattice units
  box = distances are defined in simulation box units
```

### 16.11.2 Examples

```
change_box all xy final -2.0 z final 0.0 5.0 boundary p p f remap units box
change_box all x scale 1.1 y volume z volume remap
```

### 16.11.3 Description

Change the volume and/or shape and/or boundary conditions for the simulation box. Orthogonal simulation boxes have 3 adjustable size parameters (x,y,z). Triclinic (non-orthogonal) simulation boxes have 6 adjustable size/shape parameters (x,y,z,xy,xz,yz). Any or all of them can be adjusted independently by this command. Thus it can be used to expand or contract a box, or to apply a shear strain to a non-orthogonal box. It can also be used to change the boundary conditions for the simulation box, similar to the *boundary* command.

The size and shape of the initial simulation box are specified by the *create\_box* or *read\_data* or *read\_restart* command used to setup the simulation. The size and shape may be altered by subsequent runs, e.g. by use of the *fix npt* or *fix deform* commands. The *create\_box*, *read\_data*, and *read\_restart* commands also determine whether the simulation box is orthogonal or triclinic and their doc pages explain the meaning of the xy,xz,yz tilt factors.

See the *Howto triclinic* page for a geometric description of triclinic boxes, as defined by LAMMPS, and how to transform these parameters to and from other commonly used triclinic representations.

The keywords used in this command are applied sequentially to the simulation box and the atoms in it, in the order specified.

Before the sequence of keywords are invoked, the current box size/shape is stored, in case a *remap* keyword is used to map the atom coordinates from a previously stored box size/shape to the current one.

After all the keywords have been processed, any shrink-wrap boundary conditions are invoked (see the *boundary* command) which may change simulation box boundaries, and atoms are migrated to new owning processors.

---

**Note:** This means that you cannot use the *change\_box* command to enlarge a shrink-wrapped box, e.g. to make room to insert more atoms via the *create\_atoms* command, because the simulation box will be re-shrink-wrapped before the *change\_box* command completes. Instead you could do something like this, assuming the simulation box is non-periodic and atoms extend from 0 to 20 in all dimensions:

---

```
change_box all x final -10 20
create_atoms 1 single -5 5 5      # this will fail to insert an atom

change_box all x final -10 20 boundary f s s
create_atoms 1 single -5 5 5
change_box all boundary s s s    # this will work
```

---

**Note:** Unlike the earlier “displace\_box” version of this command, atom remapping is NOT performed by default. This command allows remapping to be done in a more general way, exactly when you specify it (zero or more times) in the

sequence of transformations. Thus if you do not use the *remap* keyword, atom coordinates will not be changed even if the box size/shape changes. If a uniformly strained state is desired, the *remap* keyword should be specified.

---

**Note:** It is possible to lose atoms with this command. E.g. by changing the box without remapping the atoms, and having atoms end up outside of non-periodic boundaries. It is also possible to alter bonds between atoms straddling a boundary in bad ways. E.g. by converting a boundary from periodic to non-periodic. It is also possible when remapping atoms to put them (nearly) on top of each other. E.g. by converting a boundary from non-periodic to periodic. All of these will typically lead to bad dynamics and/or generate error messages.

---

**Note:** The simulation box size/shape can be changed by arbitrarily large amounts by this command. This is not a problem, except that the mapping of processors to the simulation box is not changed from its initial 3d configuration; see the *processors* command. Thus, if the box size/shape changes dramatically, the mapping of processors to the simulation box may not end up as optimal as the initial mapping attempted to be. You may wish to re-balance the atoms by using the *balance* command if that is the case.

---

**Note:** You cannot use this command after reading a restart file (and before a run is performed) if the restart file stored per-atom information from a fix and any of the specified keywords change the box size or shape or boundary conditions. This is because atoms may be moved to new processors and the restart info will not migrate with them. LAMMPS will generate an error if this could happen. Only the *ortho* and *triclinic* keywords do not trigger this error. One solution is to perform a “run 0” command before using the *change\_box* command. This clears the per-atom restart data, whether it has been re-assigned to a new fix or not.

---

**Note:** Because the keywords used in this command are applied one at a time to the simulation box and the atoms in it, care must be taken with triclinic cells to avoid exceeding the limits on skew after each transformation in the sequence. If skew is exceeded before the final transformation this can be avoided by changing the order of the sequence, or breaking the transformation into two or more smaller transformations. For more information on the allowed limits for box skew see the discussion on triclinic boxes on [Howto triclinic](#) doc page.

---

For the *x*, *y*, and *z* parameters, this is the meaning of their styles and values.

For style *final*, the final lo and hi box boundaries of a dimension are specified. The values can be in lattice or box distance units. See the discussion of the units keyword below.

For style *delta*, plus or minus changes in the lo/hi box boundaries of a dimension are specified. The values can be in lattice or box distance units. See the discussion of the units keyword below.

For style *scale*, a multiplicative factor to apply to the box length of a dimension is specified. For example, if the initial box length is 10, and the factor is 1.1, then the final box length will be 11. A factor less than 1.0 means compression.

The *volume* style changes the specified dimension in such a way that the overall box volume remains constant with respect to the operation performed by the preceding keyword. The *volume* style can only be used following a keyword that changed the volume, which is any of the *x*, *y*, *z* keywords. If the preceding keyword “key” had a *volume* style, then both it and the current keyword apply to the keyword preceding “key”. I.e. this sequence of keywords is allowed:

```
change_box all x scale 1.1 y volume z volume
```

The *volume* style changes the associated dimension so that the overall box volume is unchanged relative to its value before the preceding keyword was invoked.

If the following command is used, then the z box length will shrink by the same 1.1 factor the x box length was increased by:

```
change_box all x scale 1.1 z volume
```

If the following command is used, then the y,z box lengths will each shrink by  $\sqrt{1.1}$  to keep the volume constant. In this case, the y,z box lengths shrink so as to keep their relative aspect ratio constant:

```
change_box all x scale 1.1 y volume z volume
```

If the following command is used, then the final box will be a factor of 10% larger in x and y, and a factor of 21% smaller in z, so as to keep the volume constant:

```
change_box all x scale 1.1 z volume y scale 1.1 z volume
```

---

**Note:** For solids or liquids, when one dimension of the box is expanded, it may be physically undesirable to hold the other 2 box lengths constant since that implies a density change. For solids, adjusting the other dimensions via the *volume* style may make physical sense (just as for a liquid), but may not be correct for materials and potentials whose Poisson ratio is not 0.5.

---

For the *scale* and *volume* styles, the box length is expanded or compressed around its mid point.

---

For the *xy*, *xz*, and *yz* parameters, this is the meaning of their styles and values. Note that changing the tilt factors of a triclinic box does not change its volume.

For style *final*, the final tilt factor is specified. The value can be in lattice or box distance units. See the discussion of the *units* keyword below.

For style *delta*, a plus or minus change in the tilt factor is specified. The value can be in lattice or box distance units. See the discussion of the *units* keyword below.

All of these styles change the *xy*, *xz*, *yz* tilt factors. In LAMMPS, tilt factors (*xy*,*xz*,*yz*) for triclinic boxes are required to be no more than half the distance of the parallel box length. For example, if  $x_{lo} = 2$  and  $x_{hi} = 12$ , then the x box length is 10 and the *xy* tilt factor must be between -5 and 5. Similarly, both *xz* and *yz* must be between  $-(x_{hi}-x_{lo})/2$  and  $+(y_{hi}-y_{lo})/2$ . Note that this is not a limitation, since if the maximum tilt factor is 5 (as in this example), then configurations with tilt = ..., -15, -5, 5, 15, 25, ... are all equivalent. Any tilt factor specified by this command must be within these limits.

---

The *boundary* keyword takes arguments that have exactly the same meaning as they do for the *boundary* command. In each dimension, a single letter assigns the same style to both the lower and upper face of the box. Two letters assigns the first style to the lower face and the second style to the upper face.

The style *p* means the box is periodic; the other styles mean non-periodic. For style *f*, the position of the face is fixed. For style *s*, the position of the face is set so as to encompass the atoms in that dimension (shrink-wrapping), no matter how far they move. For style *m*, shrink-wrapping occurs, but is bounded by the current box edge in that dimension, so that the box will become no smaller. See the *boundary* command for more explanation of these style options.

Note that the “boundary” command itself can only be used before the simulation box is defined via a *read\_data* or *create\_box* or *read\_restart* command. This command allows the boundary conditions to be changed later in your input script. Also note that the *read\_restart* will change boundary conditions to match what is stored in the restart file. So if you wish to change them, you should use the *change\_box* command after the *read\_restart* command.

---

**Note:** Changing a periodic boundary to a non-periodic one will also cause the image flag for that dimension of all atoms to be reset to 0. LAMMPS will print a warning message, if that happens. Please note that this reset can lead

to undesired changes when atoms are involved in bonded interactions that straddle periodic boundaries and thus the values of the image flag differs for those atoms.

---

The *ortho* and *triclinic* keywords convert the simulation box to be orthogonal or triclinic (non-orthogonal).

The simulation box is defined as either orthogonal or triclinic when it is created via the *create\_box*, *read\_data*, or *read\_restart* commands.

These keywords allow you to toggle the existing simulation box from orthogonal to triclinic and vice versa. For example, an initial equilibration simulation can be run in an orthogonal box, the box can be toggled to triclinic, and then a *non-equilibrium MD (NEMD) simulation* can be run with deformation via the *fix deform* command.

If the simulation box is currently triclinic and has non-zero tilt in *xy*, *yz*, or *xz*, then it cannot be converted to an orthogonal box.

---

The *set* keyword saves the current box size/shape. This can be useful if you wish to use the *remap* keyword more than once or if you wish it to be applied to an intermediate box size/shape in a sequence of keyword operations. Note that the box size/shape is saved before any of the keywords are processed, i.e. the box size/shape at the time the *create\_box* command is encountered in the input script.

The *remap* keyword remaps atom coordinates from the last saved box size/shape to the current box state. For example, if you stretch the box in the *x* dimension or tilt it in the *xy* plane via the *x* and *xy* keywords, then the *remap* command will dilate or tilt the atoms to conform to the new box size/shape, as if the atoms moved with the box as it deformed.

Note that this operation is performed without regard to periodic boundaries. Also, any shrink-wrapping of non-periodic boundaries (see the *boundary* command) occurs after all keywords, including this one, have been processed.

Only atoms in the specified group are remapped.

---

The *units* keyword determines the meaning of the distance units used to define various arguments. A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for *units = real* or *metal*. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing.

---

## 16.11.4 Restrictions

If you use the *ortho* or *triclinic* keywords, then at the point in the input script when this command is issued, no *dumps* can be active, nor can a *fix deform* be active. This is because these commands test whether the simulation box is orthogonal when they are first issued. Note that these commands can be used in your script before a *change\_box* command is issued, so long as an *undump* or *unfix* command is also used to turn them off.

## 16.11.5 Related commands

*fix deform*, *boundary*

### 16.11.6 Default

The option default is units = lattice.

## 16.12 clear command

### 16.12.1 Syntax

```
clear
```

### 16.12.2 Examples

```
# (commands for 1st simulation)
clear
# (commands for 2nd simulation)
```

### 16.12.3 Description

This command deletes all atoms, restores all settings to their default values, and frees all memory allocated by LAMMPS. Once a clear command has been executed, it is almost as if LAMMPS were starting over, with only the exceptions noted below. This command enables multiple jobs to be run sequentially from one input script.

These settings are not affected by a clear command: the working directory (*shell* command), log file status (*log* command), echo status (*echo* command), and input script variables (*variable* command).

### 16.12.4 Restrictions

none

### 16.12.5 Related commands

none

### 16.12.6 Default

none

## 16.13 comm\_modify command

### 16.13.1 Syntax

```
comm_modify keyword value ...
```

- one or more keyword/value pairs may be appended
- keyword = *mode* or *cutoff* or *cutoff/multi* or *group* or *reduce/multi* or *vel*
  - mode* value = *single*, *multi*, or *multi/old* = communicate atoms within a single or ↵ multiple distances
  - cutoff* value = Rcut (distance units) = communicate atoms from this far away
  - cutoff/multi* collection value
    - collection = atom collection or collection range (supports asterisk notation)
    - value = Rcut (distance units) = communicate atoms for selected types from this ↵ far away
  - reduce/multi* arg = none = reduce number of communicated ghost atoms for multi style
  - cutoff/multi/old* type value
    - type = atom type or type range (supports asterisk notation)
    - value = Rcut (distance units) = communicate atoms for selected types from this ↵ far away
  - group* value = group-ID = only communicate atoms in the group
  - vel* value = *yes* or *no* = do or do not communicate velocity info with ghost atoms

### 16.13.2 Examples

```
comm_modify mode multi reduce/multi
comm_modify mode multi group solvent
comm_modify mode multi cutoff/multi 1 10.0 cutoff/multi 2*4 15.0
comm_modify vel yes
comm_modify mode single cutoff 5.0 vel yes
comm_modify cutoff/multi * 0.0
```

### 16.13.3 Description

This command sets parameters that affect the inter-processor communication of atom information that occurs each timestep as coordinates and other properties are exchanged between neighboring processors and stored as properties of ghost atoms.

---

**Note:** These options apply to the currently defined comm style. When you specify a *comm\_style* or *read\_restart* command, all communication settings are restored to their default or stored values, including those previously reset by a *comm\_modify* command. Thus if your input script specifies a *comm\_style* or *read\_restart* command, you should use the *comm\_modify* command after it.

---

The *mode* keyword determines whether a single or multiple cutoff distances are used to determine which atoms to communicate.

The default mode is *single* which means each processor acquires information for ghost atoms that are within a single distance from its sub-domain. The distance is by default the maximum of the neighbor cutoff across all atom type pairs.

For many systems this is an efficient algorithm, but for systems with widely varying cutoffs for different type pairs, the *multi* or *multi/old* mode can be faster. In *multi*, each atom is assigned to a collection which should correspond to a set of atoms with similar interaction cutoffs. In this case, each atom collection is assigned its own distance cutoff for communication purposes, and fewer atoms will be communicated. In *multi/old*, a similar technique is used but atoms are grouped by atom type. See the *neighbor multi* and *neighbor multi/old* commands for neighbor list construction options that may also be beneficial for simulations of this kind. The *multi* communication mode is only compatible with the *multi* neighbor style. The *multi/old* communication mode is comparable with both the *multi* and *multi/old* neighbor styles.

The *cutoff* keyword allows you to extend the ghost cutoff distance for communication mode *single*, which is the distance from the borders of a processor's sub-domain at which ghost atoms are acquired from other processors. By default the ghost cutoff = neighbor cutoff = pairwise force cutoff + neighbor skin. See the *neighbor* command for more information about the skin distance. If the specified *Rcut* is greater than the neighbor cutoff, then extra ghost atoms will be acquired. If the provided cutoff is smaller, the provided value will be ignored, the ghost cutoff is set to the neighbor cutoff and a warning will be printed. Specifying a cutoff value of 0.0 will reset any previous value to the default. If bonded interactions exist and equilibrium bond length information is available, then also a heuristic based on that bond length is computed. It is used as communication cutoff, if there is no pair style present and no *comm\_modify cutoff* command used. Otherwise a warning is printed, if this bond based estimate is larger than the communication cutoff used.

The *cutoff/multi* option is equivalent to *cutoff*, but applies to communication mode *multi* instead. Since the communication cutoffs are determined per atom collections, a collection specifier is needed and cutoff for one or multiple collections can be extended. Also ranges of collections using the usual asterisk notation can be given. Collections are indexed from 1 to N where N is the total number of collections. Note that the arguments for *cutoff/multi* are parsed right before each simulation to account for potential changes in the number of collections. Custom cutoffs are preserved between runs but if collections are redefined, one may want to re-specify the communication cutoffs. For granular pair styles, the default cutoff is set to the sum of the current maximum atomic radii for each collection. The *cutoff/multi/old* option is similar to *cutoff/multi* except it operates on atom types as opposed to collections.

The *reduce/multi* option applies to *multi* and sets the communication cutoff for a particle equal to the maximum interaction distance between particles in the same collection. This reduces the number of ghost atoms that need to be communicated. This method is only compatible with the *multi* neighbor style and requires a half neighbor list and Newton on. See the *neighbor multi* command for more information.

These are simulation scenarios in which it may be useful or even necessary to set a ghost cutoff > neighbor cutoff:

- a single polymer chain with bond interactions, but no pairwise interactions
- bonded interactions (e.g. dihedrals) extend further than the pairwise cutoff
- ghost atoms beyond the pairwise cutoff are needed for some computation

In the first scenario, a pairwise potential is not defined. Thus the pairwise neighbor cutoff will be 0.0. But ghost atoms are still needed for computing bond, angle, etc interactions between atoms on different processors, or when the interaction straddles a periodic boundary.

The appropriate ghost cutoff depends on the *newton bond* setting. For *newton bond off*, the distance needs to be the furthest distance between any two atoms in the bond, angle, etc. E.g. the distance between 1-4 atoms in a dihedral. For *newton bond on*, the distance between the central atom in the bond, angle, etc and any other atom is sufficient. E.g. the distance between 2-4 atoms in a dihedral.

In the second scenario, a pairwise potential is defined, but its neighbor cutoff is not sufficiently long enough to enable bond, angle, etc terms to be computed. As in the previous scenario, an appropriate ghost cutoff should be set.

In the last scenario, a *fix* or *compute* or *pairwise potential* needs to calculate with ghost atoms beyond the normal pairwise cutoff for some computation it performs (e.g. locate neighbors of ghost atoms in a manybody pair potential). Setting the ghost cutoff appropriately can insure it will find the needed atoms.

---

**Note:** In these scenarios, if you do not set the ghost cutoff long enough, and if there is only one processor in a periodic dimension (e.g. you are running in serial), then LAMMPS may “find” the atom it is looking for (e.g. the partner atom



in a bond), that is on the far side of the simulation box, across a periodic boundary. This will typically lead to bad dynamics (i.e. the bond length is now the simulation box length). To detect if this is happening, see the *neigh\_modify cluster* command.

---

The *group* keyword will limit communication to atoms in the specified group. This can be useful for models where no ghost atoms are needed for some kinds of particles. All atoms (not just those in the specified group) will still migrate to new processors as they move. The group specified with this option must also be specified via the *atom\_modify first* command.

The *vel* keyword enables velocity information to be communicated with ghost particles. Depending on the *atom\_style*, velocity info includes the translational velocity, angular velocity, and angular momentum of a particle. If the *vel* option is set to *yes*, then ghost atoms store these quantities; if *no* then they do not. The *yes* setting is needed by some pair styles which require the velocity state of both the I and J particles to compute a pairwise I,J interaction, as well as by some compute and fix commands.

Note that if the *fix deform* command is being used with its “remap v” option enabled, then the velocities for ghost atoms (in the fix deform group) mirrored across a periodic boundary will also include components due to any velocity shift that occurs across that boundary (e.g. due to dilation or shear).

### 16.13.4 Restrictions

Communication mode *multi* is currently only available for *comm\_style brick*.

### 16.13.5 Related commands

*comm\_style*, *neighbor*

### 16.13.6 Default

The option defaults are mode = single, group = all, cutoff = 0.0, vel = no. The cutoff default of 0.0 means that ghost cutoff = neighbor cutoff = pairwise force cutoff + neighbor skin.

## 16.14 comm\_style command

### 16.14.1 Syntax

<code>comm_style style</code>
-------------------------------

- style = *brick* or *tiled*

## 16.14.2 Examples

```
comm_style brick
comm_style tiled
```

## 16.14.3 Description

This command sets the style of inter-processor communication of atom information that occurs each timestep as coordinates and other properties are exchanged between neighboring processors and stored as properties of ghost atoms.

For the default *brick* style, the domain decomposition used by LAMMPS to partition the simulation box must be a regular 3d grid of bricks, one per processor. Each processor communicates with its 6 Cartesian neighbors in the grid to acquire information for nearby atoms.

For the *tiled* style, a more general domain decomposition can be used, as triggered by the *balance* or *fix balance* commands. The simulation box can be partitioned into non-overlapping rectangular-shaped “tiles” or varying sizes and shapes. Again there is one tile per processor. To acquire information for nearby atoms, communication must now be done with a more complex pattern of neighboring processors.

Note that this command does not actually define a partitioning of the simulation box (a domain decomposition), rather it determines what kinds of decompositions are allowed and the pattern of communication used to enable the decomposition. A decomposition is created when the simulation box is first created, via the *create\_box* or *read\_data* or *read\_restart* commands. For both the *brick* and *tiled* styles, the initial decomposition will be the same, as described by *create\_box* and *processors* commands. The decomposition can be changed via the *balance* or *fix balance* commands.

## 16.14.4 Restrictions

None

## 16.14.5 Related commands

*comm\_modify*, *processors*, *balance*, *fix balance*

## 16.14.6 Default

The default style is brick.

# 16.15 compute command

## 16.15.1 Syntax

```
compute ID group-ID style args
```

- ID = user-assigned name for the computation
- group-ID = ID of the group of atoms to perform the computation on
- style = one of a list of possible style names (see below)
- args = arguments used by a particular style

## 16.15.2 Examples

```
compute 1 all temp
compute newtemp flow temp/partial 1 1 0
compute 3 all ke/atom
```

## 16.15.3 Description

Define a computation that will be performed on a group of atoms. Quantities calculated by a compute are instantaneous values, meaning they are calculated from information about atoms on the current timestep or iteration, though a compute may internally store some information about a previous state of the system. Defining a compute does not perform a computation. Instead computes are invoked by other LAMMPS commands as needed (e.g., to calculate a temperature needed for a thermostat fix or to generate thermodynamic or dump file output). See the *Howto output* page for a summary of various LAMMPS output options, many of which involve computes.

The ID of a compute can only contain alphanumeric characters and underscores.

Computes calculate one of three styles of quantities: global, per-atom, or local. A global quantity is one or more system-wide values (e.g., the temperature of the system). A per-atom quantity is one or more values per atom (e.g., the kinetic energy of each atom). Per-atom values are set to 0.0 for atoms not in the specified compute group. Local quantities are calculated by each processor based on the atoms it owns, but there may be zero or more per atom (e.g., a list of bond distances). Computes that produce per-atom quantities have the word “atom” in their style (e.g., *ke/atom*). Computes that produce local quantities have the word “local” in their style (e.g., *bond/local*). Styles with neither “atom” or “local” in their style produce global quantities.

Note that a single compute can produce either global or per-atom or local quantities, but not both global and per-atom. It can produce local quantities in tandem with global or per-atom quantities. The compute page will explain.

Global, per-atom, and local quantities each come in three kinds: a single scalar value, a vector of values, or a 2d array of values. The doc page for each compute describes the style and kind of values it produces (e.g., a per-atom vector). Some computes produce more than one kind of a single style (e.g., a global scalar and a global vector).

When a compute quantity is accessed, as in many of the output commands discussed below, it can be referenced via the following bracket notation, where ID is the ID of the compute:

c_ID	entire scalar, vector, or array
c_ID[I]	one element of vector, one column of array
c_ID[I][J]	one element of array

In other words, using one bracket reduces the dimension of the quantity once (vector  $\rightarrow$  scalar, array  $\rightarrow$  vector). Using two brackets reduces the dimension twice (array  $\rightarrow$  scalar). Thus a command that uses scalar compute values as input can also process elements of a vector or array.

Note that commands and *variables* which use compute quantities typically do not allow for all kinds (e.g., a command may require a vector of values, not a scalar). This means there is no ambiguity about referring to a compute quantity as c\_ID even if it produces, for example, both a scalar and vector. The doc pages for various commands explain the details.

In LAMMPS, the values generated by a compute can be used in several ways:

- The results of computes that calculate a global temperature or pressure can be used by fixes that do thermostating or barostating or when atom velocities are created.

- Global values can be output via the *thermo\_style custom* or *fix ave/time* command. Or the values can be referenced in a *variable equal* or *variable atom* command.
- Per-atom values can be output via the *dump custom* command. Or they can be time-averaged via the *fix ave/atom* command or reduced by the *compute reduce* command. Or the per-atom values can be referenced in an *atom-style variable*.
- Local values can be reduced by the *compute reduce* command, or histogrammed by the *fix ave/histo* command, or output by the *dump local* command.

The results of computes that calculate global quantities can be either “intensive” or “extensive” values. Intensive means the value is independent of the number of atoms in the simulation (e.g., temperature). Extensive means the value scales with the number of atoms in the simulation (e.g., total rotational kinetic energy). *Thermodynamic output* will normalize extensive values by the number of atoms in the system, depending on the “thermo\_modify norm” setting. It will not normalize intensive values. If a compute value is accessed in another way (e.g., by a *variable*), you may want to know whether it is an intensive or extensive value. See the page for individual computes for further info.

---

LAMMPS creates its own computes internally for thermodynamic output. Three computes are always created, named “thermo\_temp”, “thermo\_press”, and “thermo\_pe”, as if these commands had been invoked in the input script:

```
compute thermo_temp all temp
compute thermo_press all pressure thermo_temp
compute thermo_pe all pe
```

Additional computes for other quantities are created if the thermo style requires it. See the documentation for the *thermo\_style* command.

Fixes that calculate temperature or pressure, i.e. for thermostatting or barostatting, may also create computes. These are discussed in the documentation for specific *fix* commands.

In all these cases, the default computes LAMMPS creates can be replaced by computes defined by the user in the input script, as described by the *thermo\_modify* and *fix modify* commands.

Properties of either a default or user-defined compute can be modified via the *compute\_modify* command.

Computes can be deleted with the *uncompute* command.

Code for new computes can be added to LAMMPS; see the *Modify* page for details. The results of their calculations accessed in the various ways described above.

---

Each compute style has its own page which describes its arguments and what it does. Here is an alphabetic list of compute styles available in LAMMPS. They are also listed in more compact form on the *Commands compute* doc page.

There are also additional accelerated compute styles included in the LAMMPS distribution for faster performance on CPUs, GPUs, and KNLs. The individual style names on the *Commands compute* page are followed by one or more of (g,i,k,o,t) to indicate which accelerated styles exist.

- *ackland/atom* - determines the local lattice structure based on the Ackland formulation
- *adf* - angular distribution function of triples of atoms
- *aggregate/atom* - aggregate ID for each atom
- *angle* - energy of each angle sub-style
- *angle/local* - theta and energy of each angle
- *angmom/chunk* - angular momentum for each chunk

- *ave/sphere/atom* - compute local density and temperature around each atom
- *basal/atom* - calculates the hexagonal close-packed “c” lattice vector of each atom
- *body/local* - attributes of body sub-particles
- *bond* - energy of each bond sub-style
- *bond/local* - distance and energy of each bond
- *born/matrix* - second derivative of potential with respect to strain
- *centro/atom* - centro-symmetry parameter for each atom
- *centroid/stress/atom* - centroid based stress tensor for each atom
- *chunk/atom* - assign chunk IDs to each atom
- *chunk/spread/atom* - spreads chunk values to each atom in chunk
- *cluster/atom* - cluster ID for each atom
- *cna/atom* - common neighbor analysis (CNA) for each atom
- *cnp/atom* - common neighborhood parameter (CNP) for each atom
- *com* - center of mass of group of atoms
- *com/chunk* - center of mass for each chunk
- *contact/atom* - contact count for each spherical particle
- *coord/atom* - coordination number for each atom
- *damage/atom* - Peridynamic damage for each atom
- *dihedral* - energy of each dihedral sub-style
- *dihedral/local* - angle of each dihedral
- *dilatation/atom* - Peridynamic dilatation for each atom
- *dipole* - dipole vector and total dipole
- *dipole/chunk* - dipole vector and total dipole for each chunk
- *displace/atom* - displacement of each atom
- *dpd* - total values of internal conductive energy, internal mechanical energy, chemical energy, and harmonic average of internal temperature
- *dpd/atom* - per-particle values of internal conductive energy, internal mechanical energy, chemical energy, and internal temperature
- *edpd/temp/atom* - per-atom temperature for each eDPD particle in a group
- *efield/atom* - electric field at each atom
- *entropy/atom* - pair entropy fingerprint of each atom
- *erotate/asphere* - rotational energy of aspherical particles
- *erotate/rigid* - rotational energy of rigid bodies
- *erotate/sphere* - rotational energy of spherical particles
- *erotate/sphere/atom* - rotational energy for each spherical particle
- *event/displace* - detect event on atom displacement
- *fabric* - calculates fabric tensors from pair interactions

- *fep* - compute free energies for alchemical transformation from perturbation theory
- *fep/ta* - compute free energies for a test area perturbation
- *force/tally* - force between two groups of atoms via the tally callback mechanism
- *fragment/atom* - fragment ID for each atom
- *global/atom* - assign global values to each atom from arrays of global values
- *group/group* - energy/force between two groups of atoms
- *gyration* - radius of gyration of group of atoms
- *gyration/chunk* - radius of gyration for each chunk
- *gyration/shape* - shape parameters from gyration tensor
- *gyration/shape/chunk* - shape parameters from gyration tensor for each chunk
- *heat/flux* - heat flux through a group of atoms
- *heat/flux/tally* - heat flux through a group of atoms via the tally callback mechanism
- *heat/flux/virial/tally* - virial heat flux between two groups via the tally callback mechanism
- *hexorder/atom* - bond orientational order parameter  $q_6$
- *hma* - harmonically mapped averaging for atomic crystals
- *improper* - energy of each improper sub-style
- *improper/local* - angle of each improper
- *inertia/chunk* - inertia tensor for each chunk
- *ke* - translational kinetic energy
- *ke/atom* - kinetic energy for each atom
- *ke/atom/eff* - per-atom translational and radial kinetic energy in the electron force field model
- *ke/eff* - kinetic energy of a group of nuclei and electrons in the electron force field model
- *ke/rigid* - translational kinetic energy of rigid bodies
- *mliap* - gradients of energy and forces with respect to model parameters and related quantities for training machine learning interatomic potentials
- *momentum* - translational momentum
- *msd* - mean-squared displacement of group of atoms
- *msd/chunk* - mean-squared displacement for each chunk
- *msd/nongauss* - MSD and non-Gaussian parameter of group of atoms
- *nbond/atom* - calculates number of bonds per atom
- *omega/chunk* - angular velocity for each chunk
- *orientorder/atom* - Steinhardt bond orientational order parameters  $Q_l$
- *pair* - values computed by a pair style
- *pair/local* - distance/energy/force of each pairwise interaction
- *pe* - potential energy
- *pe/atom* - potential energy for each atom

- *mesont* - Nanotube bending, stretching, and intertube energies
- *pe/mol/tally* - potential energy between two groups of atoms separated into intermolecular and intramolecular components via the tally callback mechanism
- *pe/tally* - potential energy between two groups of atoms via the tally callback mechanism
- *plasticity/atom* - Peridynamic plasticity for each atom
- *pressure* - total pressure and pressure tensor
- *pressure/uef* - pressure tensor in the reference frame of an applied flow field
- *property/atom* - convert atom attributes to per-atom vectors/arrays
- *property/chunk* - extract various per-chunk attributes
- *property/local* - convert local attributes to localvectors/arrays
- *ptm/atom* - determines the local lattice structure based on the Polyhedral Template Matching method
- *rdf* - radial distribution function  $g(r)$  histogram of group of atoms
- *reduce* - combine per-atom quantities into a single global value
- *reduce/chunk* - reduce per-atom quantities within each chunk
- *reduce/region* - same as compute reduce, within a region
- *rigid/local* - extract rigid body attributes
- *saed* - electron diffraction intensity on a mesh of reciprocal lattice nodes
- *slice* - extract values from global vector or array
- *smd/contact/radius* - contact radius for Smooth Mach Dynamics
- *smd/damage* - damage status of SPH particles in Smooth Mach Dynamics
- *smd/hourglass/error* - error associated with approximated relative separation in Smooth Mach Dynamics
- *smd/internal/energy* - per-particle enthalpy in Smooth Mach Dynamics
- *smd/plastic/strain* - equivalent plastic strain per particle in Smooth Mach Dynamics
- *smd/plastic/strain/rate* - time rate of the equivalent plastic strain in Smooth Mach Dynamics
- *smd/rho* - per-particle mass density in Smooth Mach Dynamics
- *smd/tlsph/defgrad* - deformation gradient in Smooth Mach Dynamics
- *smd/tlsph/dt* - CFL-stable time increment per particle in Smooth Mach Dynamics
- *smd/tlsph/num/neighs* - number of particles inside the smoothing kernel radius for Smooth Mach Dynamics
- *smd/tlsph/shape* - current shape of the volume of a particle for Smooth Mach Dynamics
- *smd/tlsph/strain* - Green-Lagrange strain tensor for Smooth Mach Dynamics
- *smd/tlsph/strain/rate* - rate of strain for Smooth Mach Dynamics
- *smd/tlsph/stress* - per-particle Cauchy stress tensor for SPH particles
- *smd/triangle/vertices* - coordinates of vertices corresponding to the triangle elements of a mesh for Smooth Mach Dynamics
- *smd/ulsph/effm* - per-particle effective shear modulus
- *smd/ulsph/num/neighs* - number of neighbor particles inside the smoothing kernel radius for Smooth Mach Dynamics

- *smd/ulsph/strain* - logarithmic strain tensor for Smooth Mach Dynamics
- *smd/ulsph/strain/rate* - logarithmic strain rate for Smooth Mach Dynamics
- *smd/ulsph/stress* - per-particle Cauchy stress tensor and von Mises equivalent stress in Smooth Mach Dynamics
- *smd/vol* - per-particle volumes and their sum in Smooth Mach Dynamics
- *snap* - gradients of SNAP energy and forces with respect to linear coefficients and related quantities for fitting SNAP potentials
- *sna/atom* - bispectrum components for each atom
- *sna/grid* - global array of bispectrum components on a regular grid
- *sna/grid/local* - local array of bispectrum components on a regular grid
- *snad/atom* - derivative of bispectrum components for each atom
- *snav/atom* - virial contribution from bispectrum components for each atom
- *sph/e/atom* - per-atom internal energy of Smooth-Particle Hydrodynamics atoms
- *sph/rho/atom* - per-atom density of Smooth-Particle Hydrodynamics atoms
- *sph/t/atom* - per-atom internal temperature of Smooth-Particle Hydrodynamics atoms
- *spin* - magnetic quantities for a system of atoms having spins
- *stress/atom* - stress tensor for each atom
- *stress/cartesian* - stress tensor in cartesian coordinates
- *stress/cylinder* - stress tensor in cylindrical coordinates
- *stress/mop* - normal components of the local stress tensor using the method of planes
- *stress/mop/profile* - profile of the normal components of the local stress tensor using the method of planes
- *stress/spherical* - stress tensor in spherical coordinates
- *stress/tally* - stress between two groups of atoms via the tally callback mechanism
- *tdpd/cc/atom* - per-atom chemical concentration of a specified species for each tDPD particle
- *temp* - temperature of group of atoms
- *temp/asphere* - temperature of aspherical particles
- *temp/body* - temperature of body particles
- *temp/chunk* - temperature of each chunk
- *temp/com* - temperature after subtracting center-of-mass velocity
- *temp/cs* - temperature based on the center-of-mass velocity of atom pairs that are bonded to each other
- *temp/deform* - temperature excluding box deformation velocity
- *temp/deform/eff* - temperature excluding box deformation velocity in the electron force field model
- *temp/drude* - temperature of Core–Drude pairs
- *temp/eff* - temperature of a group of nuclei and electrons in the electron force field model
- *temp/partial* - temperature excluding one or more dimensions of velocity
- *temp/profile* - temperature excluding a binned velocity profile
- *temp/ramp* - temperature excluding ramped velocity component



- *temp/region* - temperature of a region of atoms
- *temp/region/eff* - temperature of a region of nuclei and electrons in the electron force field model
- *temp/rotate* - temperature of a group of atoms after subtracting out their center-of-mass and angular velocities
- *temp/sphere* - temperature of spherical particles
- *temp/uef* - kinetic energy tensor in the reference frame of an applied flow field
- *ti* - thermodynamic integration free energy values
- *torque/chunk* - torque applied on each chunk
- *vacf* - velocity auto-correlation function of group of atoms
- *vcm/chunk* - velocity of center-of-mass for each chunk
- *viscosity/cos* - velocity profile under cosine-shaped acceleration
- *voronoi/atom* - Voronoi volume and neighbors for each atom
- *xrd* - X-ray diffraction intensity on a mesh of reciprocal lattice nodes

### 16.15.4 Restrictions

none

### 16.15.5 Related commands

*uncompute*, *compute\_modify*, *fix ave/atom*, *fix ave/time*, *fix ave/histo*

### 16.15.6 Default

none

## 16.16 compute\_modify command

### 16.16.1 Syntax

```
compute_modify compute-ID keyword value ...
```

- compute-ID = ID of the compute to modify
- one or more keyword/value pairs may be listed
- keyword = *extra/dof* or *extra* or *dynamic/dof* or *dynamic*
  - extra/dof* value = N
    - N = # of extra degrees of freedom to subtract
    - extra* syntax is identical to *extra/dof*, will be disabled at some point
  - dynamic/dof* value = *yes* or *no*
    - yes/no* = do or do not re-compute the number of degrees of freedom (DOF)
      - ↪ contributing to the temperature
    - dynamic* syntax is identical to *dynamic/dof*, will be disabled at some point

## 16.16.2 Examples

```
compute_modify myTemp extra/dof 0
compute_modify newtemp dynamic/dof yes extra/dof 600
```

## 16.16.3 Description

Modify one or more parameters of a previously defined compute. Not all compute styles support all parameters.

The *extra/dof* or *extra* keyword refers to how many degrees of freedom are subtracted (typically from  $3N$ ) as a normalizing factor in a temperature computation. Only computes that compute a temperature use this option. The default is 2 or 3 for *2d* or *3d* systems, which is a correction factor for an ensemble of velocities with zero total linear momentum. For compute temp/partial, if one or more velocity components are excluded, the value used for *extra* is scaled accordingly. You can use a negative number for the *extra* parameter if you need to add degrees-of-freedom. See the *compute temp/asphere* command for an example.

The *dynamic/dof* or *dynamic* keyword determines whether the number of atoms  $N$  in the compute group and their associated degrees of freedom (DOF) are re-computed each time a temperature is computed. Only compute styles that calculate a temperature use this option. By default,  $N$  and their DOF are assumed to be constant. If you are adding atoms or molecules to the system (see the *fix pour*, *fix deposit*, and *fix gcmc* commands) or expect atoms or molecules to be lost (e.g., due to exiting the simulation box or via *fix evaporate*), then this option should be used to ensure the temperature is correctly normalized.

---

**Note:** The *extra* and *dynamic* keywords should not be used as they are deprecated (March 2017) and will eventually be disabled. Instead, use the equivalent *extra/dof* and *dynamic/dof* keywords.

---

## 16.16.4 Restrictions

none

## 16.16.5 Related commands

*compute*

## 16.16.6 Default

The option defaults are *extra/dof* = 2 or 3 for 2d or 3d systems and *dynamic/dof* = *no*.

## 16.17 create\_atoms command

### 16.17.1 Syntax

```
create_atoms type style args keyword values ...
```

- type = atom type (1-Ntypes) of atoms to create (offset for molecule creation)
- style = *box* or *region* or *single* or *mesh* or *random*

*box* args = none  
*region* args = region-ID  
     region-ID = particles will only be created if contained in the region  
*single* args = x y z  
     x,y,z = coordinates of a single particle (distance units)  
*mesh* args = STL-file  
     STL-file = file with triangle mesh in STL format  
*random* args = N seed region-ID  
     N = number of particles to create  
     seed = random # seed (positive integer)  
     region-ID = create atoms within this region, use NULL for entire simulation box

- zero or more keyword/value pairs may be appended
- keyword = *mol* or *basis* or *ratio* or *subset* or *remap* or *var* or *set* or *rotate* or *overlap* or *maxtry* or *units*

*mol* values = template-ID seed  
     template-ID = ID of molecule template specified in a separate *molecule* command  
     seed = random # seed (positive integer)  
*basis* values = M itype  
     M = which basis atom  
     itype = atom type (1-N) to assign to this basis atom  
*ratio* values = frac seed  
     frac = fraction of lattice sites (0 to 1) to populate randomly  
     seed = random # seed (positive integer)  
*subset* values = Nsubset seed  
     Nsubset = # of lattice sites to populate randomly  
     seed = random # seed (positive integer)  
*remap* value = yes or no  
*var* value = name = variable name to evaluate for test of atom creation  
*set* values = dim name  
     dim = x or y or z  
     name = name of variable to set with x, y, or z atom position  
*radscale* value = factor  
     factor = scale factor for setting atom radius  
*meshmode* values = mode arg  
     mode = *bisect* or *grand*  
     *bisect* arg = radthresh  
         radthresh = threshold value for *mesh* to determine when to split triangles.  
     ↪(distance units)  
     *grand* arg = density  
         density = minimum number density for atoms place on *mesh* triangles (inverse.  
     ↪distance squared units)  
*rotate* values = theta Rx Ry Rz  
     theta = rotation angle for single molecule (degrees)  
     Rx,Ry,Rz = rotation vector for single molecule  
*overlap* value = Doverlap  
     Doverlap = only insert if at least this distance from all existing atoms  
*maxtry* value = Ntry  
     Ntry = number of attempts to insert a particle before failure  
*units* value = *lattice* or *box*  
     *lattice* = the geometry is defined in lattice units  
     *box* = the geometry is defined in simulation box units

## 16.17.2 Examples

```

create_atoms 1 box
create_atoms 3 region regsphere basis 2 3
create_atoms 3 region regsphere basis 2 3 ratio 0.5 74637
create_atoms 3 single 0 0 5
create_atoms 1 box var v set x xpos set y ypos
create_atoms 2 random 50 12345 NULL overlap 2.0 maxtry 50
create_atoms 1 mesh open_box.stl meshmode grand 0.1 units box
create_atoms 1 mesh funnel.stl meshmode bisect 4.0 units box radscale 0.9

```

## 16.17.3 Description

This command creates atoms (or molecules) within the simulation box, either on a lattice, or a single atom (or molecule), or on a surface defined by a triangulated mesh, or a random collection of atoms (or molecules). It is an alternative to reading in atom coordinates explicitly via a *read\_data* or *read\_restart* command. A simulation box must already exist, which is typically created via the *create\_box* command. Before using this command, a lattice must also be defined using the *lattice* command, unless you specify the *single* style with *units = box* or the *random* style. For the remainder of this doc page, a created atom or molecule is referred to as a “particle”.

If created particles are individual atoms, they are assigned the specified atom *type*, though this can be altered via the *basis* keyword as discussed below. If molecules are being created, the type of each atom in the created molecule is specified in the file read by the *molecule* command, and those values are added to the specified atom *type* (e.g., if *type* = 2 and the file specifies atom types 1, 2, and 3, then each created molecule will have atom types 3, 4, and 5).

For the *box* style, the *create\_atoms* command fills the entire simulation box with particles on the lattice. If your simulation box is periodic, you should insure its size is a multiple of the lattice spacings, to avoid unwanted atom overlaps at the box boundaries. If your box is periodic and a multiple of the lattice spacing in a particular dimension, LAMMPS is careful to put exactly one particle at the boundary (on either side of the box), not zero or two.

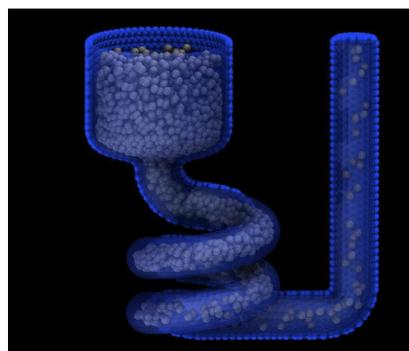
For the *region* style, a geometric volume is filled with particles on the lattice. This volume is what is both inside the simulation box and also consistent with the region volume. See the *region* command for details. Note that a region can be specified so that its “volume” is either inside or outside its geometric boundary. Also note that if a region is the same size as a periodic simulation box (in some dimension), LAMMPS does NOT implement the same logic described above for the *box* style, to insure exactly one particle at periodic boundaries. If this is desired, you should either use the *box* style, or tweak the region size to get precisely the particles you want.

For the *single* style, a single particle is added to the system at the specified coordinates. This can be useful for debugging purposes or to create a tiny system with a handful of particles at specified positions.

New in version 2Jun2022.

For the *mesh* style, a file with a triangle mesh in *STL format* is read and one or more particles are placed into the area of each triangle. The reader supports both ASCII and binary files conforming to the format on the Wikipedia page. Binary STL files (e.g. as frequently offered for 3d-printing) can also be first converted to ASCII for editing with the *stl\_bin2txt tool*. The use of the *units box* option is required. There are two algorithms available for placing atoms: *bisect* and *grand*. They can be selected via the *meshmode* option; *bisect* is the default. If the atom style allows it, the radius will be set to a value depending on the algorithm and the value of the *radscale* parameter (see below), and the atoms created from the mesh are assigned a new molecule ID.

In *bisect* mode a particle is created at the center of each triangle unless the average distance of the triangle vertices from its center is larger than the



*radthresh* value (default is lattice spacing in x-direction). In case the average distance is over the threshold, the triangle is recursively split into two halves along the the longest side until the threshold is reached. There will be at least one sphere per triangle. The value of *radthresh* is set as an argument to *meshmode bisect*. The average distance of the vertices from the center is also used to set the radius.

In *grand* mode a quasi-random sequence is used to distribute particles on mesh triangles using an approach by (*Roberts*). Particles are added to the triangle until the minimum number density is met or exceeded such that every triangle will have at least one particle. The minimum number density is set as an argument to the *grand* option. The radius will be set so that the sum of the area of the radius of the particles created in place of a triangle will be equal to the area of that triangle.

---

**Note:** The atom placement algorithms in the *mesh* style benefit from meshes where triangles are close to equilateral. It is therefore recommended to pre-process STL files to optimize the mesh accordingly. There are multiple open source and commercial software tools available with the capability to generate optimized meshes.

---



---

**Note:** In most cases the atoms created in *mesh* style will become an immobile or rigid object that would not be time integrated or moved by *fix move* or *fix rigid*. For computational efficiency *and* to avoid undesired contributions to pressure and potential energy due to close contacts, it is usually beneficial to exclude computing interactions between the created particles using *neigh\_modify exclude*.

---

For the *random* style, *N* particles are added to the system at randomly generated coordinates, which can be useful for generating an amorphous system. The particles are created one by one using the specified random number *seed*, resulting in the same set of particle coordinates, independent of how many processors are being used in the simulation. Unless the *overlap* keyword is specified, particles created by the *random* style will typically be highly overlapped. Various additional criteria can be used to accept or reject a random particle insertion; see the keyword discussion below. Multiple attempts per particle are made (see the *maxtry* keyword) until the insertion is either successful or fails. If this command fails to add all requested *N* particles, a warning will be output.

If the *region-ID* argument is specified as NULL, then the randomly created particles will be anywhere in the simulation box. If a *region-ID* is specified, a geometric volume is filled that is both inside the simulation box and is also consistent with the region volume. See the *region* command for details. Note that a region can be specified so that its “volume” is either inside or outside its geometric boundary.

Note that the *create\_atoms* command adds particles to those that already exist. This means it can be used to add particles to a system previously read in from a data or restart file. Or the *create\_atoms* command can be used multiple times, to add multiple sets of particles to the simulation. For example, grain boundaries can be created, by interleaving the *create\_atoms* command with *lattice* commands specifying different orientations.

When this command is used, care should be taken to insure the resulting system does not contain particles that are highly overlapped. Such overlaps will cause many interatomic potentials to compute huge energies and forces, leading to bad dynamics. There are several strategies to avoid this problem:

- Use the *delete\_atoms overlap* command after *create\_atoms*. For example, this strategy can be used to overlay and surround a large protein molecule with a volume of water molecules, then delete water molecules that overlap with the protein atoms.
- For the *random* style, use the optional *overlap* keyword to avoid overlaps when each new particle is created.
- Before running dynamics on an overlapped system, perform an *energy minimization*. Or run initial dynamics with *pair\_style soft* or with *fix nve/limit* to un-overlap the particles, before running normal dynamics.

---

**Note:** You cannot use any of the styles explained above to create atoms that are outside the simulation box; they will just be ignored by LAMMPS. This is true even if you are using shrink-wrapped box boundaries, as specified by the *boundary* command. However, you can first use the *change\_box* command to temporarily expand the box, then add

atoms via `create_atoms`, then finally use `change_box` command again if needed to re-shrink-wrap the new atoms. See the [change\\_box](#) doc page for an example of how to do this, using the `create_atoms single` style to insert a new atom outside the current simulation box.

---

Individual atoms are inserted by this command, unless the `mol` keyword is used. It specifies a *template-ID* previously defined using the `molecule` command, which reads a file that defines the molecule. The coordinates, atom types, charges, etc, as well as any bond/angle/etc and special neighbor information for the molecule can be specified in the molecule file. See the `molecule` command for details. The only settings required to be in this file are the coordinates and types of atoms in the molecule.

Using a lattice to add molecules, e.g. via the `box` or `region` or `single` styles, is exactly the same as adding atoms on lattice points, except that entire molecules are added at each point, i.e. on the point defined by each basis atom in the unit cell as it tiles the simulation box or region. This is done by placing the geometric center of the molecule at the lattice point, and (by default) giving the molecule a random orientation about the point. The random *seed* specified with the `mol` keyword is used for this operation, and the random numbers generated by each processor are different. This means the coordinates of individual atoms (in the molecules) will be different when running on different numbers of processors, unlike when atoms are being created in parallel.

Note that with random rotations, it may be important to use a lattice with a large enough spacing that adjacent molecules will not overlap, regardless of their relative orientations. See the description of the `rotate` keyword below, which overrides the default random orientation and inserts all molecules at a specified orientation.

---

**Note:** If the `create_box` command is used to create the simulation box, followed by the `create_atoms` command with its `mol` option for adding molecules, then you typically need to use the optional keywords allowed by the `create_box` command for extra bonds (angles,etc) or extra special neighbors. This is because by default, the `create_box` command sets up a non-molecular system that does not allow molecules to be added.

---

This is the meaning of the other optional keywords.

The `basis` keyword is only used when atoms (not molecules) are being created. It specifies an atom type that will be assigned to specific basis atoms as they are created. See the `lattice` command for specifics on how basis atoms are defined for the unit cell of the lattice. By default, all created atoms are assigned the argument *type* as their atom type.

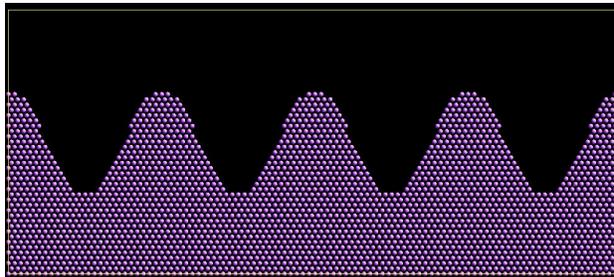
The `ratio` and `subset` keywords can be used in conjunction with the `box` or `region` styles to limit the total number of particles inserted. The lattice defines a set of  $N_{latt}$  eligible sites for inserting particles, which may be limited by the `region` style or the `var` and `set` keywords. For the `ratio` keyword, only the specified fraction of them ( $0 \leq f \leq 1$ ) will be assigned particles. For the `subset` keyword only the specified  $N_{subset}$  of them will be assigned particles. In both cases the assigned lattice sites are chosen randomly. An iterative algorithm is used that insures the correct number of particles are inserted, in a perfectly random fashion. Which lattice sites are selected will change with the number of processors used.

The `remap` keyword only applies to the `single` style. If it is set to *yes*, then if the specified position is outside the simulation box, it will mapped back into the box, assuming the relevant dimensions are periodic. If it is set to *no*, no remapping is done and no particle is created if its position is outside the box.

The `var` and `set` keywords can be used together to provide a criterion for accepting or rejecting the addition of an individual atom, based on its coordinates. They apply to all styles except `single`. The *name* specified for the `var` keyword is the name of an *equal-style variable* that should evaluate to a zero or non-zero value based on one or two or three variables that will store the *x*, *y*, or *z* coordinates of an atom (one variable per coordinate). If used, these other variables must be *internal-style variables* defined in the input script; their initial numeric value can be anything. They must be internal-style variables, because this command resets their values directly. The `set` keyword is used to identify the names of these other variables, one variable for the *x*-coordinate of a created atom, one for *y*, and one for *z*.

When an atom is created, its  $(x, y, z)$  coordinates become the values for any *set* variable that is defined. The *var* variable is then evaluated. If the returned value is 0.0, the atom is not created. If it is non-zero, the atom is created.

As an example, these commands can be used in a 2d simulation, to create a sinusoidal surface. Note that the surface is “rough” due to individual lattice points being “above” or “below” the mathematical expression for the sinusoidal curve. If a finer lattice were used, the sinusoid would appear to be “smoother”. Also note the use of the “xlat” and “ylat” *thermo\_style* keywords, which converts lattice spacings to distance.



```

dimension      2
variable      x equal 100
variable      y equal 25
lattice       hex 0.8442
region        box block 0 $x 0 $y -0.5 0.5
create_box    1 box

variable      xx internal 0.0
variable      yy internal 0.0
variable      v equal "(0.2*v_y*ylat * cos(v_xx/xlat * 2.0*PI*4.0/v_x) + 0.5*v_y*ylat - v_
->yy) > 0.0"
create_atoms  1 box var v set x xx set y yy
write_dump    all atom sinusoid.lammpstrj

```

The *rotate* keyword allows specification of the orientation at which molecules are inserted. The axis of rotation is determined by the rotation vector  $(R_x, R_y, R_z)$  that goes through the insertion point. The specified *theta* determines the angle of rotation around that axis. Note that the direction of rotation for the atoms around the rotation axis is consistent with the right-hand rule: if your right-hand’s thumb points along  $R$ , then your fingers wrap around the axis in the direction of rotation.

The *radscale* keyword only applies to the *mesh* style and adjusts the radius of created particles (see above), provided this is supported by the atom style. Its value is a prefactor (must be  $> 0.0$ , default is 1.0) that is applied to the atom radius inferred from the size of the individual triangles in the triangle mesh that the particle corresponds to.

New in version 2Jun2022.

The *overlap* keyword only applies to the *random* style. It prevents newly created particles from being created closer than the specified *Doverlap* distance from any other particle. When the particles being created are molecules, the radius of the molecule (from its geometric center) is added to *Doverlap*. If particles have finite size (see *atom\_style sphere* for example) *Doverlap* should be specified large enough to include the particle size in the non-overlapping criterion.

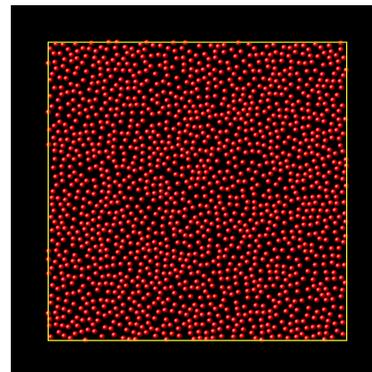
**Note:** Checking for overlaps is a costly  $\mathcal{O}(N(N + M))$  operation for inserting  $N$  new particles into a system with  $M$  existing particles. This is because distances to all  $M$  existing particles are computed for each new particle that is added. Thus the intended use of this keyword is to add relatively small numbers of particles to systems that remain at a relatively low density even after the new particles are created. Careful use of the *maxtry* keyword in combination with *overlap* is recommended. See the discussion above about systems with overlapped particles for alternate strategies that allow for overlapped insertions.

The *maxtry* keyword only applies to the *random* style. It limits the number of attempts to generate valid coordinates for a single new particle that satisfy all requirements imposed by the *region*, *var*, and *overlap* keywords. The default is 10 attempts per particle before the loop over the requested  $N$  particles advances to the next particle. Note that if

insertion success is unlikely (e.g., inserting new particles into a dense system using the *overlap* keyword), setting the *maxtry* keyword to a large value may result in this command running for a long time.

Here is an example for the *random* style using these commands

```
units          lj
dimension      2
region         box block 0 50 0 50 -0.5 0.5
create_box     1 box
create_atoms   1 random 2000 13487 NULL overlap 1.0 maxtry 50
pair_style     lj/cut 2.5
pair_coeff      1 1 1.0 1.0 2.5
```



to produce a system as shown in the image with 1520 particles (out of 2000 requested) that are moderately dense and which have no overlaps sufficient to prevent the LJ pair\_style from running properly (because the overlap criterion is 1.0). The create\_atoms command ran for 0.3 s on a single CPU core.

The *units* keyword determines the meaning of the distance units used to specify the coordinates of the one particle created by the *single* style, or the overlap distance *Overlap* by the *overlap* keyword. A *box* value selects standard distance units as defined by the *units* command (e.g., Å for units = *real* or *metal*. A *lattice* value means the distance units are in lattice spacings.

Atom IDs are assigned to created atoms in the following way. The collection of created atoms are assigned consecutive IDs that start immediately following the largest atom ID existing before the create\_atoms command was invoked. This is done by the processor's communicating the number of atoms they each own, the first processor numbering its atoms from 1 to  $N_1$ , the second processor from  $N_1 + 1$  to  $N_2$ , and so on, where  $N_1$  is the number of atoms owned by the first processor,  $N_2$  is the number owned by the second processor, and so forth. Thus, when the same simulation is performed on different numbers of processors, there is no guarantee a particular created atom will be assigned the same ID in both simulations. If molecules are being created, molecule IDs are assigned to created molecules in a similar fashion.

Aside from their ID, atom type, and *xyz* position, other properties of created atoms are set to default values, depending on which quantities are defined by the chosen *atom style*. See the *atom style* command for more details. See the *set* and *velocity* commands for info on how to change these values.

- charge = 0.0
- dipole moment magnitude = 0.0
- diameter = 1.0
- shape = 0.0 0.0 0.0
- density = 1.0
- volume = 1.0
- velocity = 0.0 0.0 0.0
- angular velocity = 0.0 0.0 0.0
- angular momentum = 0.0 0.0 0.0
- quaternion = (1,0,0,0)
- bonds, angles, dihedrals, impropers = none



If molecules are being created, these defaults can be overridden by values specified in the file read by the *molecule* command. That is, the file typically defines bonds (angles, etc.) between atoms in the molecule, and can optionally define charges on each atom.

Note that the *sphere* atom style sets the default particle diameter to 1.0 as well as the density. This means the mass for the particle is not 1.0, but is  $\frac{\pi}{6}d^3 = 0.5236$ , where  $d$  is the diameter. When using the *mesh* style, the particle diameter is adjusted from the size of the individual triangles in the triangle mesh.

Note that the *ellipsoid* atom style sets the default particle shape to (0.0 0.0 0.0) and the density to 1.0, which means it is a point particle, not an ellipsoid, and has a mass of 1.0.

Note that the *peri* style sets the default volume and density to 1.0 and thus also set the mass for the particle to 1.0.

The *set* command can be used to override many of these default settings.

## 16.17.4 Restrictions

An *atom\_style* must be previously defined to use this command.

A rotation vector specified for a single molecule must be in the z-direction for a 2d model.

## 16.17.5 Related commands

*lattice, region, create\_box, read\_data, read\_restart*

## 16.17.6 Default

The default for the *basis* keyword is that all created atoms are assigned the argument *type* as their atom type (when single atoms are being created). The other defaults are *remap* = no, *rotate* = random, *radscale* = 1.0, *radthresh* = x-lattice spacing, *overlap* not checked, *maxtry* = 10, and *units* = lattice.

(Roberts) R. Roberts (2019) “Evenly Distributing Points in a Triangle.” Extreme Learning. <http://extremelearning.com.au/evenly-distributing-points-in-a-triangle/>

## 16.18 create\_bonds command

### 16.18.1 Syntax

```
create_bonds style args ... keyword value ...
```

- style = *many* or *single/bond* or *single/angle* or *single/dihedral* or *single/improper*

*many* args = group-ID group2-ID btype rmin rmax

group-ID = ID of first group

group2-ID = ID of second group, bonds will be between atoms in the 2 groups

btype = bond type of created bonds

rmin = minimum distance between pair of atoms to bond together

rmax = maximum distance between pair of atoms to bond together

*single/bond* args = btype batom1 batom2

```
btype = bond type of new bond
batom1,batom2 = atom IDs for two atoms in bond
single/angle args = aatype aatom1 aatom2 aatom3
  aatype = angle type of new angle
  aatom1,aatom2,aatom3 = atom IDs for three atoms in angle
single/dihedral args = dtype datom1 datom2 datom3 datom4
  dtype = dihedral type of new dihedral
  datom1,datom2,datom3,datom4 = atom IDs for four atoms in dihedral
single/improper args = itype iatom1 iatom2 iatom3 iatom4
  itype = improper type of new improper
  iatom1,iatom2,iatom3,iatom4 = atom IDs for four atoms in improper
```

- zero or more keyword/value pairs may be appended
- keyword = *special*

*special* value = *yes* or *no*

## 16.18.2 Examples

```
create_bonds many all all 1 1.0 1.2
create_bonds many surf solvent 3 2.0 2.4
create_bonds single/bond 1 1 2
create_bonds single/angle 5 52 98 107 special no
create_bonds single/dihedral 2 4 19 27 101
create_bonds single/improper 3 23 26 31 57
```

## 16.18.3 Description

Create bonds between pairs of atoms that meet a specified distance criteria. Or create a single bond, angle, dihedral or improper between 2, 3, or 4 specified atoms.

The new bond (angle, dihedral, improper) interactions will then be computed during a simulation by the bond (angle, dihedral, improper) potential defined by the *bond\_style*, *bond\_coeff*, *angle\_style*, *angle\_coeff*, *dihedral\_style*, *dihedral\_coeff*, *improper\_style*, *improper\_coeff* commands.

The *many* style is useful for adding bonds to a system (e.g., between nearest neighbors in a lattice of atoms) without having to enumerate all the bonds in the data file read by the *read\_data* command.

The *single* styles are useful for adding bonds, angles, dihedrals, and improvers to a system incrementally, then continuing a simulation.

Note that this command does not auto-create any angle, dihedral, or improper interactions when a bond is added, nor does it auto-create any bonds when an angle, dihedral, or improper is added. It also will not auto-create any angles when a dihedral or improper is added. Thus, the flexibility of this command is limited. It can be used several times to create different types of bond at different distances, but it cannot typically auto-create all the bonds or angles or dihedrals or improvers that would normally be defined in a data file for a complex system of molecules.

---

**Note:** If the system has no bonds (angles, dihedrals, improvers) to begin with, or if more bonds per atom are being added than currently exist, then you must ensure that the number of bond types and the maximum number of bonds per atom are set to large enough values, and similarly for angles, dihedrals, and improvers, otherwise an error may occur when too many bonds (angles, dihedrals, improvers) are added to an atom. If the *read\_data* command is used to define the system, these parameters can be set via the “bond types” and “extra bond per atom” fields in the header section of the data file. If the *create\_box* command is used to define the system, these two parameters can be set via its optional

*bond/types* and *extra/bond/per/atom* arguments, and similarly for angles, dihedrals, and improper. See the doc pages for these two commands for details.

The *many* style will create bonds between pairs of atoms  $I, J$ , where  $I$  is in one of the two specified groups and  $J$  is in the other. The two groups can be the same (e.g., group “all”). The created bonds will be of bond type *btype*, where *btype* must be a value between 1 and the number of bond types defined.

For a bond to be created, an  $I, J$  pair of atoms must be a distance  $D$  apart such that  $r_{\min} \leq D \leq r_{\max}$ .

The following settings must have been made in an input script before this style is used:

- special\_bonds weight for 1–2 interactions must be 0.0
- a *pair\_style* must be defined
- no *kpace\_style* defined
- minimum *pair\_style* cutoff + *neighbor* skin  $\geq r_{\max}$

These settings are required so that a neighbor list can be created to search for nearby atoms. Pairs of atoms that are already bonded cannot appear in the neighbor list, to avoid creation of duplicate bonds. The neighbor list for all atom type pairs must also extend to a distance that encompasses the *rmax* for new bonds to create.

**Note:** If you want to create bonds between pairs of 1–3 or 1–4 atoms in the current bond topology, then you need to use *special\_bonds lj 0 1 1* to insure those pairs appear in the neighbor list. They will not appear with the default special\_bonds settings, which are zero for 1–2, 1–3, and 1–4 atoms. 1–3 or 1–4 atoms are those which are two hops or three hops apart in the bond topology.

An additional requirement for this style is that your system must be ready to perform a simulation. This means, for example, that all *pair\_style* coefficients be set via the *pair\_coeff* command. A *bond\_style* command and all bond coefficients must also be set, even if no bonds exist before this command is invoked. This is because the building of neighbor list requires initialization and setup of a simulation, similar to what a *run* command would require.

Note that you can change any of these settings after this command executes (e.g., if you wish to use long-range Coulombic interactions) via the *kpace\_style* command for your subsequent simulation.

The *single/bond* style creates a single bond of type *btype* between two atoms with IDs *batom1* and *batom2*. *Btype* must be a value between 1 and the number of bond types defined.

The *single/angle* style creates a single angle of type *atype* between three atoms with IDs *aatom1*, *aatom2*, and *aatom3*. The ordering of the atoms is the same as in the *Angles* section of a data file read by the *read\_data* command (i.e., the three atoms are ordered linearly within the angle; the central atom is *aatom2*). *Atype* must be a value between 1 and the number of angle types defined.

The *single/dihedral* style creates a single dihedral of type *dtype* between four atoms with IDs *datom1*, *datom2*, *datom3*, and *datom4*. The ordering of the atoms is the same as in the *Dihedrals* section of a data file read by the *read\_data* command. I.e. the 4 atoms are ordered linearly within the dihedral. *dtype* must be a value between 1 and the number of dihedral types defined.

The *single/improper* style creates a single improper of type *itype* between four atoms with IDs *iatom1*, *iatom2*, *iatom3*, and *iatom4*. The ordering of the atoms is the same as in the *Improper* section of a data file read by the *read\_data* command. I.e. the 4 atoms are ordered linearly within the improper. *itype* must be a value between 1 and the number of improper types defined.

The keyword *special* controls whether an internal list of special bonds is created after one or more bonds, or a single angle, dihedral, or improper is added to the system.

The default value is *yes*. A value of *no* cannot be used with the *many* style.

This is an expensive operation since the bond topology for the system must be walked to find all 1–2, 1–3, and 1–4 interactions to store in an internal list, which is used when pairwise interactions are weighted; see the *special\_bonds* command for details.

Thus if you are adding a few bonds or a large list of angles all at the same time, by using this command repeatedly, it is more efficient to only trigger the internal list to be created once, after the last bond (or angle, or dihedral, or improper) is added:

```
create_bonds single/bond 5 52 98 special no
create_bonds single/bond 5 73 74 special no
...
create_bonds single/bond 5 17 386 special no
create_bonds single/bond 4 112 183 special yes
```

Note that you **must** ensure the internal list is rebuilt after the last bond (angle, dihedral, improper) is added, *before* performing a simulation. Otherwise, pairwise interactions will not be properly excluded or weighted. LAMMPS does **not** check that you have done this correctly.

---

## 16.18.4 Restrictions

This command cannot be used with molecular systems defined using molecule template files via the *molecule* and *atom\_style template* commands.

## 16.18.5 Related commands

*create\_atoms*, *delete\_bonds*

## 16.18.6 Default

The keyword default is *special = yes*.

# 16.19 create\_box command

## 16.19.1 Syntax

```
create_box N region-ID keyword value ...
```

- N = # of atom types to use in this simulation
- region-ID = ID of region to use as simulation domain
- zero or more keyword/value pairs may be appended
- keyword = *bond/types* or *angle/types* or *dihedral/types* or *improper/types* or *extra/bond/per/atom* or *extra/angle/per/atom* or *extra/dihedral/per/atom* or *extra/improper/per/atom* or *extra/special/per/atom*

```

bond/types value = # of bond types
angle/types value = # of angle types
dihedral/types value = # of dihedral types
improper/types value = # of improper types
extra/bond/per/atom value = # of bonds per atom
extra/angle/per/atom value = # of angles per atom
extra/dihedral/per/atom value = # of dihedrals per atom
extra/improper/per/atom value = # of impropers per atom
extra/special/per/atom value = # of special neighbors per atom

```

## 16.19.2 Examples

```

create_box 2 mybox
create_box 2 mybox bond/types 2 extra/bond/per/atom 1

```

## 16.19.3 Description

This command creates a simulation box based on the specified region. Thus a *region* command must first be used to define a geometric domain. It also partitions the simulation box into a regular 3d grid of rectangular bricks, one per processor, based on the number of processors being used and the settings of the *processors* command. The partitioning can later be changed by the *balance* or *fix balance* commands.

The argument N is the number of atom types that will be used in the simulation.

If the region is not of style *prism*, then LAMMPS encloses the region (block, sphere, etc.) with an axis-aligned orthogonal bounding box which becomes the simulation domain.

If the region is of style *prism*, LAMMPS creates a non-orthogonal simulation domain shaped as a parallelepiped with triclinic symmetry. As defined by the *region prism* command, the parallelepiped has its “origin” at  $(x_{lo}, y_{lo}, z_{lo})$  and is defined by three edge vectors starting from the origin given by  $\vec{a} = (x_{hi} - x_{lo}, 0, 0)$ ;  $\vec{b} = (xy, y_{hi} - y_{lo}, 0)$ ; and  $\vec{c} = (xz, yz, z_{hi} - z_{lo})$ . The parameters *xy*, *xz*, and *yz* can be 0.0 or positive or negative values and are called “tilt factors” because they are the amount of displacement applied to faces of an originally orthogonal box to transform it into the parallelepiped.

By default, a *prism* region used with the *create\_box* command must have tilt factors (*xy*, *xz*, *yz*) that do not skew the box more than half the distance of the parallel box length. For example, if  $x_{lo} = 2$  and  $x_{hi} = 12$ , then the *x* box length is 10 and the *xy* tilt factor must be between  $-5$  and  $5$ . Similarly, both *xz* and *yz* must be between  $-(x_{hi} - x_{lo})/2$  and  $+(y_{hi} - y_{lo})/2$ . Note that this is not a limitation, since if the maximum tilt factor is 5 (as in this example), then configurations with tilt =  $\dots, -15, -5, 5, 15, 25, \dots$  are all geometrically equivalent. If you wish to define a box with tilt factors that exceed these limits, you can use the *box tilt* command, with a setting of *large*; a setting of *small* is the default.

See the *Howto triclinic* page for a geometric description of triclinic boxes, as defined by LAMMPS, and how to transform these parameters to and from other commonly used triclinic representations.

When a *prism* region is used, the simulation domain should normally be periodic in the dimension that the tilt is applied to, which is given by the second dimension of the tilt factor (e.g., *y* for *xy* tilt). This is so that pairs of atoms interacting across that boundary will have one of them shifted by the tilt factor. Periodicity is set by the *boundary* command. For example, if the *xy* tilt factor is non-zero, then the *y* dimension should be periodic. Similarly, the *z* dimension should be periodic if *xz* or *yz* is non-zero. LAMMPS does not require this periodicity, but you may lose atoms if this is not the case.

Note that if your simulation will tilt the box (e.g., via the *fix deform* command), the simulation box must be set up to be triclinic, even if the tilt factors are initially 0.0. You can also change an orthogonal box to a triclinic box or vice versa by using the *change box* command with its *ortho* and *triclinic* options.

**Note:** If the system is non-periodic (in a dimension), then you should not make the lo/hi box dimensions (as defined in your *region* command) radically smaller/larger than the extent of the atoms you eventually plan to create (e.g., via the *create\_atoms* command). For example, if your atoms extend from 0 to 50, you should not specify the box bounds as -10000 and 10000. This is because as described above, LAMMPS uses the specified box size to lay out the 3d grid of processors. A huge (mostly empty) box will be sub-optimal for performance when using “fixed” boundary conditions (see the *boundary* command). When using “shrink-wrap” boundary conditions (see the *boundary* command), a huge (mostly empty) box may cause a parallel simulation to lose atoms the first time that LAMMPS shrink-wraps the box around the atoms.

---

The optional keywords can be used to create a system that allows for bond (angle, dihedral, improper) interactions, or for molecules with special 1-2, 1-3, or 1-4 neighbors to be added later. These optional keywords serve the same purpose as the analogous keywords that can be used in a data file which are recognized by the *read\_data* command when it sets up a system.

Note that if these keywords are not used, then the *create\_box* command creates an atomic (non-molecular) simulation that does not allow bonds between pairs of atoms to be defined, or a *bond potential* to be specified, or for molecules with special neighbors to be added to the system by commands such as *create\_atoms mol*, *fix deposit* or *fix pour*.

As an example, see the examples/deposit/in.deposit.molecule script, which deposits molecules onto a substrate. Initially there are no molecules in the system, but they are added later by the *fix deposit* command. The *create\_box* command in the script uses the *bond/types* and *extra/bond/per/atom* keywords to allow this. If the added molecule contained more than one special bond (allowed by default), an *extra/special/per/atom* keyword would also need to be specified.

---

## 16.19.4 Restrictions

An *atom\_style* and *region* must have been previously defined to use this command.

## 16.19.5 Related commands

*read\_data*, *create\_atoms*, *region*

## 16.19.6 Default

none

## 16.20 delete\_atoms command

### 16.20.1 Syntax

`delete_atoms style args keyword value ...`

- *style* = *group* or *region* or *overlap* or *random* or *variable*

```

group args = group-ID
region args = region-ID
overlap args = cutoff group1-ID group2-ID
  cutoff = delete one atom from pairs of atoms within the cutoff (distance units)
  group1-ID = one atom in pair must be in this group
  group2-ID = other atom in pair must be in this group
random args = ranstyle value eflag group-ID region-ID seed
  ranstyle = fraction or count
  for fraction:
    value = fraction (0.0 to 1.0) of eligible atoms to delete
    eflag = no for fast approximate deletion, yes for exact deletion
  for count:
    value = number of atoms to delete
    eflag = no for warning if count > eligible atoms, yes for error
  group-ID = group within which to perform deletions
  region-ID = region within which to perform deletions
               or NULL to only impose the group criterion
  seed = random number seed (positive integer)
variable args = variable-name

```

- zero or more keyword/value pairs may be appended
- keyword = *compress* or *bond* or *mol*

```

compress value = no or yes
bond value = no or yes
mol value = no or yes

```

## 16.20.2 Examples

```

delete_atoms group edge
delete_atoms region sphere compress no
delete_atoms overlap 0.3 all all
delete_atoms overlap 0.5 solvent colloid
delete_atoms random fraction 0.1 yes all cube 482793 bond yes
delete_atoms random fraction 0.3 no polymer NULL 482793 bond yes
delete_atoms random count 500 no ions NULL 482793
delete_atoms variable checkers

```

## 16.20.3 Description

Delete the specified atoms. This command can be used, for example, to carve out voids from a block of material or to delete created atoms that are too close to each other (e.g., at a grain boundary).

For style *group*, all atoms belonging to the group are deleted.

For style *region*, all atoms in the region volume are deleted. Additional atoms can be deleted if they are in a molecule for which one or more atoms were deleted within the region; see the *mol* keyword discussion below.

For style *overlap* pairs of atoms whose distance of separation is within the specified cutoff distance are searched for, and one of the 2 atoms is deleted. Only pairs where one of the two atoms is in the first group specified and the other atom is in the second group are considered. The atom that is in the first group is the one that is deleted.

Note that it is OK for the two group IDs to be the same (e.g., *group all*), or for some atoms to be members of both groups. In these cases, either atom in the pair may be deleted. Also note that if there are atoms which are members of both

groups, the only guarantee is that at the end of the deletion operation, enough deletions will have occurred that no atom pairs within the cutoff will remain (subject to the group restriction). There is no guarantee that the minimum number of atoms will be deleted, or that the same atoms will be deleted when running on different numbers of processors.

For style *random* a subset of eligible atoms are deleted. Which atoms to delete are chosen randomly using the specified random number *seed*. Which atoms are deleted may vary when running on different numbers of processors.

For *ranstyle = fraction*, the specified fractional *value* (0.0 to 1.0) of eligible atoms are deleted. If *eflag* is set to *no*, then the number of deleted atoms will be approximate, but the operation will be fast. If *eflag* is set to *yes*, then the number deleted will match the requested fraction, but for large systems the selection of deleted atoms may take additional time to determine.

For *ranstyle = count*, the specified integer *value* is the number of eligible atoms are deleted. If *eflag* is set to *no*, then if the requested number is larger then the number of eligible atoms, a warning is issued and only the eligible atoms are deleted instead of the requested *value*. If *eflag* is set to *yes*, an error is triggered instead and LAMMPS will exit. For large systems the selection of atoms to delete may take additional time to determine, the same as for requesting an exact fraction with *pstyle = fraction*.

Which atoms are eligible for deletion for style *random* is determined by the specified *group-ID* and *region-ID*. To be eligible, an atom must be in both the specified group and region. If *group-ID = all*, there is effectively no group criterion. If *region-ID* is specified as NULL, no region criterion is imposed.

For style *variable*, all atoms for which the atom-style variable with the given name evaluates to non-zero will be deleted. Additional atoms can be deleted if they are in a molecule for which one or more atoms were deleted within the region; see the *mol* keyword discussion below. This option allows complex selections of atoms not covered by the other options listed above.

---

Here is the meaning of the optional keywords.

If the *compress* keyword is set to *yes*, then after atoms are deleted, then atom IDs are re-assigned so that they run from 1 to the number of atoms in the system. Note that this is not done for molecular systems (see the *atom\_style* command), regardless of the *compress* setting, since it would foul up the bond connectivity that has already been assigned. However, the *reset\_atom\_ids* command can be used after this command to accomplish the same thing.

Note that the re-assignment of IDs is not really a compression, where gaps in atom IDs are removed by decrementing atom IDs that are larger. Instead the IDs for all atoms are erased, and new IDs are assigned so that the atoms owned by individual processors have consecutive IDs, as the *create\_atoms* command explains.

A molecular system with fixed bonds, angles, dihedrals, or improper interactions, is one where the topology of the interactions is typically defined in the data file read by the *read\_data* command, and where the interactions themselves are defined with the *bond\_style*, *angle\_style*, etc. commands. If you delete atoms from such a system, you must be careful not to end up with bonded interactions that are stored by remaining atoms but which include deleted atoms. This will cause LAMMPS to generate a “missing atoms” error when the bonded interaction is computed. The *bond* and *mol* keywords offer two ways to do that.

If the *bond* keyword is set to *yes* then any bond or angle or dihedral or improper interaction that includes a deleted atom is also removed from the lists of such interactions stored by non-deleted atoms. Note that simply deleting interactions due to dangling bonds (e.g., at a surface) may result in an inaccurate or invalid model for the remaining atoms.

If the *mol* keyword is set to *yes*, then for every atom that is deleted, all other atoms in the same molecule (with the same molecule ID) will also be deleted. This is not done for atoms with molecule ID = 0, since such an ID is assumed to flag isolated atoms that are not part of molecules.

---

**Note:** The molecule deletion operation is invoked after all individual atoms have been deleted using the rules described above for each style. This means additional atoms may be deleted that are not in the group or region, that are not required by the overlap cutoff criterion, or that will create a higher fraction of porosity than was requested.

---



## 16.20.4 Restrictions

The *overlap* styles requires inter-processor communication to acquire ghost atoms and build a neighbor list. This means that your system must be ready to perform a simulation before using this command (force fields setup, atom masses set, etc.). Since a neighbor list is used to find overlapping atom pairs, it also means that you must define a *pair style* with the minimum force cutoff distance between any pair of atoms types (plus the *neighbor* skin)  $\geq$  the specified overlap cutoff.

If the *special\_bonds* command is used with a setting of 0, then a pair of bonded atoms (1–2, 1–3, or 1–4) will not appear in the neighbor list, and thus will not be considered for deletion by the *overlap* styles. You probably do not want to delete one atom in a bonded pair anyway.

The *bond yes* option cannot be used with molecular systems defined using molecule template files via the *molecule* and *atom\_style template* commands.

## 16.20.5 Related commands

*create\_atoms*, *reset\_atom\_ids*

## 16.20.6 Default

The option defaults are *compress* = yes, *bond* = no, *mol* = no.

## 16.21 delete\_bonds command

### 16.21.1 Syntax

```
delete_bonds group-ID style arg keyword ...
```

- group-ID = group ID
- style = *multi* or *atom* or *bond* or *angle* or *dihedral* or *improper* or *stats*
  - multi* arg = none
  - atom* arg = an atom type or range of types (see below)
  - bond* arg = a bond type or range of types (see below)
  - angle* arg = an angle type or range of types (see below)
  - dihedral* arg = a dihedral type or range of types (see below)
  - improper* arg = an improper type or range of types (see below)
  - stats* arg = none
- zero or more keywords may be appended
- keyword = *any* or *undo* or *remove* or *special*

## 16.21.2 Examples

```
delete_bonds frozen multi remove
delete_bonds all atom 4 special
delete_bonds all bond 0*3 special
delete_bonds all stats
```

## 16.21.3 Description

Turn off (or on) molecular topology interactions (i.e., bonds, angles, dihedrals, and/or improper). This command is useful for deleting interactions that have been previously turned off by bond-breaking potentials. It is also useful for turning off topology interactions between frozen or rigid atoms. Pairwise interactions can be turned off via the *neigh\_modify exclude* command. The *fix shake* command also effectively turns off certain bond and angle interactions.

For all styles, by default, an interaction is only turned off (or on) if all the atoms involved are in the specified group. See the *any* keyword to change the behavior.

Several of the styles (*atom*, *bond*, *angle*, *dihedral*, *improper*) take a *type* as an argument. The specified *type* should be an integer from 0 to  $N$ , where  $N$  is the number of relevant types (atom types, bond types, etc.). A value of 0 is only relevant for style *bond*; see details below. In all cases, a wildcard asterisk can be used in place of or in conjunction with the *type* argument to specify a range of types. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the number of types, then an asterisk with no numeric values means all types from 0 to  $N$ . A leading asterisk means all types from 0 to  $n$  (inclusive). A trailing asterisk means all types from  $m$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive). Note that it is fine to include a type of 0 for non-bond styles; it will simply be ignored.

For style *multi* all bond, angle, dihedral, and improper interactions of any type, involving atoms in the group, are turned off.

Style *atom* is the same as style *multi* except that in addition, one or more of the atoms involved in the bond, angle, dihedral, or improper interaction must also be of the specified atom type.

For style *bond*, only bonds are candidates for turn-off, and the bond must also be of the specified type. Styles *angle*, *dihedral*, and *improper* are treated similarly.

For style *bond*, you can set the type to 0 to delete bonds that have been previously broken by a bond-breaking potential (which sets the bond type to 0 when a bond is broken); for example, see the *bond\_style quartic* command.

For style *stats* no interactions are turned off (or on); the status of all interactions in the specified group is simply reported. This is useful for diagnostic purposes if bonds have been turned off by a bond-breaking potential during a previous run.

The default behavior of the *delete\_bonds* command is to turn off interactions by toggling their type to a negative value, but not to permanently remove the interaction. For example, a *bond\_type* of 2 is set to  $-2$ . The neighbor list creation routines will not include such an interaction in their interaction lists. The default is also to not alter the list of 1–2, 1–3, or 1–4 neighbors computed by the *special\_bonds* command and used to weight pairwise force and energy calculations. This means that pairwise computations will proceed as if the bond (or angle, etc.) were still turned on.

Several keywords can be appended to the argument list to alter the default behaviors.

The *any* keyword changes the requirement that all atoms in the bond (angle, etc) must be in the specified group in order to turn off the interaction. Instead, if any of the atoms in the interaction are in the specified group, it will be turned off (or on if the *undo* keyword is used).

The *undo* keyword inverts the *delete\_bonds* command so that the specified bonds, angles, etc are turned on if they are currently turned off. This means a negative value is toggled to positive. For example, for style *angle*, if *type* is specified as 2, then all angles with current type =  $-2$  are reset to type = 2. Note that the *fix shake* command also sets bond and angle types negative, so this option should not be used on those interactions.

The *remove* keyword is invoked at the end of the `delete_bonds` operation. It causes turned-off bonds (angles, etc.) to be removed from each atom's data structure and then adjusts the global bond (angle, etc.) counts accordingly. Removal is a permanent change; removed bonds cannot be turned back on via the *undo* keyword. Removal does not alter the pairwise 1–2, 1–3, or 1–4 weighting list.

The *special* keyword is invoked at the end of the `delete_bonds` operation, after (optional) removal. It re-computes the pairwise 1–2, 1–3, 1–4 weighting list. The weighting list computation treats turned-off bonds the same as turned-on. Thus, turned-off bonds must be removed if you wish to change the weighting list.

Note that the choice of *remove* and *special* options affects how 1–2, 1–3, 1–4 pairwise interactions will be computed across bonds that have been modified by the `delete_bonds` command.

## 16.21.4 Restrictions

This command requires inter-processor communication to acquire ghost atoms, to coordinate the deleting of bonds, angles, etc. between atoms shared by multiple processors. This means that your system must be ready to perform a simulation before using this command (force fields setup, atom masses set, etc.). Just as would be needed to run dynamics, the force field you define should define a cutoff (e.g., through a *pair\_style* command) which is long enough for a processor to acquire the ghost atoms its needs to compute bond, angle, etc. interactions.

If deleted bonds (or angles, etc.) are removed but the 1–2, 1–3, and 1–4 weighting list is not recomputed, this can cause a later *fix shake* command to fail due to an atom's bonds being inconsistent with the weighting list. This should only happen if the group used in the `fix` command includes both atoms in the bond, in which case you probably should be recomputing the weighting list.

## 16.21.5 Related commands

*neigh\_modify* `exclude`, *special\_bonds*, *fix shake*

## 16.21.6 Default

none

## 16.22 dielectric command

### 16.22.1 Syntax

<code>dielectric</code> value
-------------------------------

- value = dielectric constant

## 16.22.2 Examples

```
dielectric 2.0
```

## 16.22.3 Description

Set the dielectric constant for Coulombic interactions (pairwise and long-range) to this value. The constant is unitless, since it is used to reduce the strength of the interactions. The value is used in the denominator of the formulas for Coulombic interactions (e.g., a value of 4.0 reduces the Coulombic interactions to 25% of their default strength). See the *pair\_style* command for more details.

## 16.22.4 Restrictions

none

## 16.22.5 Related commands

*pair\_style*

## 16.22.6 Default

```
dielectric 1.0
```

## 16.23 dihedral\_coeff command

### 16.23.1 Syntax

```
dihedral_coeff N args
```

- N = numeric dihedral type (see asterisk form below) or alphanumeric type label
- args = coefficients for one or more dihedral types

### 16.23.2 Examples

```
dihedral_coeff 1 80.0 1 3  
dihedral_coeff * 80.0 1 3 0.5  
dihedral_coeff 2* 80.0 1 3 0.5  
  
labelmap dihedral 1 backbone  
dihedral_coeff backbone 80.0 1 3
```

### 16.23.3 Description

Specify the dihedral force field coefficients for one or more dihedral types. The number and meaning of the coefficients depends on the dihedral style. Dihedral coefficients can also be set in the data file read by the `read_data` command or in a restart file.

$N$  can be specified in one of two ways. An explicit numeric value can be used, as in the first example above. Or  $N$  can be an alphanumeric type label, which is a string defined by the `labelmap` command or in a corresponding section of a data file read by the `read_data` command.

For numeric values only, a wild-card asterisk can be used to set the coefficients for multiple dihedral types. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If  $N$  is the number of dihedral types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $n$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive).

Note that using a `dihedral_coeff` command can override a previous setting for the same dihedral type. For example, these commands set the coeffs for all dihedral types, then overwrite the coeffs for just dihedral type 2:

```
dihedral_coeff * 80.0 1 3
dihedral_coeff 2 200.0 1 3
```

A line in a data file that specifies dihedral coefficients uses the exact same format as the arguments of the `dihedral_coeff` command in an input script, except that wild-card asterisks should not be used since coefficients for all  $N$  types must be listed in the file. For example, under the “Dihedral Coeffs” section of a data file, the line that corresponds to the first example above would be listed as

```
1 80.0 1 3
```

The `dihedral_style class2` is an exception to this rule, in that an additional argument is used in the input script to allow specification of the cross-term coefficients. See its doc page for details.

---

**Note:** When comparing the formulas and coefficients for various LAMMPS dihedral styles with dihedral equations defined by other force fields, note that some force field implementations divide/multiply the energy prefactor  $K$  by the multiple number of torsions that contain the  $J$ - $K$  bond in an  $I$ - $J$ - $K$ - $L$  torsion. LAMMPS does not do this (i.e., the listed dihedral equation applies to each individual dihedral). Thus, you need to define  $K$  appropriately to account for this difference, if necessary.

---

The list of all dihedral styles defined in LAMMPS is given on the `dihedral_style` doc page. They are also listed in more compact form on the `Commands dihedral` doc page.

On either of those pages, click on the style to display the formula it computes and its coefficients as specified by the associated `dihedral_coeff` command.

---

## 16.23.4 Restrictions

This command must come after the simulation box is defined by a *read\_data*, *read\_restart*, or *create\_box* command.

A dihedral style must be defined before any dihedral coefficients are set, either in the input script or in a data file.

## 16.23.5 Related commands

*dihedral\_style*

## 16.23.6 Default

none

# 16.24 dihedral\_style command

## 16.24.1 Syntax

```
dihedral_style style
```

- style = *none* or *zero* or *hybrid* or *charmm* or *charmmfsw* or *class2* or *osine/shift/exp* or *fourier* or *harmonic* or *helix* or *multi/harmonic* or *nharmonic* or *opls* or *spherical* or *table* or *table/cut*

## 16.24.2 Examples

```
dihedral_style harmonic
dihedral_style multi/harmonic
dihedral_style hybrid harmonic charmm
```

## 16.24.3 Description

Set the formula(s) LAMMPS uses to compute dihedral interactions between quadruplets of atoms, which remain in force for the duration of the simulation. The list of dihedral quadruplets is read in by a *read\_data* or *read\_restart* command from a data or restart file.

Hybrid models where dihedrals are computed using different dihedral potentials can be setup using the *hybrid* dihedral style.

The coefficients associated with a dihedral style can be specified in a data or restart file or via the *dihedral\_coeff* command.

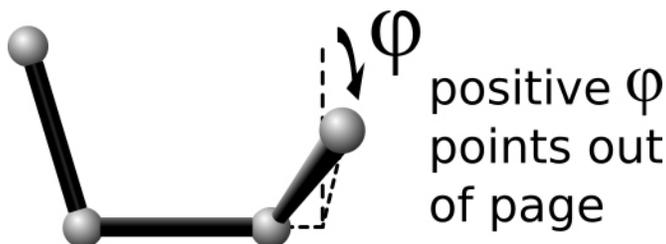
All dihedral potentials store their coefficient data in binary restart files which means *dihedral\_style* and *dihedral\_coeff* commands do not need to be re-specified in an input script that restarts a simulation. See the *read\_restart* command for details on how to do this. The one exception is that *dihedral\_style hybrid* only stores the list of sub-styles in the restart file; dihedral coefficients need to be re-specified.

---

**Note:** When both a dihedral and pair style is defined, the *special\_bonds* command often needs to be used to turn off (or weight) the pairwise interaction that would otherwise exist between four bonded atoms.

---

In the formulas listed for each dihedral style,  $\phi$  is the torsional angle defined by the quadruplet of atoms. This angle has a sign convention as shown in this diagram:



where the  $I, J, K, L$  ordering of the four atoms that define the dihedral is from left to right.

This sign convention effects several of the dihedral styles listed below (e.g., charmm, helix) in the sense that the energy formula depends on the sign of  $\phi$ , which may be reflected in the value of the coefficients you specify.

---

**Note:** When comparing the formulas and coefficients for various LAMMPS dihedral styles with dihedral equations defined by other force fields, note that some force field implementations divide/multiply the energy prefactor  $K$  by the multiple number of torsions that contain the  $J-K$  bond in an  $I-J-K-L$  torsion. LAMMPS does not do this (i.e., the listed dihedral equation applies to each individual dihedral). Thus, you need to define  $K$  appropriately via the `dihedral_coeff` command to account for this difference if necessary.

---

Here is an alphabetic list of dihedral styles defined in LAMMPS. Click on the style to display the formula it computes and coefficients specified by the associated `dihedral_coeff` command.

Click on the style to display the formula it computes, any additional arguments specified in the `dihedral_style` command, and coefficients specified by the associated `dihedral_coeff` command.

There are also additional accelerated pair styles included in the LAMMPS distribution for faster performance on CPUs, GPUs, and KNLs. The individual style names on the [Commands dihedral](#) page are followed by one or more of (g,i,k,o,t) to indicate which accelerated styles exist.

- *none* - turn off dihedral interactions
- *zero* - topology but no interactions
- *hybrid* - define multiple styles of dihedral interactions
- *charmm* - CHARMM dihedral
- *charmmfsw* - CHARMM dihedral with force switching
- *class2* - COMPASS (class 2) dihedral
- *cosine/shift/exp* - dihedral with exponential in spring constant
- *fourier* - dihedral with multiple cosine terms
- *harmonic* - harmonic dihedral
- *helix* - helix dihedral
- *multi/harmonic* - dihedral with 5 harmonic terms
- *nharmonic* - same as multi-harmonic with N terms
- *opls* - OPLS dihedral
- *quadratic* - dihedral with quadratic term in angle
- *spherical* - dihedral which includes angle terms to avoid singularities

- *table* - tabulated dihedral
  - *table/cut* - tabulated dihedral with analytic cutoff
- 

## 16.24.4 Restrictions

Dihedral styles can only be set for atom styles that allow dihedrals to be defined.

Most dihedral styles are part of the MOLECULE package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info. The doc pages for individual dihedral potentials tell if it is part of a package.

## 16.24.5 Related commands

*dihedral\_coeff*

## 16.24.6 Default

dihedral\_style none

# 16.25 dimension command

## 16.25.1 Syntax

```
dimension N
```

- N = 2 or 3

## 16.25.2 Examples

```
dimension 2
```

## 16.25.3 Description

Set the dimensionality of the simulation. By default LAMMPS runs 3d simulations. To run a 2d simulation, this command should be used prior to setting up a simulation box via the *create\_box* or *read\_data* commands. Restart files also store this setting.

See the discussion on the *Howto 2d* page for additional instructions on how to run 2d simulations.

---

**Note:** Some models in LAMMPS treat particles as finite-size spheres or ellipsoids, as opposed to point particles. In 2d, the particles will still be spheres or ellipsoids, not circular disks or ellipses, meaning their moment of inertia will be the same as in 3d.

---



## 16.25.4 Restrictions

This command must be used before the simulation box is defined by a *read\_data* or *create\_box* command.

## 16.25.5 Related commands

*fix enforce2d*

## 16.25.6 Default

```
dimension 3
```

# 16.26 displace\_atoms command

## 16.26.1 Syntax

```
displace_atoms group-ID style args keyword value ...
```

- group-ID = ID of group of atoms to displace
- style = *move* or *ramp* or *random* or *rotate*
  - move* args = delx dely delz  
delx,dely,delz = distance to displace in each dimension (distance units)  
any of delx,dely,delz can be a variable (see below)
  - ramp* args = ddim dlo dhi dim clo chi  
ddim = x or y or z  
dlo,dhi = displacement distance between dlo and dhi (distance units)  
dim = x or y or z  
clo,chi = lower and upper bound of domain to displace (distance units)
  - random* args = dx dy dz seed  
dx,dy,dz = random displacement magnitude in each dimension (distance units)  
seed = random # seed (positive integer)
  - rotate* args = Px Py Pz Rx Ry Rz theta  
Px,Py,Pz = origin point of axis of rotation (distance units)  
Rx,Ry,Rz = axis of rotation vector  
theta = angle of rotation (degrees)
- zero or more keyword/value pairs may be appended
  - keyword = *units*  
units value = *box* or *lattice*

## 16.26.2 Examples

```
displace_atoms top move 0 -5 0 units box
displace_atoms flow ramp x 0.0 5.0 y 2.0 20.5
```

## 16.26.3 Description

Displace a group of atoms. This can be used to move atoms a large distance before beginning a simulation or to randomize atoms initially on a lattice. For example, in a shear simulation, an initial strain can be imposed on the system. Or two groups of atoms can be brought into closer proximity.

The *move* style displaces the group of atoms by the specified 3d displacement vector. Any of the three quantities defining the vector components can be specified as an equal-style or atom-style *variable*. If the value is a variable, it should be specified as *v\_name*, where name is the variable name. In this case, the variable will be evaluated, and its value(s) used for the displacement(s). The scale factor implied by the *units* keyword will also be applied to the variable result.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Atom-style variables can specify the same formulas as equal-style variables but can also include per-atom values, such as atom coordinates or per-atom values read from a file. Note that if the variable references other *compute* or *fix* commands, those values must be up-to-date for the current timestep. See the “Variable Accuracy” section of the *variable* doc page for more details.

The *ramp* style displaces atoms a variable amount in one dimension depending on the atom’s coordinate in a (possibly) different dimension. For example, the second example command displaces atoms in the *x*-direction an amount between 0.0 and 5.0 distance units. Each atom’s displacement depends on the fractional distance its *y* coordinate is between 2.0 and 20.5. Atoms with *y*-coordinates outside those bounds will be moved the minimum (0.0) or maximum (5.0) amount.

The *random* style independently moves each atom in the group by a random displacement, uniformly sampled from a value between  $-dx$  and  $+dx$  in the *x* dimension, and similarly for *y* and *z*. Random numbers are used in such a way that the displacement of a particular atom is the same, regardless of how many processors are being used.

The *rotate* style rotates each atom in the group by the angle *theta* around a rotation axis  $R = (R_x, R_y, R_z)$  that goes through a point  $P = (P_x, P_y, P_z)$ . The direction of rotation for the atoms around the rotation axis is consistent with the right-hand rule: if your right-hand thumb points along *R*, then your fingers wrap around the axis in the direction of positive *theta*.

If the defined *atom\_style* assigns an orientation to each atom (*atom styles* ellipsoid, line, tri, body), then that property is also updated appropriately to correspond to the atom’s rotation.

Distance units for displacements and the origin point of the *rotate* style are determined by the setting of *box* or *lattice* for the *units* keyword. *Box* means distance units as defined by the *units* command (e.g., Å for *real* or *metal* units). *Lattice* means distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing.

---

**Note:** Care should be taken not to move atoms on top of other atoms. After the move, atoms are remapped into the periodic simulation box if needed, and any shrink-wrap boundary conditions (see the *boundary* command) are enforced which may change the box size. Other than this effect, this command does not change the size or shape of the simulation box. See the *change\_box* command if that effect is desired.

---

**Note:** Atoms can be moved arbitrarily long distances by this command. If the simulation box is non-periodic and shrink-wrapped (see the *boundary* command), this can change its size or shape. This is not a problem, except that the mapping of processors to the simulation box is not changed by this command from its initial 3d configuration; see the

*processors* command. Thus, if the box size/shape changes dramatically, the mapping of processors to the simulation box may not end up as optimal as the initial mapping attempted to be.

---

## 16.26.4 Restrictions

For a 2d simulation, only rotations around the a vector parallel to the  $z$ -axis are allowed.

## 16.26.5 Related commands

*lattice*, *change\_box*, *fix move*

## 16.26.6 Default

The option defaults are units = lattice.

## 16.27 dump command

### 16.28 dump vtk command

### 16.29 dump h5md command

### 16.30 dump molfile command

### 16.31 dump netcdf command

### 16.32 dump image command

### 16.33 dump movie command

### 16.34 dump atom/adios command

### 16.35 dump custom/adios command

### 16.36 dump cfg/uef command

#### 16.36.1 Syntax

<code>dump ID group-ID style N file args</code>
---

- ID = user-assigned name for the dump
- group-ID = ID of the group of atoms to be dumped
- style = *atom* or *atom/adios* or *atom/gz* or *atom/zstd* or *atom/mpiio* or *cfg* or *cfg/gz* or *cfg/zstd* or *cfg/mpiio* or *cfg/uef* or *custom* or *custom/gz* or *custom/zstd* or *custom/mpiio* or *custom/adios* or *dcd* or *h5md* or *image* or *local* or *local/gz* or *local/zstd* or *molfile* or *movie* or *netcdf* or *netcdf/mpiio* or *vtk* or *xtc* or *xyz* or *xyz/gz* or *xyz/zstd* or *xyz/mpiio* or *yaml*
- N = dump on timesteps which are multiples of N
- file = name of file to write dump info to
- args = list of arguments for a particular style

*atom* args = none

*atom/adios* args = none, discussed on [dump atom/adios](#) page

*atom/gz* args = none

*atom/zstd* args = none

*atom/mpiio* args = none

*cfg* args = same as *custom* args, see below

*cfg/gz* args = same as *custom* args, see below

*cfg/zstd* args = same as *custom* args, see below

*cfg/mpiio* args = same as *custom* args, see below

*cfg/uef* args = same as *custom* args, discussed on [dump cfg/uef](#) page

*custom*, *custom/gz*, *custom/zstd*, *custom/mpiio* args = see below

*custom/adios* args = same as *custom* args, discussed on [dump custom/adios](#) page

*dcd* args = none

*h5md* args = discussed on [dump h5md](#) page

*image* args = discussed on [dump image](#) page

*local*, *local/gz*, *local/zstd* args = see below

*molfile* args = discussed on [dump molfile](#) page

*movie* args = discussed on [dump image](#) page

*netcdf* args = discussed on [dump netcdf](#) page

*netcdf/mpiio* args = discussed on [dump netcdf](#) page

*vtk* args = same as *custom* args, see below, also [dump vtk](#) page

*xtc* args = none

*xyz* args = none

*xyz/gz* args = none

*xyz/zstd* args = none

*xyz/mpiio* args = none

*yaml* args = same as *custom* args, see below

- *custom* or *custom/gz* or *custom/zstd* or *custom/mpiio* or *cfg* or *cfg/gz* or *cfg/zstd* or *cfg/mpiio* or *cfg/uef* or *netcdf* or *netcdf/mpiio* or *yaml* args = list of atom attributes

```
possible attributes = id, mol, proc, procp1, type, element, mass,
                    x, y, z, xs, ys, zs, xu, yu, zu,
                    xsu, ysu, zsu, ix, iy, iz,
                    vx, vy, vz, fx, fy, fz,
                    q, mux, muy, muz, mu,
                    radius, diameter, omegax, omegay, omegaz,
                    angmomx, angmomy, angmomz, tqx, tqy, tqz,
                    c_ID, c_ID[I], f_ID, f_ID[I], v_name,
                    i_name, d_name, i2_name[I], d2_name[I]
```

id = atom ID

mol = molecule ID

```

proc = ID of processor that owns atom
procp1 = ID+1 of processor that owns atom
type = atom type
element = name of atom element, as defined by dump_modify command
mass = atom mass
x,y,z = unscaled atom coordinates
xs,ys,zs = scaled atom coordinates
xu,yu,zu = unwrapped atom coordinates
xsu,ysu,zsu = scaled unwrapped atom coordinates
ix,iy,iz = box image that the atom is in
vx,vy,vz = atom velocities
fx,fy,fz = forces on atoms
q = atom charge
mux,muy,muz = orientation of dipole moment of atom
mu = magnitude of dipole moment of atom
radius,diameter = radius,diameter of spherical particle
omegax,omegay,omegaz = angular velocity of spherical particle
angmomx,angmomy,angmomz = angular momentum of aspherical particle
tqx,tqy,tqz = torque on finite-size particles
c_ID = per-atom vector calculated by a compute with ID
c_ID[I] = Ith column of per-atom array calculated by a compute with ID, I can
↔include wildcard (see below)
f_ID = per-atom vector calculated by a fix with ID
f_ID[I] = Ith column of per-atom array calculated by a fix with ID, I can include
↔wildcard (see below)
v_name = per-atom vector calculated by an atom-style variable with name
i_name = custom integer vector with name
d_name = custom floating point vector with name
i2_name[I] = Ith column of custom integer array with name, I can include wildcard
↔(see below)
d2_name[I] = Ith column of custom floating point vector with name, I can include
↔wildcard (see below)

```

- *local* or *local/gz* or *local/zstd* args = list of local attributes

```

possible attributes = index, c_ID, c_ID[I], f_ID, f_ID[I]
index = enumeration of local values
c_ID = local vector calculated by a compute with ID
c_ID[I] = Ith column of local array calculated by a compute with ID, I can
↔include wildcard (see below)
f_ID = local vector calculated by a fix with ID
f_ID[I] = Ith column of local array calculated by a fix with ID, I can include
↔wildcard (see below)

```

## 16.36.2 Examples

```
dump myDump all atom 100 dump.lampstrj
dump myDump all atom/mpiio 100 dump.atom.mpio
dump myDump all atom/gz 100 dump.atom.gz
dump myDump all atom/zstd 100 dump.atom.zst
dump 2 subgroup atom 50 dump.run.bin
dump 2 subgroup atom/mpiio 50 dump.run.mpio.bin
dump 4a all custom 100 dump.myforce.* id type x y vx fx
dump 4a all custom 100 dump.myvel.lammpsbin id type x y z vx vy vz
dump 4b flow custom 100 dump.%myforce id type c_myF[3] v_ke
dump 4b flow custom 100 dump.%myforce id type c_myF[*] v_ke
dump 2 inner cfg 10 dump.snap.*.cfg mass type xs ys zs vx vy vz
dump snap all cfg 100 dump.config.*.cfg mass type xs ys zs id type c_Stress[2]
dump 1 all xtc 1000 file.xtc
```

## 16.36.3 Description

Dump a snapshot of atom quantities to one or more files once every  $N$  timesteps in one of several styles. The *image* and *movie* styles are the exception: the *image* style renders a JPG, PNG, or PPM image file of the atom configuration every  $N$  timesteps while the *movie* style combines and compresses them into a movie file; both are discussed in detail on the [dump image](#) page. The timesteps on which dump output is written can also be controlled by a variable. See the [dump\\_modify every](#) command.

Only information for atoms in the specified group is dumped. The [dump\\_modify thresh and region and refresh](#) commands can also alter what atoms are included. Not all styles support these options; see details on the [dump\\_modify](#) doc page.

As described below, the filename determines the kind of output (text or binary or gzipped, one big file or one per timestep, one big file or multiple smaller files).

---

**Note:** Because periodic boundary conditions are enforced only on timesteps when neighbor lists are rebuilt, the coordinates of an atom written to a dump file may be slightly outside the simulation box. Re-neighbor timesteps will not typically coincide with the timesteps dump snapshots are written. See the [dump\\_modify pbc](#) command if you wish to force coordinates to be strictly inside the simulation box.

---

**Note:** Unless the [dump\\_modify sort](#) option is invoked, the lines of atom information written to dump files (typically one line per atom) will be in an indeterminate order for each snapshot. This is even true when running on a single processor, if the [atom\\_modify sort](#) option is on, which it is by default. In this case atoms are re-ordered periodically during a simulation, due to spatial sorting. It is also true when running in parallel, because data for a single snapshot is collected from multiple processors, each of which owns a subset of the atoms.

---

For the *atom*, *custom*, *cfg*, and *local* styles, sorting is off by default. For the *dcd*, *xtc*, *xyz*, and *molfile* styles, sorting by atom ID is on by default. See the [dump\\_modify](#) page for details.

The *atom/gz*, *cfg/gz*, *custom/gz*, *local/gz*, and *xyz/gz* styles are identical in command syntax to the corresponding styles without “gz”, however, they generate compressed files using the zlib library. Thus the filename suffix “.gz” is mandatory. This is an alternative approach to writing compressed files via a pipe, as done by the regular dump styles, which may be required on clusters where the interface to the high-speed network disallows using the `fork()` library call (which is needed for a pipe). For the remainder of this page, you should thus consider the *atom* and *atom/gz* styles (etc.) to be inter-changeable, with the exception of the required filename suffix.

Similarly, the *atom/zstd*, *cfg/zstd*, *custom/zstd*, *local/zstd*, and *xyz/zstd* styles are identical to the *gz* styles, but use the Zstd compression library instead and require the “.zst” suffix. See the [dump\\_modify](#) page for details on how to control the compression level in both variants.

As explained below, the *atom/mpio*, *cfg/mpio*, *custom/mpio*, and *xyz/mpio* styles are identical in command syntax and in the format of the dump files they create, to the corresponding styles without “mpio”, except the single dump file they produce is written in parallel via the MPI-IO library. For the remainder of this page, you should thus consider the *atom* and *atom/mpio* styles (etc.) to be inter-changeable. The one exception is how the filename is specified for the MPI-IO styles, as explained below.

**Warning:** The MPIIO package is currently unmaintained and has become unreliable. Use with caution.

The precision of values output to text-based dump files can be controlled by the [dump\\_modify format](#) command and its options.

The *style* keyword determines what atom quantities are written to the file and in what format. Settings made via the [dump\\_modify](#) command can also alter the format of individual values and the file itself.

The *atom*, *local*, and *custom* styles create files in a simple text format that is self-explanatory when viewing a dump file. Some of the LAMMPS post-processing tools described on the [Tools](#) doc page, including [Pizza.py](#), work with this format, as does the [rerun](#) command.

For post-processing purposes the *atom*, *local*, and *custom* text files are self-describing in the following sense.

The dimensions of the simulation box are included in each snapshot. For an orthogonal simulation box this information is formatted as:

```
ITEM: BOX BOUNDS xx yy zz
xlo xhi
ylo yhi
zlo zhi
```

where *xlo,xhi* are the maximum extents of the simulation box in the *x*-dimension, and similarly for *y* and *z*. The “*xx yy zz*” terms are six characters that encode the style of boundary for each of the six simulation box boundaries (*xlo,xhi*; *ylo,yhi*; and *zlo,zhi*). Each of the six characters is one of *p* (periodic), *f* (fixed), *s* (shrink wrap), or *m* (shrink wrapped with a minimum value). See the [boundary](#) command for details.

For triclinic simulation boxes (non-orthogonal), an orthogonal bounding box which encloses the triclinic simulation box is output, along with the three tilt factors (*xy, xz, yz*) of the triclinic box, formatted as follows:

```
ITEM: BOX BOUNDS xy xz yz xx yy zz
xlo_bound xhi_bound xy
ylo_bound yhi_bound xz
zlo_bound zhi_bound yz
```

The presence of the text “*xy xz yz*” in the ITEM line indicates that the three tilt factors will be included on each of the three following lines. This bounding box is convenient for many visualization programs. The meaning of the six character flags for “*xx yy zz*” is the same as above.

Note that the first two numbers on each line are now *xlo\_bound* instead of *xlo*, etc. because they represent a bounding box. See the [Howto triclinic](#) page for a geometric description of triclinic boxes, as defined by LAMMPS, simple formulas for how the six bounding box extents (*xlo\_bound*, *xhi\_bound*, etc.) are calculated from the triclinic parameters, and how to transform those parameters to and from other commonly used triclinic representations.

The “ITEM: ATOMS” line in each snapshot lists column descriptors for the per-atom lines that follow. For example, the descriptors would be “id type xs ys zs” for the default *atom* style, and would be the atom attributes you specify in the dump command for the *custom* style.

For style *atom*, atom coordinates are written to the file, along with the atom ID and atom type. By default, atom coords are written in a scaled format (from 0 to 1). That is, an *x* value of 0.25 means the atom is at a location 1/4 of the distance from *xlo* to *xhi* of the box boundaries. The format can be changed to unscaled coords via the *dump\_modify* settings. Image flags can also be added for each atom via *dump\_modify*.

Style *custom* allows you to specify a list of atom attributes to be written to the dump file for each atom. Possible attributes are listed above and will appear in the order specified. You cannot specify a quantity that is not defined for a particular simulation—such as *q* for atom style *bond*, since that atom style does not assign charges. Dumps occur at the very end of a timestep, so atom attributes will include effects due to fixes that are applied during the timestep. An explanation of the possible dump custom attributes is given below.

For style *local*, local output generated by *computes* and *fixes* is used to generate lines of output that is written to the dump file. This local data is typically calculated by each processor based on the atoms it owns, but there may be zero or more entities per atom (e.g., a list of bond distances). An explanation of the possible dump local attributes is given below. Note that by using input from the *compute property/local* command with *dump local*, it is possible to generate information on bonds, angles, etc. that can be cut and pasted directly into a data file read by the *read\_data* command.

Style *cfg* has the same command syntax as style *custom* and writes extended CFG format files, as used by the AtomEye visualization package. Since the extended CFG format uses a single snapshot of the system per file, a wildcard “\*” must be included in the filename, as discussed below. The list of atom attributes for style *cfg* must begin with either “mass type xs ys zs” or “mass type xsu ysu zsu” since these quantities are needed to write the CFG files in the appropriate format (though the “mass” and “type” fields do not appear explicitly in the file). Any remaining attributes will be stored as “auxiliary properties” in the CFG files. Note that you will typically want to use the *dump\_modify element* command with CFG-formatted files, to associate element names with atom types, so that AtomEye can render atoms appropriately. When unwrapped coordinates *xsu*, *ysu*, and *zsu* are requested, the nominal AtomEye periodic cell dimensions are expanded by a large factor UNWRAPEXPAND = 10.0, which ensures atoms that are displayed correctly for up to UNWRAPEXPAND/2 periodic boundary crossings in any direction. Beyond this, AtomEye will rewrap the unwrapped coordinates. The expansion causes the atoms to be drawn farther away from the viewer, but it is easy to zoom the atoms closer, and the interatomic distances are unaffected.

The *dcd* style writes DCD files, a standard atomic trajectory format used by the CHARMM, NAMD, and XPLOR molecular dynamics packages. DCD files are binary and thus may not be portable to different machines. The number of atoms per snapshot cannot change with the *dcd* style. The *unwrap* option of the *dump\_modify* command allows DCD coordinates to be written “unwrapped” by the image flags for each atom. Unwrapped means that if the atom has passed through a periodic boundary one or more times, the value is printed for what the coordinate would be if it had not been wrapped back into the periodic box. Note that these coordinates may thus be far outside the box size stored with the snapshot.

The *xtc* style writes XTC files, a compressed trajectory format used by the GROMACS molecular dynamics package, and described [here](#). The precision used in XTC files can be adjusted via the *dump\_modify* command. The default value of 1000 means that coordinates are stored to 1/1000 nanometer accuracy. XTC files are portable binary files written in the NFS XDR data format, so that any machine which supports XDR should be able to read them. The number of atoms per snapshot cannot change with the *xtc* style. The *unwrap* option of the *dump\_modify* command allows XTC coordinates to be written “unwrapped” by the image flags for each atom. Unwrapped means that if the atom has passed through a periodic boundary one or more times, the value is printed for what the coordinate would be if it had not been wrapped back into the periodic box. Note that these coordinates may thus be far outside the box size stored with the snapshot.

The *xyz* style writes XYZ files, which is a simple text-based coordinate format that many codes can read. Specifically it has a line with the number of atoms, then a comment line that is usually ignored followed by one line per atom with the atom type and the *x*-, *y*-, and *z*-coordinate of that atom. You can use the *dump\_modify element* option to change the output from using the (numerical) atom type to an element name (or some other label). This will help many visualization programs to guess bonds and colors.



New in version 4May2022.

Dump style *yaml* has the same command syntax as style *custom* and writes YAML format files that can be easily parsed by a variety of data processing tools and programming languages. Each timestep will be written as a YAML “document” (i.e., starts with “—” and ends with “...” ). The style supports writing one file per timestep through the “\*” wildcard but not multi-processor outputs with the “%” token in the filename. In addition to per-atom data, *thermo* data can be included in the *yaml* style dump file using the *dump\_modify thermo yes*. The data included in the dump file uses the “thermo” tag and is otherwise identical to data specified by the *thermo\_style* command.

Below is an example for a YAML format dump created by the following commands.

```
dump out all yaml 100 dump.yaml id type x y z vx vy vz ix iy iz
dump_modify out time yes units yes thermo yes format 1 %5d format "% 10.6e"
```

The tags “time”, “units”, and “thermo” are optional and enabled by the *dump\_modify* command. The list under the “box” tag has three lines for orthogonal boxes and four lines for triclinic boxes, where the first three are the box boundaries and the fourth the three tilt factors (*xy*, *xz*, *yz*). The “thermo” data follows the format of the *yaml* thermo style. The “keywords” tag lists the per-atom properties contained in the “data” columns, which contain a list with one line per atom. The keywords may be renamed using the *dump\_modify* command same as for the *custom* dump style.

```
---
creator: LAMMPS
timestep: 0
units: lj
time: 0
natoms: 4000
boundary: [ p, p, p, p, p, p, ]
thermo:
  - keywords: [ Step, Temp, E_pair, E_mol, TotEng, Press, ]
  - data: [ 0, 0, -27093.472213010766, 0, 0, 0, ]
box:
  - [ 0, 16.795961913825074 ]
  - [ 0, 16.795961913825074 ]
  - [ 0, 16.795961913825074 ]
  - [ 0, 0, 0 ]
keywords: [ id, type, x, y, z, vx, vy, vz, ix, iy, iz, ]
data:
  - [ 1, 1, 0.000000e+00, 0.000000e+00, 0.000000e+00, -1.841579e-01, -9.
↪710036e-01, -2.934617e+00, 0, 0, 0, ]
  - [ 2, 1, 8.397981e-01, 8.397981e-01, 0.000000e+00, -1.799591e+00, 2.
↪127197e+00, 2.298572e+00, 0, 0, 0, ]
  - [ 3, 1, 8.397981e-01, 0.000000e+00, 8.397981e-01, -1.807682e+00, -9.
↪585130e-01, 1.605884e+00, 0, 0, 0, ]

  [...]
...
---
timestep: 100
units: lj
time: 0.5

  [...]
...
---
```

Note that *atom*, *custom*, *dcd*, *xtc*, and *xyz* style dump files can be read directly by [VMD](#), a popular molecular viewing program.

---

Dumps are performed on timesteps that are a multiple of  $N$  (including timestep 0) and on the last timestep of a minimization if the minimization converges. Note that this means a dump will not be performed on the initial timestep after the dump command is invoked, if the current timestep is not a multiple of  $N$ . This behavior can be changed via the *dump\_modify first* command, which can also be useful if the dump command is invoked after a minimization ended on an arbitrary timestep.

The value of  $N$  can be changed between runs by using the *dump\_modify every* command (not allowed for *dcd* style). The *dump\_modify every* command also allows a variable to be used to determine the sequence of timesteps on which dump files are written. In this mode a dump on the first timestep of a run will also not be written unless the *dump\_modify first* command is used.

If you instead want to dump snapshots based on simulation time (in time units of the *units command* command), the *dump\_modify every/time* command can be used. This can be useful when the timestep size varies during a simulation run, e.g. by use of the *fix dt/reset* command.

The specified filename determines how the dump file(s) is written. The default is to write one large text file, which is opened when the dump command is invoked and closed when an *undump* command is used or when LAMMPS exits. For the *dcd* and *xtc* styles, this is a single large binary file.

Dump filenames can contain two wildcard characters. If a “\*” character appears in the filename, then one file per snapshot is written and the “\*” character is replaced with the timestep value. For example, *tmp.dump.\** becomes *tmp.dump.0*, *tmp.dump.10000*, *tmp.dump.20000*, etc. This option is not available for the *dcd* and *xtc* styles. Note that the *dump\_modify pad* command can be used to insure all timestep numbers are the same length (e.g., 00010), which can make it easier to read a series of dump files in order with some post-processing tools.

If a “%” character appears in the filename, then each of  $P$  processors writes a portion of the dump file, and the “%” character is replaced with the processor ID from 0 to  $P - 1$ . For example, *tmp.dump.%* becomes *tmp.dump.0*, *tmp.dump.1*, ... *tmp.dump.math:P-1*, etc. This creates smaller files and can be a fast mode of output on parallel machines that support parallel I/O for output. This option is **not** available for the *dcd*, *xtc*, *xyz*, and *yaml* styles.

By default,  $P$  is the the number of processors, meaning one file per processor, but  $P$  can be set to a smaller value via the *nfile* or *fileper* keywords of the *dump\_modify* command. These options can be the most efficient way of writing out dump files when running on large numbers of processors.

Note that using the “\*” and “%” characters together can produce a large number of small dump files!

For the *atom/mpii*, *cfg/mpii*, *custom/mpii*, and *xyz/mpii* styles, a single dump file is written in parallel via the MPI-IO library, which is part of the MPI standard for versions 2.0 and above. Using MPI-IO requires two steps. First, build LAMMPS with its MPIIO package installed, viz.,

```
make yes-mpii    # installs the MPIIO package
make mpi        # build LAMMPS for your platform
```

Second, use a dump filename which contains “.mpii”. Note that it does not have to end in “.mpii”, just contain those characters. Unlike MPI-IO restart files, which must be both written and read using MPI-IO, the dump files produced by these MPI-IO styles are identical in format to the files produced by their non-MPI-IO style counterparts. This means you can write a dump file using MPI-IO and use the *read\_dump* command or perform other post-processing, just as if the dump file was not written using MPI-IO.

**Warning:** The MPIIO package is currently unmaintained and has become unreliable. Use with caution.

Note that MPI-IO dump files are one large file which all processors write to. You thus cannot use the “%” wildcard character described above in the filename since that specifies generation of multiple files. You can use the “.bin” or

“.lammpsbin” suffix described below in an MPI-IO dump file; again this file will be written in parallel and have the same binary format as if it were written without MPI-IO.

If the filename ends with “.bin” or “.lammpsbin”, the dump file (or files, if “\*” or “%” is also used) is written in binary format. A binary dump file will be about the same size as a text version, but will typically write out much faster. Of course, when post-processing, you will need to convert it back to text format (see the *binary2txt tool*) or write your own code to read the binary file. The format of the binary file can be understood by looking at the `tools/binary2txt.cpp` file. This option is only available for the *atom* and *custom* styles.

If the filename ends with “.gz”, the dump file (or files, if “\*” or “%” is also used) is written in gzipped format. A gzipped dump file will be about 3× smaller than the text version, but will also take longer to write. This option is not available for the *dcd* and *xtc* styles.

Note that in the discussion which follows, for styles which can reference values from a compute or fix or custom atom property, like the *custom*, *cfg*, or *local* styles, the bracketed index *i* can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If *N* is the number of columns in the array, then an asterisk with no numeric values means all column indices from 1 to *N*. A leading asterisk means all indices from 1 to *n* (inclusive). A trailing asterisk means all indices from *m* to *N* (inclusive). A middle asterisk means all indices from *m* to *n* (inclusive).

Using a wildcard is the same as if the individual columns of the array had been listed one by one. For example, these two dump commands are equivalent, since the *compute stress/atom* command creates a per-atom array with six columns:

```
compute myPress all stress/atom NULL
dump 2 all custom 100 tmp.dump id myPress[*]
dump 2 all custom 100 tmp.dump id myPress[1] myPress[2] myPress[3] &
                                     myPress[4] myPress[5] myPress[6]
```

This section explains the local attributes that can be specified as part of the *local* style.

The *index* attribute can be used to generate an index number from 1 to *N* for each line written into the dump file, where *N* is the total number of local datums from all processors, or lines of output that will appear in the snapshot. Note that because data from different processors depend on what atoms they currently own, and atoms migrate between processor, there is no guarantee that the same index will be used for the same info (e.g., a particular bond) in successive snapshots.

The *c\_ID* and *c\_ID[I]* attributes allow local vectors or arrays calculated by a *compute* to be output. The ID in the attribute should be replaced by the actual ID of the compute that has been defined previously in the input script. See the *compute* command for details. There are computes for calculating local information such as indices, types, and energies for bonds and angles.

Note that computes which calculate global or per-atom quantities, as opposed to local quantities, cannot be output in a dump local command. Instead, global quantities can be output by the *thermo\_style custom* command, and per-atom quantities can be output by the dump custom command.

If *c\_ID* is used as an attribute, then the local vector calculated by the compute is printed. If *c\_ID[i]* is used, then *i* must be in the range from 1 – *M*, which will print the *i*th column of the local array with *M* columns calculated by the compute. See the discussion above for how *i* can be specified with a wildcard asterisk to effectively specify multiple values.

The *f\_ID* and *f\_ID[I]* attributes allow local vectors or arrays calculated by a *fix* to be output. The ID in the attribute should be replaced by the actual ID of the fix that has been defined previously in the input script.

If *f\_ID* is used as an attribute, then the local vector calculated by the fix is printed. If *f\_ID[i]* is used, then *i* must be in the range 1–*M*, which will print the *i*th column of the local with *M* columns calculated by the fix. See the discussion above for how *i* can be specified with a wildcard asterisk to effectively specify multiple values.

Here is an example of how to dump bond info for a system, including the distance and energy of each bond:

```
compute 1 all property/local batom1 batom2 btype
compute 2 all bond/local dist eng
dump 1 all local 1000 tmp.dump index c_1[1] c_1[2] c_1[3] c_2[1] c_2[2]
```

This section explains the atom attributes that can be specified as part of the *custom* and *cfg* styles.

The *id*, *mol*, *proc*, *procp1*, *type*, *element*, *mass*, *vx*, *vy*, *vz*, *fx*, *fy*, *fz*, *q* attributes are self-explanatory.

*Id* is the atom ID. *Mol* is the molecule ID, included in the data file for molecular systems. *Proc* is the ID of the processor (0 to  $N_{\text{procs}} - 1$ ) that currently owns the atom. *Procp1* is the proc ID+1, which can be convenient in place of a *type* attribute (1 to  $N_{\text{types}}$ ) for coloring atoms in a visualization program. *Type* is the atom type (1 to  $N_{\text{types}}$ ). *Element* is typically the chemical name of an element, which you must assign to each type via the *dump\_modify element* command. More generally, it can be any string you wish to associated with an atom type. *Mass* is the atom mass. The quantities *vx*, *vy*, *vz*, *fx*, *fy*, *fz*, and *q* are components of atom velocity and force and atomic charge.

There are several options for outputting atom coordinates. The *x*, *y*, and *z* attributes write atom coordinates “unscaled”, in the appropriate distance *units* (Å, σ, etc.). Use *xs*, *ys*, and *zs* if you want the coordinates “scaled” to the box size so that each value is 0.0 to 1.0. If the simulation box is triclinic (tilted), then all atom coords will still be between 0.0 and 1.0. The actual unscaled (*x*, *y*, *z*) coordinate is  $x_s a + y_s b + z_s c$ , where (*a*, *b*, *c*) are the non-orthogonal vectors of the simulation box edges, as discussed on the *Howto triclinic* page.

Use *xu*, *yu*, and *zu* if you want the coordinates “unwrapped” by the image flags for each atom. Unwrapped means that if the atom has passed through a periodic boundary one or more times, the value is printed for what the coordinate would be if it had not been wrapped back into the periodic box. Note that using *xu*, *yu*, and *zu* means that the coordinate values may be far outside the box bounds printed with the snapshot. Using *xsu*, *ysu*, and *zsu* is similar to using *xu*, *yu*, and *zu*, except that the unwrapped coordinates are scaled by the box size. Atoms that have passed through a periodic boundary will have the corresponding coordinate increased or decreased by 1.0.

The image flags can be printed directly using the *ix*, *iy*, and *iz* attributes. For periodic dimensions, they specify which image of the simulation box the atom is considered to be in. An image of 0 means it is inside the box as defined. A value of 2 means add 2 box lengths to get the true value. A value of -1 means subtract 1 box length to get the true value. LAMMPS updates these flags as atoms cross periodic boundaries during the simulation.

The *mux*, *muy*, and *muz* attributes are specific to dipolar systems defined with an atom style of *dipole*. They give the orientation of the atom’s point dipole moment. The *mu* attribute gives the magnitude of the atom’s dipole moment.

The *radius* and *diameter* attributes are specific to spherical particles that have a finite size, such as those defined with an atom style of *sphere*.

The *omegax*, *omegay*, and *omegaz* attributes are specific to finite-size spherical particles that have an angular velocity. Only certain atom styles, such as *sphere*, define this quantity.

The *angmomx*, *angmomy*, and *angmomz* attributes are specific to finite-size aspherical particles that have an angular momentum. Only the *ellipsoid* atom style defines this quantity.

The *txx*, *tqy*, and *tqz* attributes are for finite-size particles that can sustain a rotational torque due to interactions with other particles.

The *c\_ID* and *c\_ID[I]* attributes allow per-atom vectors or arrays calculated by a *compute* to be output. The ID in the attribute should be replaced by the actual ID of the compute that has been defined previously in the input script. See the *compute* command for details. There are computes for calculating the per-atom energy, stress, centro-symmetry parameter, and coordination number of individual atoms.

Note that computes which calculate global or local quantities, as opposed to per-atom quantities, cannot be output in a dump custom command. Instead, global quantities can be output by the *thermo\_style custom* command, and local quantities can be output by the dump local command.

If *c\_ID* is used as an attribute, then the per-atom vector calculated by the compute is printed. If *c\_ID[i]* is used, then *i* must be in the range from 1 to *M*, which will print the *i*th column of the per-atom array with *M* columns calculated by the compute. See the discussion above for how *i* can be specified with a wildcard asterisk to effectively specify multiple values.

The *f\_ID* and *f\_ID[I]* attributes allow vector or array per-atom quantities calculated by a *fix* to be output. The ID in the attribute should be replaced by the actual ID of the fix that has been defined previously in the input script. The *fix ave/atom* command is one that calculates per-atom quantities. Since it can time-average per-atom quantities produced by any *compute*, *fix*, or atom-style *variable*, this allows those time-averaged results to be written to a dump file.

If *f\_ID* is used as an attribute, then the per-atom vector calculated by the fix is printed. If *f\_ID[i]* is used, then *i* must be in the range from 1 to *M*, which will print the *i*th column of the per-atom array with *M* columns calculated by the fix. See the discussion above for how *i* can be specified with a wildcard asterisk to effectively specify multiple values.

The *v\_name* attribute allows per-atom vectors calculated by a *variable* to be output. The name in the attribute should be replaced by the actual name of the variable that has been defined previously in the input script. Only an atom-style variable can be referenced, since it is the only style that generates per-atom values. Variables of style *atom* can reference individual atom attributes, per-atom attributes, thermodynamic keywords, or invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of creating quantities to output to a dump file.

The *i\_name*, *d\_name*, *i2\_name*, *d2\_name* attributes refer to per-atom integer and floating-point vectors or arrays that have been added via the *fix property/atom* command. When that command is used specific names are given to each attribute which are the “name” portion of these keywords. For arrays *i2\_name* and *d2\_name*, the column of the array must also be included following the name in brackets (e.g., *d2\_xyz[i]*, *i2\_mySpin[i]*, where *i* is in the range from 1 to *M*, where *M* is the number of columns in the custom array). See the discussion above for how *i* can be specified with a wildcard asterisk to effectively specify multiple values.

See the [Modify](#) page for information on how to add new compute and fix styles to LAMMPS to calculate per-atom quantities which could then be output into dump files.

---

## 16.36.4 Restrictions

To write gzipped dump files, you must either compile LAMMPS with the `-DLAMMPS_GZIP` option or use the styles from the COMPRESS package. See the [Build settings](#) page for details.

While a dump command is active (i.e., has not been stopped by using the [undump command](#)), no commands may be used that will change the timestep (e.g., [reset timestep](#)). LAMMPS will terminate with an error otherwise.

The *atom/gz*, *cfg/gz*, *custom/gz*, and *xyz/gz* styles are part of the COMPRESS package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

The *atom/mpiio*, *cfg/mpiio*, *custom/mpiio*, and *xyz/mpiio* styles are part of the MPIIO package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

The *xtc*, *dcd*, and *yaml* styles are part of the EXTRA-DUMP package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 16.36.5 Related commands

*dump atom/adios*, *dump custom/adios*, *dump cfg/uef*, *dump h5md*, *dump image*, *dump molfile*, *dump\_modify*, *undump*, *write\_dump*

### 16.36.6 Default

The defaults for the *image* and *movie* styles are listed on the *dump image* page.

## 16.37 dump atom/adios command

## 16.38 dump custom/adios command

### 16.38.1 Syntax

```
dump ID group-ID atom/adios N file.bp
dump ID group-ID custom/adios N file.bp args
```

- ID = user-assigned name for the dump
- group-ID = ID of the group of atoms to be imaged
- adios = style of dump command (other styles *atom* or *cfg* or *dcd* or *xtc* or *xyz* or *local* or *custom* are discussed on the *dump* doc page)
- N = dump every this many timesteps
- file.bp = name of file/stream to write to
- args = same options as in *dump custom* command

### 16.38.2 Examples

```
dump adios1 all atom/adios 100 atoms.bp
dump 4a all custom/adios 100 dump_adios.bp id v_p x y z
dump 2 subgroup custom/adios 100 dump_adios.bp mass type xs ys zs vx vy vz
```

### 16.38.3 Description

Dump a snapshot of atom coordinates every  $N$  timesteps in the ADIOS-based “BP” file format, or using different I/O solutions in ADIOS, to a stream that can be read on-line by another program. ADIOS-BP files are binary, portable, and self-describing.

---

**Note:** To be able to use ADIOS, a file `adios2_config.xml` with specific configuration settings is expected in the current working directory. If the file is not present, LAMMPS will try to create a minimal default file. Please refer to the ADIOS documentation for details on how to adjust this file for optimal performance and desired features.

---

Use from `write_dump`:

It is possible to use these dump styles with the *write\_dump* command. In this case, the sub-intervals must not be set at all. The *write\_dump* command can be used to create a new file at each individual dump.

```
dump 4      all atom/adios 100 dump.bp
write_dump all atom/adios singledump.bp
```

### 16.38.4 Restrictions

The number of atoms per snapshot **can** change with the adios style. When using the ADIOS tool ‘bpls’ to list the content of a .bp file, bpls will print `__` for the size of the output table indicating that its size is changing every step.

The *atom/adios* and *custom/adios* dump styles are part of the ADIOS package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 16.38.5 Related commands

*dump*, *dump\_modify*, *undump*

## 16.39 dump cfg/uef command

### 16.39.1 Syntax

```
dump ID group-ID cfg/uef N file mass type xs ys zs args
```

- ID = user-assigned name for the dump
  - group-ID = ID of the group of atoms to be dumped
  - N = dump every this many timesteps
  - file = name of file to write dump info to
- args = same as args for *dump custom*

### 16.39.2 Examples

```
dump 1 all cfg/uef 10 dump.*.cfg mass type xs ys zs
dump 2 all cfg/uef 100 dump.*.cfg mass type xs ys zs id c_stress
```

### 16.39.3 Description

This command is used to dump atomic coordinates in the reference frame of the applied flow field when *fix nvt/uef* or *fix npt/uef* is used. Only the atomic coordinates and frame-invariant scalar quantities will be in the flow frame. If velocities are selected as output, for example, they will not be in the same reference frame as the atomic positions.

### 16.39.4 Restrictions

This fix is part of the UEF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This command can only be used when *fix nvt/uef* or *fix npt/uef* is active.

### 16.39.5 Related commands

*dump, fix nvt/uef*

### 16.39.6 Default

none

## 16.40 dump h5md command

### 16.40.1 Syntax

```
dump ID group-ID h5md N file.h5 args
```

- ID = user-assigned name for the dump
- group-ID = ID of the group of atoms to be imaged
- *h5md* = style of dump command (other styles *atom* or *cfg* or *dcd* or *xtc* or *xyz* or *local* or *custom* are discussed on the *dump* doc page)
- N = dump every this many timesteps
- file.h5 = name of file to write to
- args = *position* options or *image* or *velocity* options or *force* options or *species* options or *file\_from* ID or *box* value or *create\_group* value or *author* value = list of data elements to dump, with their dump “sub-intervals”

*position* options

*image*

*velocity* options

*force* options

*species* options

*file\_from* ID = do not open a new file, re-use the already opened file from dump ID

*box* value = *yes* or *no*

*create\_group* value = *yes* or *no*

*author* value = quoted string



Note that at least one element must be specified and that *image* may only be present if *position* is specified first.

For the elements *position*, *velocity*, *force* and *species*, a sub-interval may be specified to write the data only every  $N_{\text{element}}$  iterations of the dump (i.e. every  $N * N_{\text{element}}$  time steps). This is specified by this option directly following the element declaration:

```
options = every N_element
```

## 16.40.2 Examples

```
dump h5md1 all h5md 100 dump_h5md.h5 position image
dump h5md1 all h5md 100 dump_h5md.h5 position velocity every 10
dump h5md1 all h5md 100 dump_h5md.h5 velocity author "John Doe"
```

## 16.40.3 Description

Dump a snapshot of atom coordinates every  $N$  timesteps in the HDF5 based H5MD file format (*de Buyl*). HDF5 files are binary, portable and self-describing. This dump style will write only one file, on the root node.

Several dumps may write to the same file, by using *file\_from* and referring to a previously defined dump. Several groups may also be stored within the same file by defining several dumps. A dump that refers (via *file\_from*) to an already open dump ID and that concerns another particle group must specify *create\_group yes*.

Each data element is written every  $N * N_{\text{element}}$  steps. For *image*, no sub-interval is needed as it must be present at the same interval as *position*. *image* must be given after *position* in any case. The box information (edges in each dimension) is stored at the same interval than the *position* element, if present. Else it is stored every  $N$  steps.

---

**Note:** Because periodic boundary conditions are enforced only on timesteps when neighbor lists are rebuilt, the coordinates of an atom written to a dump file may be slightly outside the simulation box.

---

### Use from `write_dump`:

It is possible to use this dump style with the *write\_dump* command. In this case, the sub-intervals must not be set at all. The *write\_dump* command can be used either to create a new file or to add current data to an existing dump file by using the *file\_from* keyword.

Typically, the *species* data is fixed. The following two commands store the position data every 100 timesteps, with the image data, and store once the species data in the same file.

```
dump h5md1 all h5md 100 dump.h5 position image
write_dump all h5md dump.h5 file_from h5md1 species
```

## 16.40.4 Restrictions

The number of atoms per snapshot cannot change with the `h5md` style. The position data is stored wrapped (box boundaries not enforced, see note above). Only orthogonal domains are currently supported. This is a limitation of the present `dump h5md` command and not of H5MD itself.

The `h5md` dump style is part of the H5MD package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. It also requires (i) building the `ch5md` library provided with LAMMPS (See the [Build package](#) page for more info.) and (ii) having the [HDF5](#) library installed (C bindings are sufficient) on your system. The library `ch5md` is compiled with the `h5cc` wrapper provided by the HDF5 library.

---

## 16.40.5 Related commands

*dump*, *dump\_modify*, *undump*

---

(**de Buyl**) de Buyl, Colberg and Hofling, H5MD: A structured, efficient, and portable file format for molecular data, *Comp. Phys. Comm.* 185(6), 1546-1553 (2014) - [[arXiv:1308.6382](#)].

## 16.41 dump image command

## 16.42 dump movie command

(see below for *dump\_modify options* specific to dump image/movie)

### 16.42.1 Syntax

```
dump ID group-ID style N file color diameter keyword value ...
```

- ID = user-assigned name for the dump
- group-ID = ID of the group of atoms to be imaged
- style = *image* or *movie* = style of dump command (other styles such as *atom* or *cfg* or *dcd* or *xtc* or *xyz* or *local* or *custom* are discussed on the *dump* doc page)
- N = dump every this many timesteps
- file = name of file to write image to
- color = atom attribute that determines color of each atom
- diameter = atom attribute that determines size of each atom
- zero or more keyword/value pairs may be appended
- keyword = *atom* or *adiam* or *bond* or *line* or *tri* or *body* or *fix* or *size* or *view* or *center* or *up* or *zoom* or *box* or *axes* or *subbox* or *shiny* or *ssao*

*atom* = *yes* or *no* = do or do not draw atoms  
*adiam* size = numeric value for atom diameter (distance units)  
*bond* values = color width = color and width of bonds  
    color = *atom* or *type* or *none*  
    width = number or *atom* or *type* or *none*  
        number = numeric value for bond width (distance units)  
*line* = color width  
    color = *type*  
    width = numeric value for line width (distance units)  
*tri* = color tflag width  
    color = *type*  
    tflag = 1 for just triangle, 2 for just tri edges, 3 for both  
    width = numeric value for triangle edge width (distance units)  
*body* = color bflag1 bflag2  
    color = *type*  
    bflag1,bflag2 = 2 numeric flags to affect how bodies are drawn  
*fix* = fixID color fflag1 fflag2  
    fixID = ID of fix that generates objects to draw  
    color = *type*  
    fflag1,fflag2 = 2 numeric flags to affect how fix objects are drawn  
*size* values = width height = size of images  
    width = width of image in # of pixels  
    height = height of image in # of pixels  
*view* values = theta phi = view of simulation box  
    theta = view angle from +z axis (degrees)  
    phi = azimuthal view angle (degrees)  
    theta or phi can be a variable (see below)  
*center* values = flag Cx Cy Cz = center point of image  
    flag = *s* for static, *d* for dynamic  
    Cx,Cy,Cz = center point of image as fraction of box dimension (0.5 = center of  
    ↳box)  
    Cx,Cy,Cz can be variables (see below)  
*up* values = Ux Uy Uz = direction that is "up" in image  
    Ux,Uy,Uz = components of up vector  
    Ux,Uy,Uz can be variables (see below)  
*zoom* value = zfactor = size that simulation box appears in image  
    zfactor = scale image size by factor > 1 to enlarge, factor < 1 to shrink  
    zfactor can be a variable (see below)  
*box* values = *yes/no* diam = draw outline of simulation box  
    *yes/no* = do or do not draw simulation box lines  
    diam = diameter of box lines as fraction of shortest box length  
*axes* values = axes length diam = draw xyz axes  
    axes = *yes* or *no* = do or do not draw xyz axes lines next to simulation box  
    length = length of axes lines as fraction of respective box lengths  
    diam = diameter of axes lines as fraction of shortest box length  
*\*subbox* values = lines diam = draw outline of processor sub-domains  
    lines = *yes* or *no* = do or do not draw sub-domain lines  
    diam = diameter of sub-domain lines as fraction of shortest box length  
*shiny* value = sfactor = shininess of spheres and cylinders  
    sfactor = shininess of spheres and cylinders from 0.0 to 1.0  
*ssao* value = shading seed dfactor = SSAO depth shading  
    shading = *yes* or *no* = turn depth shading on/off  
    seed = random # seed (positive integer)  
    dfactor = strength of shading from 0.0 to 1.0

## 16.43 dump\_modify options for dump image/movie

### 16.43.1 Syntax

`dump_modify dump-ID keyword values ...`

- these keywords apply only to the *image* and *movie* styles and are documented on this page
- keyword = *acolor* or *adiam* or *amap* or *backcolor* or *bcolor* or *bdiam* or *boxcolor* or *color* or *bitrate* or *framerate*
- see the *dump modify* doc page for more general keywords

*acolor* args = type color  
type = atom type or range of types (see below)  
color = name of color or color1/color2/...

*adiam* args = type diam  
type = atom type or range of types (see below)  
diam = diameter of atoms of that type (distance units)

*amap* args = lo hi style delta N entry1 entry2 ... entryN  
lo = number or *min* = lower bound of range of color map  
hi = number or *max* = upper bound of range of color map  
style = 2 letters = *c* or *d* or *s* plus *a* or *f*  
  *c* for continuous  
  *d* for discrete  
  *s* for sequential  
  *a* for absolute  
  *f* for fractional  
delta = binsize (only used for style *s*, otherwise ignored)  
binsize = range is divided into bins of this width  
N = # of subsequent entries  
entry = value color (for continuous style)  
  value = number or *min* or *max* = single value within range  
  color = name of color used for that value  
entry = lo hi color (for discrete style)  
  lo/hi = number or *min* or *max* = lower/upper bound of subset of range  
  color = name of color used for that subset of values  
entry = color (for sequential style)  
  color = name of color used for a bin of values

*backcolor* arg = color  
color = name of color for background

*bcolor* args = type color  
type = bond type or range of types (see below)  
color = name of color or color1/color2/...

*bdiam* args = type diam  
type = bond type or range of types (see below)  
diam = diameter of bonds of that type (distance units)

*boxcolor* arg = color  
color = name of color for simulation box lines and processor sub-domain lines

*color* args = name R G B  
name = name of color  
R,G,B = red/green/blue numeric values from 0.0 to 1.0

*bitrate* arg = rate  
rate = target bitrate for movie in kbps

*framerate* arg = fps

fps = frames per second for movie

### 16.43.2 Examples

```
dump d0 all image 100 dump.*.jpg type type
dump d1 mobile image 500 snap.*.png element element ssao yes 4539 0.6
dump d2 all image 200 img-*.ppm type type zoom 2.5 adiam 1.5 size 1280 720
dump m0 all movie 1000 movie.mpg type type size 640 480
dump m1 all movie 1000 movie.avi type type size 640 480
dump m2 all movie 100 movie.m4v type type zoom 1.8 adiam v_value size 1280 720

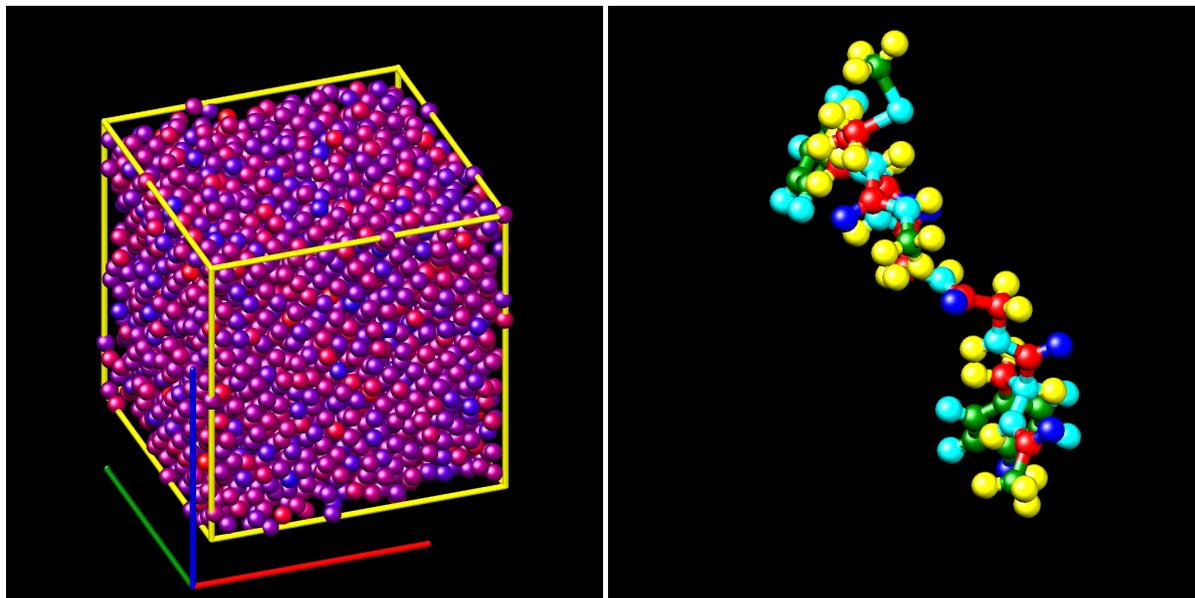
dump_modify 1 amap min max cf 0.0 3 min green 0.5 yellow max blue boxcolor red
```

### 16.43.3 Description

Dump a high-quality rendered image of the atom configuration every  $N$  timesteps and save the images either as a sequence of JPEG or PNG or PPM files, or as a single movie file. The options for this command as well as the *dump\_modify* command control what is included in the image or movie and how it appears. A series of such images can easily be manually converted into an animated movie of your simulation or the process can be automated without writing the intermediate files using the dump movie style; see further details below. Other dump styles store snapshots of numerical data associated with atoms in various formats, as discussed on the *dump* doc page.

Note that a set of images or a movie can be made after a simulation has been run, using the *rerun* command to read snapshots from an existing dump file, and using these dump commands in the rerun script to generate the images/movie.

Here are two sample images, rendered as  $1024 \times 1024$  JPEG files.



Only atoms in the specified group are rendered in the image. The *dump\_modify region and thresh* commands can also alter what atoms are included in the image. The filename suffix determines whether a JPEG, PNG, or PPM file is created with the *image* dump style. If the suffix is “.jpg” or “.jpeg”, then a JPEG format file is created, if the suffix is “.png”, then a PNG format is created, else a PPM (aka NETPBM) format file is created. The JPEG and PNG files are binary; PPM has a text mode header followed by binary data. JPEG images have lossy compression, PNG has lossless

compression, and PPM files are uncompressed but can be compressed with gzip, if LAMMPS has been compiled with `-DLAMMPS_GZIP` and a “.gz” suffix is used.

Similarly, the format of the resulting movie is chosen with the *movie* dump style. This is handled by the underlying FFmpeg converter and thus details have to be looked up in the [FFmpeg documentation](#). Typical examples are: .avi, .mpg, .m4v, .mp4, .mkv, .flv, .mov, .gif. Additional settings of the movie compression like bitrate and framerate can be set using the `dump_modify` command as described below.

To write out JPEG and PNG format files, you must build LAMMPS with support for the corresponding JPEG or PNG library. To convert images into movies, LAMMPS has to be compiled with the `-DLAMMPS_FFMPEG` flag. See the [Build settings](#) page for details.

---

**Note:** Because periodic boundary conditions are enforced only on timesteps when neighbor lists are rebuilt, the coordinates of an atom in the image may be slightly outside the simulation box.

---

Dumps are performed on timesteps that are a multiple of  $N$  (including timestep 0) and on the last timestep of a minimization if the minimization converges. Note that this means a dump will not be performed on the initial timestep after the dump command is invoked, if the current timestep is not a multiple of  $N$ . This behavior can be changed via the `dump_modify first` command, which can be useful if the dump command is invoked after a minimization ended on an arbitrary timestep.  $N$  can be changed between runs by using the `dump_modify every` command.

Dump *image* filenames must contain a wildcard character “\*” so that one image file per snapshot is written. The “\*” character is replaced with the timestep value. For example, `tmp.dump.*.jpg` becomes `tmp.dump.0.jpg`, `tmp.dump.10000.jpg`, `tmp.dump.20000.jpg`, etc. Note that the `dump_modify pad` command can be used to insure all timestep numbers are the same length (e.g., 00010), which can make it easier to convert a series of images into a movie in the correct ordering.

Dump *movie* filenames on the other hand, must not have any wildcard character since only one file combining all images into a single movie will be written by the movie encoder.

---

The *color* and *diameter* settings determine the color and size of atoms rendered in the image. They can be any atom attribute defined for the `dump custom` command, including *type* and *element*. This includes per-atom quantities calculated by a *compute*, *fix*, or *variable*, which are prefixed by “c\_”, “f\_”, or “v\_”, respectively. Note that the *diameter* setting can be overridden with a numeric value applied to all atoms by the optional *adiam* keyword.

If *type* is specified for the *color* setting, then the color of each atom is determined by its atom type. By default the mapping of types to colors is as follows:

- type 1 = red
- type 2 = green
- type 3 = blue
- type 4 = yellow
- type 5 = aqua
- type 6 = cyan

and repeats itself for types  $> 6$ . This mapping can be changed by the “`dump_modify acolor`” command, as described below.

If *type* is specified for the *diameter* setting then the diameter of each atom is determined by its atom type. By default all types have diameter 1.0. This mapping can be changed by the “`dump_modify adiam`” command, as described below.

If *element* is specified for the *color* and/or *diameter* setting, then the color and/or diameter of each atom is determined by which element it is, which in turn is specified by the element-to-type mapping specified by the “`dump_modify`

element” command, as described below. By default every atom type is C (carbon). Every element has a color and diameter associated with it, which is the same as the colors and sizes used by the [AtomEye](#) visualization package.

If other atom attributes are used for the *color* or *diameter* settings, they are interpreted in the following way.

If “vx”, for example, is used as the *color* setting, then the color of the atom will depend on the x-component of its velocity. The association of a per-atom value with a specific color is determined by a “color map”, which can be specified via the `dump_modify` command, as described below. The basic idea is that the atom-attribute will be within a range of values, and every value within the range is mapped to a specific color. Depending on how the color map is defined, that mapping can take place via interpolation so that a value of -3.2 is halfway between “red” and “blue”, or discretely so that the value of -3.2 is “orange”.

If “vx”, for example, is used as the *diameter* setting, then the atom will be rendered using the x-component of its velocity as the diameter. If the per-atom value  $\leq 0.0$ , then the atom will not be drawn. Note that finite-size spherical particles, as defined by *atom\_style sphere* define a per-particle radius or diameter, which can be used as the *diameter* setting.

---

The various keywords listed above control how the image is rendered. As listed below, all of the keywords have defaults, most of which you will likely not need to change. As described below, the `dump modify` command also has options specific to the dump image style, particularly for assigning colors to atoms, bonds, and other image features.

---

The *atom* keyword allow you to turn off the drawing of all atoms, if the specified value is *no*. Note that this will not turn off the drawing of particles that are represented as lines, triangles, or bodies, as discussed below. These particles can be drawn separately if the *line*, *tri*, or *body* keywords are used.

The *adiam* keyword allows you to override the *diameter* setting to set a single numeric *size*. All atoms will be drawn with that diameter, e.g. 1.5, which is in whatever distance *units* the input script defines, e.g. Angstroms.

---

The *bond* keyword allows to you to alter how bonds are drawn. A bond is only drawn if both atoms in the bond are being drawn due to being in the specified group and due to other selection criteria (e.g. region, threshold settings of the `dump_modify` command). By default, bonds are drawn if they are defined in the input data file as read by the `read_data` command. Using *none* for both the bond *color* and *width* value will turn off the drawing of all bonds.

If *atom* is specified for the bond *color* value, then each bond is drawn in 2 halves, with the color of each half being the color of the atom at that end of the bond.

If *type* is specified for the *color* value, then the color of each bond is determined by its bond type. By default the mapping of bond types to colors is as follows:

- type 1 = red
- type 2 = green
- type 3 = blue
- type 4 = yellow
- type 5 = aqua
- type 6 = cyan

and repeats itself for bond types  $> 6$ . This mapping can be changed by the “`dump_modify bcolor`” command, as described below.

The bond *width* value can be a numeric value or *atom* or *type* (or *none* as indicated above).

If a numeric value is specified, then all bonds will be drawn as cylinders with that diameter, e.g. 1.0, which is in whatever distance *units* the input script defines, e.g. Angstroms.

If *atom* is specified for the *width* value, then each bond will be drawn with a width corresponding to the minimum diameter of the 2 atoms in the bond.

If *type* is specified for the *width* value then the diameter of each bond is determined by its bond type. By default all types have diameter 0.5. This mapping can be changed by the “dump\_modify bdiam” command, as described below.

---

The *line* keyword can be used when *atom\_style line* is used to define particles as line segments, and will draw them as lines. If this keyword is not used, such particles will be drawn as spheres, the same as if they were regular atoms. The only setting currently allowed for the *color* value is *type*, which will color the lines according to the atom type of the particle. By default the mapping of types to colors is as follows:

- type 1 = red
- type 2 = green
- type 3 = blue
- type 4 = yellow
- type 5 = aqua
- type 6 = cyan

and repeats itself for types > 6. There is not yet an option to change this via the dump\_modify command.

The line *width* can only be a numeric value, which specifies that all lines will be drawn as cylinders with that diameter, e.g. 1.0, which is in whatever distance *units* the input script defines, e.g. Angstroms.

---

The *tri* keyword can be used when *atom\_style tri* is used to define particles as triangles, and will draw them as triangles or edges (3 lines) or both, depending on the setting for *tflag*. If edges are drawn, the *width* setting determines the diameters of the line segments. If this keyword is not used, triangle particles will be drawn as spheres, the same as if they were regular atoms. The only setting currently allowed for the *color* value is *type*, which will color the triangles according to the atom type of the particle. By default the mapping of types to colors is as follows:

- type 1 = red
- type 2 = green
- type 3 = blue
- type 4 = yellow
- type 5 = aqua
- type 6 = cyan

and repeats itself for types > 6. There is not yet an option to change this via the dump\_modify command.

---

The *body* keyword can be used when *atom\_style body* is used to define body particles with internal state (e.g. sub-particles), and will drawn them in a manner specific to the body style. If this keyword is not used, such particles will be drawn as spheres, the same as if they were regular atoms.

The [Howto body](#) page describes the body styles LAMMPS currently supports, and provides more details as to the kind of body particles they represent and how they are drawn by this dump image command. For all the body styles, individual atoms can be either a body particle or a usual point (non-body) particle. Non-body particles will be drawn the same way they would be as a regular atom. The *bflag1* and *bflag2* settings are numerical values which are passed to the body style to affect how the drawing of a body particle is done. See the [Howto body](#) page for a description of what these parameters mean for each body style.

The only setting currently allowed for the *color* value is *type*, which will color the body particles according to the atom type of the particle. By default the mapping of types to colors is as follows:



- type 1 = red
- type 2 = green
- type 3 = blue
- type 4 = yellow
- type 5 = aqua
- type 6 = cyan

and repeats itself for types > 6. There is not yet an option to change this via the `dump_modify` command.

---

The `fix` keyword can be used with a `fix` that produces objects to be drawn.

The `fflag1` and `fflag2` settings are numerical values which are passed to the `fix` to affect how the drawing of its objects is done. See the individual `fix` page for a description of what these parameters mean for a particular `fix`.

The only setting currently allowed for the `color` value is `type`, which will color the `fix` objects according to their type. By default the mapping of types to colors is as follows:

- type 1 = red
- type 2 = green
- type 3 = blue
- type 4 = yellow
- type 5 = aqua
- type 6 = cyan

and repeats itself for types > 6. There is not yet an option to change this via the `dump_modify` command.

---

The `size` keyword sets the width and height of the created images, i.e. the number of pixels in each direction.

---

The `view`, `center`, `up`, and `zoom` values determine how 3d simulation space is mapped to the 2d plane of the image. Basically they control how the simulation box appears in the image.

All of the `view`, `center`, `up`, and `zoom` values can be specified as numeric quantities, whose meaning is explained below. Any of them can also be specified as an *equal-style variable*, by using `v_name` as the value, where “name” is the variable name. In this case the variable will be evaluated on the timestep each image is created to create a new value. If the equal-style variable is time-dependent, this is a means of changing the way the simulation box appears from image to image, effectively doing a pan or fly-by view of your simulation.

The `view` keyword determines the viewpoint from which the simulation box is viewed, looking towards the `center` point. The `theta` value is the vertical angle from the +z axis, and must be an angle from 0 to 180 degrees. The `phi` value is an azimuthal angle around the z axis and can be positive or negative. A value of 0.0 is a view along the +x axis, towards the `center` point. If `theta` or `phi` are specified via variables, then the variable values should be in degrees.

The `center` keyword determines the point in simulation space that will be at the center of the image. `Cx`, `Cy`, and `Cz` are specified as fractions of the box dimensions, so that (0.5,0.5,0.5) is the center of the simulation box. These values do not have to be between 0.0 and 1.0, if you want the simulation box to be offset from the center of the image. Note, however, that if you choose strange values for `Cx`, `Cy`, or `Cz` you may get a blank image. Internally, `Cx`, `Cy`, and `Cz` are converted into a point in simulation space. If `flag` is set to “s” for static, then this conversion is done once, at the time the `dump` command is issued. If `flag` is set to “d” for dynamic then the conversion is performed every time a new image is created. If the box size or shape is changing, this will adjust the center point in simulation space.

The *up* keyword determines what direction in simulation space will be “up” in the image. Internally it is stored as a vector that is in the plane perpendicular to the view vector implied by the *theta* and *pni* values, and which is also in the plane defined by the view vector and user-specified up vector. Thus this internal vector is computed from the user-specified *up* vector as

```
up_internal = view cross (up cross view)
```

This means the only restriction on the specified *up* vector is that it cannot be parallel to the *view* vector, implied by the *theta* and *phi* values.

The *zoom* keyword scales the size of the simulation box as it appears in the image. The default *zfactor* value of 1 should display an image mostly filled by the atoms in the simulation box. A *zfactor* > 1 will make the simulation box larger; a *zfactor* < 1 will make it smaller. *Zfactor* must be a value > 0.0.

---

The *box* keyword determines if and how the simulation box boundaries are rendered as thin cylinders in the image. If *no* is set, then the box boundaries are not drawn and the *diam* setting is ignored. If *yes* is set, the 12 edges of the box are drawn, with a diameter that is a fraction of the shortest box length in x,y,z (for 3d) or x,y (for 2d). The color of the box boundaries can be set with the “dump\_modify boxcolor” command.

The *axes* keyword determines if and how the coordinate axes are rendered as thin cylinders in the image. If *no* is set, then the axes are not drawn and the *length* and *diam* settings are ignored. If *yes* is set, 3 thin cylinders are drawn to represent the x,y,z axes in colors red,green,blue. The origin of these cylinders will be offset from the lower left corner of the box by 10%. The *length* setting determines how long the cylinders will be as a fraction of the respective box lengths. The *diam* setting determines their thickness as a fraction of the shortest box length in x,y,z (for 3d) or x,y (for 2d).

The *subbox* keyword determines if and how processor sub-domain boundaries are rendered as thin cylinders in the image. If *no* is set (default), then the sub-domain boundaries are not drawn and the *diam* setting is ignored. If *yes* is set, the 12 edges of each processor sub-domain are drawn, with a diameter that is a fraction of the shortest box length in x,y,z (for 3d) or x,y (for 2d). The color of the sub-domain boundaries can be set with the “dump\_modify boxcolor” command.

---

The *shiny* keyword determines how shiny the objects rendered in the image will appear. The *sfactor* value must be a value  $0.0 \leq sfactor \leq 1.0$ , where *sfactor* = 1 is a highly reflective surface and *sfactor* = 0 is a rough non-shiny surface.

The *ssao* keyword turns on/off a screen space ambient occlusion (SSAO) model for depth shading. If *yes* is set, then atoms further away from the viewer are darkened via a randomized process, which is perceived as depth. The calculation of this effect can increase the cost of computing the image by roughly 2x. The strength of the effect can be scaled by the *dfactor* parameter. If *no* is set, no depth shading is performed.

---

A series of JPEG, PNG, or PPM images can be converted into a movie file and then played as a movie using commonly available tools. Using dump style *movie* automates this step and avoids the intermediate step of writing (many) image snapshot file. But LAMMPS has to be compiled with -DLAMMPS\_FFmpeg and an FFmpeg executable have to be installed.

To manually convert JPEG, PNG or PPM files into an animated GIF or MPEG or other movie file you can use:

- a) Use the ImageMagick convert program.

```
% convert *.jpg foo.gif  
% convert -loop 1 *.ppm foo.mpg
```

Animated GIF files from ImageMagick are not optimized. You can use a program like gifsicle to optimize and thus massively shrink them. MPEG files created by ImageMagick are in MPEG-1 format with a rather inefficient compression and low quality compared to more modern compression styles like MPEG-4, H.264, VP8, VP9, H.265 and so on.

- b) Use QuickTime.

Select “Open Image Sequence” under the File menu Load the images into QuickTime to animate them Select “Export” under the File menu Save the movie as a QuickTime movie (\*.mov) or in another format. QuickTime can generate very high quality and efficiently compressed movie files. Some of the supported formats require to buy a license and some are not readable on all platforms until specific runtime libraries are installed.

- c) Use FFmpeg

FFmpeg is a command line tool that is available on many platforms and allows extremely flexible encoding and decoding of movies.

```
cat snap.*.jpg | ffmpeg -y -f image2pipe -c:v mjpeg -i - -b:v 2000k movie.m4v
cat snap.*.ppm | ffmpeg -y -f image2pipe -c:v ppm -i - -b:v 2400k movie.avi
```

Front ends for FFmpeg exist for multiple platforms. For more information see the [FFmpeg homepage](#)

Play the movie:

- a) Use your browser to view an animated GIF movie.

Select “Open File” under the File menu Load the animated GIF file

- b) Use the freely available mplayer or ffplay tool to view a movie. Both are available for multiple OSES and support a large variety of file formats and decoders.

```
% mplayer foo.mpg
% ffplay bar.avi
```

- c) Use the [Pizza.py animate tool](#), which works directly on a series of image files.

```
a = animate("foo*.jpg")
```

- d) QuickTime and other Windows- or MacOS-based media players can obviously play movie files directly. Similarly for corresponding tools bundled with Linux desktop environments. However, due to licensing issues with some file formats, the formats may require installing additional libraries, purchasing a license, or may not be supported.

#### 16.43.4 Dump\_modify keywords for dump image and dump movie

The following dump\_modify keywords apply only to the dump image and dump movie styles. Any keyword that works with dump image also works with dump movie, since the movie is simply a collection of images. Some of the keywords only affect the dump movie style. The descriptions give details.

The *acolor* keyword can be used with the dump image command, when its atom color setting is *type*, to set the color that atoms of each type will be drawn in the image.

The specified *type* should be an integer from 1 to Ntypes = the number of atom types. A wildcard asterisk can be used in place of or in conjunction with the *type* argument to specify a range of atom types. This takes the form “\*” or “\*n” or

“n\*” or “m\*n”. If  $N$  = the number of atom types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $n$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive).

The specified *color* can be a single color which is any of the 140 pre-defined colors (see below) or a color name defined by the “dump\_modify color” command, as described below. Or it can be two or more colors separated by a “/” character, e.g. red/green/blue. In the former case, that color is assigned to all the specified atom types. In the latter case, the list of colors are assigned in a round-robin fashion to each of the specified atom types.

---

The *adiam* keyword can be used with the dump image command, when its atom diameter setting is *type*, to set the size that atoms of each type will be drawn in the image. The specified *type* should be an integer from 1 to  $N_{types}$ . As with the *acolor* keyword, a wildcard asterisk can be used as part of the *type* argument to specify a range of atom types. The specified *diam* is the size in whatever distance *units* the input script is using, e.g. Angstroms.

---

The *amap* keyword can be used with the dump image command, with its *atom* keyword, when its atom setting is an atom-attribute, to setup a color map. The color map is used to assign a specific RGB (red/green/blue) color value to an individual atom when it is drawn, based on the atom’s attribute, which is a numeric value, e.g. its x-component of velocity if the atom-attribute “vx” was specified.

The basic idea of a color map is that the atom-attribute will be within a range of values, and that range is associated with a series of colors (e.g. red, blue, green). An atom’s specific value (vx = -3.2) can then mapped to the series of colors (e.g. halfway between red and blue), and a specific color is determined via an interpolation procedure.

There are many possible options for the color map, enabled by the *amap* keyword. Here are the details.

The *lo* and *hi* settings determine the range of values allowed for the atom attribute. If numeric values are used for *lo* and/or *hi*, then values that are lower/higher than that value are set to the value. I.e. the range is static. If *lo* is specified as *min* or *hi* as *max* then the range is dynamic, and the lower and/or upper bound will be calculated each time an image is drawn, based on the set of atoms being visualized.

The *style* setting is two letters, such as “ca”. The first letter is either “c” for continuous, “d” for discrete, or “s” for sequential. The second letter is either “a” for absolute, or “f” for fractional.

A continuous color map is one in which the color changes continuously from value to value within the range. A discrete color map is one in which discrete colors are assigned to sub-ranges of values within the range. A sequential color map is one in which discrete colors are assigned to a sequence of sub-ranges of values covering the entire range.

An absolute color map is one in which the values to which colors are assigned are specified explicitly as values within the range. A fractional color map is one in which the values to which colors are assigned are specified as a fractional portion of the range. For example if the range is from -10.0 to 10.0, and the color red is to be assigned to atoms with a value of 5.0, then for an absolute color map the number 5.0 would be used. But for a fractional map, the number 0.75 would be used since 5.0 is 3/4 of the way from -10.0 to 10.0.

The *delta* setting must be specified for all styles, but is only used for the sequential style; otherwise the value is ignored. It specifies the bin size to use within the range for assigning consecutive colors to. For example, if the range is from -10.0 to 10.0 and a *delta* of 1.0 is used, then 20 colors will be assigned to the range. The first will be from  $-10.0 \leq color1 < -9.0$ , then second from  $-9.0 \leq color2 < -8.0$ , etc.

The *N* setting is how many entries follow. The format of the entries depends on whether the color map style is continuous, discrete or sequential. In all cases the *color* setting can be any of the 140 pre-defined colors (see below) or a color name defined by the dump\_modify color option.

For continuous color maps, each entry has a *value* and a *color*. The *value* is either a number within the range of values or *min* or *max*. The *value* of the first entry must be *min* and the *value* of the last entry must be *max*. Any entries in between must have increasing values. Note that numeric values can be specified either as absolute numbers or as fractions (0.0 to 1.0) of the range, depending on the “a” or “f” in the style setting for the color map.

Here is how the entries are used to determine the color of an individual atom, given the value  $X$  of its atom attribute.  $X$  will fall between 2 of the entry values. The color of the atom is linearly interpolated (in each of the RGB values) between the 2 colors associated with those entries. For example, if  $X = -5.0$  and the two surrounding entries are “red” at  $-10.0$  and “blue” at  $0.0$ , then the atom’s color will be halfway between “red” and “blue”, which happens to be “purple”.

For discrete color maps, each entry has a *lo* and *hi* value and a *color*. The *lo* and *hi* settings are either numbers within the range of values or *lo* can be *min* or *hi* can be *max*. The *lo* and *hi* settings of the last entry must be *min* and *max*. Other entries can have any *lo* and *hi* values and the sub-ranges of different values can overlap. Note that numeric *lo* and *hi* values can be specified either as absolute numbers or as fractions (0.0 to 1.0) of the range, depending on the “a” or “f” in the style setting for the color map.

Here is how the entries are used to determine the color of an individual atom, given the value  $X$  of its atom attribute. The entries are scanned from first to last. The first time that  $lo \leq X \leq hi$ ,  $X$  is assigned the color associated with that entry. You can think of the last entry as assigning a default color (since it will always be matched by  $X$ ), and the earlier entries as colors that override the default. Also note that no interpolation of a color RGB is done. All atoms will be drawn with one of the colors in the list of entries.

For sequential color maps, each entry has only a *color*. Here is how the entries are used to determine the color of an individual atom, given the value  $X$  of its atom attribute. The range is partitioned into  $N$  bins of width *binsize*. Thus  $X$  will fall in a specific bin from 1 to  $N$ , say the  $M$ th bin. If it falls on a boundary between 2 bins, it is considered to be in the higher of the 2 bins. Each bin is assigned a color from the  $E$  entries. If  $E < N$ , then the colors are repeated. For example if 2 entries with colors red and green are specified, then the odd numbered bins will be red and the even bins green. The color of the atom is the color of its bin. Note that the sequential color map is really a shorthand way of defining a discrete color map without having to specify where all the bin boundaries are.

Here is an example of using a sequential color map to color all the atoms in individual molecules with a different color. See the examples/pour/in.pour.2d.molecule input script for an example of how this is used.

```
variable colors string &
    "red green blue yellow white &
    purple pink orange lime gray"
variable mol atom mol%10
dump 1 all image 250 image.*.jpg v_mol type &
    zoom 1.6 adiam 1.5
dump_modify 1 pad 5 amap 0 10 sa 1 10 ${colors}
```

In this case, 10 colors are defined, and molecule IDs are mapped to one of the colors, even if there are 1000s of molecules.

The *backcolor* sets the background color of the images. The color name can be any of the 140 pre-defined colors (see below) or a color name defined by the *dump\_modify* color option.

The *bcolor* keyword can be used with the *dump* image command, with its *bond* keyword, when its color setting is *type*, to set the color that bonds of each type will be drawn in the image.

The specified *type* should be an integer from 1 to  $N$ , where  $N$  is the number of bond types. A wildcard asterisk can be used in place of or in conjunction with the *type* argument to specify a range of bond types. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the number of bond types, then an asterisk with no numerical values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $m$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive).

The specified *color* can be a single color which is any of the 140 pre-defined colors (see below) or a color name defined by the *dump\_modify* color option. Or it can be two or more colors separated by a “/” character (e.g., red/green/blue).

In the former case, that color is assigned to all the specified bond types. In the latter case, the list of colors are assigned in a round-robin fashion to each of the specified bond types.

---

The *bdiam* keyword can be used with the *dump image* command, with its *bond* keyword, when its *diam* setting is *type*, to set the diameter that bonds of each type will be drawn in the image. The specified *type* should be an integer from 1 to *Nbondtypes*. As with the *bcolor* keyword, a wildcard asterisk can be used as part of the *type* argument to specify a range of bond types. The specified *diam* is the size in whatever distance *units* you are using (e.g., Angstroms).

---

The *bitrate* keyword can be used with the *dump movie* command to define the size of the resulting movie file and its quality via setting how many kbits per second are to be used for the movie file. Higher bitrates require less compression and will result in higher quality movies. The quality is also determined by the compression format and encoder. The default setting is 2000 kbit/s, which will result in average quality with older compression formats.

---

**Note:** Not all movie file formats supported by *dump movie* allow the *bitrate* to be set. If not, the setting is silently ignored.

---

The *boxcolor* keyword sets the color of the simulation box drawn around the atoms in each image as well as the color of processor sub-domain boundaries. See the “*dump image box*” command for how to specify that a box be drawn via the *box* keyword, and the sub-domain boundaries via the *subbox* keyword. The color name can be any of the 140 pre-defined colors (see below) or a color name defined by the *dump\_modify color* option.

---

The *color* keyword allows definition of a new color name, in addition to the 140-predefined colors (see below), and associates three red/green/blue RGB values with that color name. The color name can then be used with any other *dump\_modify* keyword that takes a color name as a value. The RGB values should each be floating point values between 0.0 and 1.0 inclusive.

When a color name is converted to RGB values, the user-defined color names are searched first, then the 140 pre-defined color names. This means you can also use the *color* keyword to overwrite one of the pre-defined color names with new RGB values.

---

The *framerate* keyword can be used with the *dump movie* command to define the duration of the resulting movie file. Movie files written by the *dump movie* command have a default frame rate of 24 frames per second and the images generated will be converted at that rate. Thus a sequence of 1000 *dump image* commands will result in a movie of about 42 seconds. To make a movie run longer you can either generate images more frequently or lower the frame rate. To speed a movie up, you can do the inverse. Using a frame rate higher than 24 is not recommended, as it will result in simply dropping the rendered images. It is more efficient to *dump image* less frequently.

---

### 16.43.5 Restrictions

To write JPEG images, you must use the `-DLAMMPS_JPEG` switch when building LAMMPS and link with a JPEG library. To write PNG images, you must use the `-DLAMMPS_PNG` switch when building LAMMPS and link with a PNG library.

To write *movie* dumps, you must use the `-DLAMMPS_FFmpeg` switch when building LAMMPS and have the FFmpeg executable available on the machine where LAMMPS is being run. Typically its name is lowercase (i.e., “`ffmpeg`”).

---

See the *Build settings* page for details.

Note that since FFmpeg is run as an external program via a pipe, LAMMPS has limited control over its execution and no knowledge about errors and warnings printed by it. Those warnings and error messages will be printed to the screen only. Due to the way image data are communicated to FFmpeg, it will often print the message

```
pipe:: Input/output error
```

which can be safely ignored. Other warnings and errors have to be addressed according to the FFmpeg documentation. One known issue is that certain movie file formats (e.g., MPEG level 1 and 2 format streams) have video bandwidth limits that can be crossed when rendering too large of image sizes. Typical warnings look like this:

```
[mpeg @ 0x98b5e0] packet too large, ignoring buffer limits to mux it
[mpeg @ 0x98b5e0] buffer underflow st=0 bufi=281407 size=285018
[mpeg @ 0x98b5e0] buffer underflow st=0 bufi=283448 size=285018
```

In this case it is recommended either to reduce the size of the image or to encode in a different format that is also supported by your copy of FFmpeg and which does not have this limitation (e.g., .avi, .mkv, mp4).

### 16.43.6 Related commands

*dump*, *dump\_modify*, *undump*

### 16.43.7 Default

The defaults for the dump image and dump movie keywords are as follows:

- adiam = not specified (use diameter setting)
- atom = yes
- bond = none none (if no bonds in system)
- bond = atom 0.5 (if bonds in system)
- size = 512 512
- view = 60 30 (for 3d)
- view = 0 0 (for 2d)
- center = s 0.5 0.5 0.5
- up = 0 0 1 (for 3d)
- up = 0 1 0 (for 2d)
- zoom = 1.0
- box = yes 0.02
- axes = no 0.0 0.0
- subbox no 0.0
- shiny = 1.0
- ssao = no

The defaults for the dump\_modify keywords specific to dump image and dump movie are as follows:

- `acolor = * red/green/blue/yellow/aqua/cyan`
  - `adiam = * 1.0`
  - `amap = min max cf 0.0 2 min blue max red`
  - `backcolor = black`
  - `bcolor = * red/green/blue/yellow/aqua/cyan`
  - `bdiam = * 0.5`
  - `bitrate = 2000`
  - `boxcolor = yellow`
  - `color = 140` color names are pre-defined as listed below
  - `framerate = 24`
- 

These are the standard 109 element names that LAMMPS pre-defines for use with the `dump image` and `dump_modify` commands.

- 1-10 = "H", "He", "Li", "Be", "B", "C", "N", "O", "F", "Ne"
  - 11-20 = "Na", "Mg", "Al", "Si", "P", "S", "Cl", "Ar", "K", "Ca"
  - 21-30 = "Sc", "Ti", "V", "Cr", "Mn", "Fe", "Co", "Ni", "Cu", "Zn"
  - 31-40 = "Ga", "Ge", "As", "Se", "Br", "Kr", "Rb", "Sr", "Y", "Zr"
  - 41-50 = "Nb", "Mo", "Tc", "Ru", "Rh", "Pd", "Ag", "Cd", "In", "Sn"
  - 51-60 = "Sb", "Te", "I", "Xe", "Cs", "Ba", "La", "Ce", "Pr", "Nd"
  - 61-70 = "Pm", "Sm", "Eu", "Gd", "Tb", "Dy", "Ho", "Er", "Tm", "Yb"
  - 71-80 = "Lu", "Hf", "Ta", "W", "Re", "Os", "Ir", "Pt", "Au", "Hg"
  - 81-90 = "Tl", "Pb", "Bi", "Po", "At", "Rn", "Fr", "Ra", "Ac", "Th"
  - 91-100 = "Pa", "U", "Np", "Pu", "Am", "Cm", "Bk", "Cf", "Es", "Fm"
  - 101-109 = "Md", "No", "Lr", "Rf", "Db", "Sg", "Bh", "Hs", "Mt"
- 

These are the 140 colors that LAMMPS pre-defines for use with the `dump image` and `dump_modify` commands. Additional colors can be defined with the `dump_modify color` command. The 3 numbers listed for each name are the RGB (red/green/blue) values. Divide each value by 255 to get the equivalent 0.0 to 1.0 value.



aliceblue = 240, 248, 255	antiquewhite = 250, 235, 215	aqua = 0, 255, 255	aquamarine = 127, 255, 212	azure = 240, 255, 255
beige = 245, 245, 220	bisque = 255, 228, 196	black = 0, 0, 0	blanchedalmond = 255, 255, 205	blue = 0, 0, 255
blueviolet = 138, 43, 226	brown = 165, 42, 42	burlywood = 222, 184, 135	cadetblue = 95, 158, 160	chartreuse = 127, 255, 0
chocolate = 210, 105, 30	coral = 255, 127, 80	cornflowerblue = 100, 149, 237	cornsilk = 255, 248, 220	crimson = 220, 20, 60
cyan = 0, 255, 255	darkblue = 0, 0, 139	darkcyan = 0, 139, 139	darkgoldenrod = 184, 134, 11	darkgray = 169, 169, 169
darkgreen = 0, 100, 0	darkkhaki = 189, 183, 107	darkmagenta = 139, 0, 139	darkolivegreen = 85, 107, 47	darkorange = 255, 140, 0
darkorchid = 153, 50, 204	darkred = 139, 0, 0	darksalmon = 233, 150, 122	darkseagreen = 143, 188, 143	darkslateblue = 72, 61, 139
darkslategray = 47, 79, 79	darkturquoise = 0, 206, 209	darkviolet = 148, 0, 211	deeppink = 255, 20, 147	deepskyblue = 0, 191, 255
dimgray = 105, 105, 105	dodgerblue = 30, 144, 255	firebrick = 178, 34, 34	floralwhite = 255, 250, 240	forestgreen = 34, 139, 34
fuchsia = 255, 0, 255	gainsboro = 220, 220, 220	ghostwhite = 248, 248, 255	gold = 255, 215, 0	goldenrod = 218, 165, 32
gray = 128, 128, 128	green = 0, 128, 0	greenyellow = 173, 255, 47	honeydew = 240, 255, 240	hotpink = 255, 105, 180
indianred = 205, 92, 92	indigo = 75, 0, 130	ivory = 255, 240, 240	khaki = 240, 230, 140	lavender = 230, 230, 250
lavenderblush = 255, 240, 245	lawngreen = 124, 252, 0	lemonchiffon = 255, 250, 205	lightblue = 173, 216, 230	lightcoral = 240, 128, 128
lightcyan = 224, 255, 255	lightgoldenrodyellow = 250, 250, 210	lightgreen = 144, 238, 144	lightgrey = 211, 211, 211	lightpink = 255, 182, 193
lightsalmon = 255, 160, 122	lightseagreen = 32, 178, 170	lightskyblue = 135, 206, 250	lightslategray = 119, 136, 153	lightsteelblue = 176, 196, 222
lightyellow = 255, 255, 224	lime = 0, 255, 0	limegreen = 50, 205, 50	linen = 250, 240, 230	magenta = 255, 0, 255
maroon = 128, 0, 0	mediumaquamarine = 102, 205, 170	mediumblue = 0, 0, 205	mediumorchid = 186, 85, 211	mediumpurple = 147, 112, 219
mediumseagreen = 60, 179, 113	mediumslateblue = 123, 104, 238	mediumspringgreen = 0, 250, 154	mediumturquoise = 72, 209, 204	mediumvioletred = 199, 21, 133
midnightblue = 25, 25, 112	mintcream = 245, 255, 250	mistyrose = 255, 228, 225	moccasin = 255, 228, 181	navajowhite = 255, 222, 173
navy = 0, 0, 128	oldlace = 253, 245, 230	olive = 128, 128, 0	olivedrab = 107, 142, 35	orange = 255, 165, 0
orangered = 255, 69, 0	orchid = 218, 112, 214	palegoldenrod = 238, 232, 170	palegreen = 152, 251, 152	paleturquoise = 175, 238, 238
palevioletred = 219, 112, 147	papayawhip = 255, 239, 213	peachpuff = 255, 239, 213	peru = 205, 133, 63	pink = 255, 192, 203
plum = 221, 160, 221	powderblue = 176, 224, 230	purple = 128, 0, 128	red = 255, 0, 0	rosybrown = 188, 143, 143
royalblue = 65, 105, 225	saddlebrown = 139, 69, 19	salmon = 250, 128, 114	sandybrown = 244, 164, 96	seagreen = 46, 139, 87
seashell = 255, 245, 238	sienna = 160, 82, 45	silver = 192, 192, 192	skyblue = 135, 206, 235	slateblue = 106, 90, 205
slategray = 112, 128, 144	snow = 255, 250, 250	springgreen = 0, 255, 127	steelblue = 70, 130, 180	tan = 210, 180, 140
teal = 0, 128, 128	thistle = 216, 191, 216	tomato = 253, 99, 71	turquoise = 64, 224, 208	violet = 238, 130, 238
white = 255, 255, 255	white = 255, 255, 255	whitesmoke = 245, 245, 245	yellow = 255, 255, 0	yellowgreen = 154, 205, 50

## 16.44 dump\_modify command

## 16.45 dump\_modify command for image/movie options

### 16.45.1 Syntax

```
dump_modify dump-ID keyword values ...
```

- dump-ID = ID of dump to modify
- one or more keyword/value pairs may be appended
- these keywords apply to various dump styles
- keyword = *append* or *at* or *balance* or *buffer* or *delay* or *element* or *every* or *every/time* or *fileper* or *first* or *flush* or *format* or *header* or *image* or *label* or *maxfiles* or *nfile* or *pad* or *pbcs* or *precision* or *region* or *refresh* or *scale* or *sfactor* or *skip* or *sort* or *tfactor* or *thermo* or *thresh* or *time* or *units* or *unwrap*

*append* arg = *yes* or *no*

*at* arg = N

N = index of frame written upon first dump

*balance* arg = *yes* or *no*

*buffer* arg = *yes* or *no*

*colname* values = ID string, or *default*

string = new column header name

ID = integer from 1 to N, or integer from -1 to -N, where N = # of quantities.

↪being output

or a custom dump keyword or reference to compute, fix, property or variable.

*delay* arg = Dstep

Dstep = delay output until this timestep

*element* args = E1 E2 ... EN, where N = # of atom types

E1,...,EN = element name (e.g., C or Fe or Ga)

*every* arg = N

N = dump on timesteps which are a multiple of N

N can be a variable (see below)

*every/time* arg = Delta

Delta = dump once every Delta interval of simulation time (time units)

Delta can be a variable (see below)

*fileper* arg = Np

Np = write one file for every this many processors

*first* arg = *yes* or *no*

*flush* arg = *yes* or *no*

*format* args = *line* string, *int* string, *float* string, ID string, or *none*

string = C-style format string

ID = integer from 1 to N, or integer from -1 to -N, where N = # of quantities.

↪being output

or a custom dump keyword or reference to compute, fix, property or variable.

*header* arg = *yes* or *no*

*yes* to write the header

*no* to not write the header

*image* arg = *yes* or *no*

*label* arg = string

string = character string (e.g., BONDS) to use in header of dump local file

*maxfiles* arg = Fmax

*Fmax* = keep only the most recent *Fmax* snapshots (one snapshot per file)  
*nfile* arg = *Nf*  
*Nf* = write this many files, one from each of *Nf* processors  
*pad* arg = *Nchar* = # of characters to convert timestep to  
*pbcs* arg = *yes* or *no* = remap atoms via periodic boundary conditions  
*precision* arg = power-of-10 value from 10 to 1000000  
*region* arg = region-ID or "none"  
*refresh* arg = *c\_ID* = compute ID that supports a refresh operation  
*scale* arg = *yes* or *no*  
*sfactor* arg = coordinate scaling factor (> 0.0)  
*skip* arg = *v\_name*  
*v\_name* = variable with name which evaluates to non-zero (skip) or 0  
*sort* arg = *off* or *id* or *N* or *-N*  
*off* = no sorting of per-atom lines within a snapshot  
*id* = sort per-atom lines by atom ID  
*N* = sort per-atom lines in ascending order by the *N*th column  
*-N* = sort per-atom lines in descending order by the *N*th column  
*tfactor* arg = time scaling factor (> 0.0)  
*thermo* arg = *yes* or *no*  
*time* arg = *yes* or *no*  
*thresh* args = attribute operator value  
attribute = same attributes (*x*,*fy*,*etotal*,*sxx*,etc) used by dump custom style  
operator = "<" or "<=" or ">" or ">=" or "==" or "!=" or "|^"  
value = numeric value to compare to, or LAST  
these 3 args can be replaced by the word "none" to turn off thresholding  
*units* arg = *yes* or *no*  
*unwrap* arg = *yes* or *no*

- these keywords apply only to the *image* and *movie* styles
- keyword = *acolor* or *adiam* or *amap* or *backcolor* or *bcolor* or *bdiam* or *boxcolor* or *color* or *bitrate* or *framerate*  
see the [dump image](#) doc page for details
- these keywords apply only to the */gz* and */zstd* dump styles
- keyword = *compression\_level*  
*compression\_level* args = level  
level = integer specifying the compression level that should be used (see below ↪  
↪ for supported levels)
- these keywords apply only to the */zstd* dump styles
- keyword = *checksum*  
*checksum* args = *yes* or *no* (add checksum at end of zst file)

## 16.45.2 Examples

```
dump_modify 1 format line "%d %d %20.15g %g %g" scale yes
dump_modify 1 format float %20.15g scale yes
dump_modify myDump image yes scale no flush yes
dump_modify 1 region mySphere thresh x < 0.0 thresh fx >= 3.2
dump_modify xtcdump precision 10000 sfactor 0.1
dump_modify 1 every 1000 nfile 20
dump_modify 1 every v_myVar
```

## 16.45.3 Description

Modify the parameters of a previously defined dump command. Not all parameters are relevant to all dump styles.

As explained on the *dump* doc page, the *atom/mpiio*, *custom/mpiio*, and *xyz/mpiio* dump styles are identical in command syntax and in the format of the dump files they create, to the corresponding styles without “mpiio”, except the single dump file they produce is written in parallel via the MPI-IO library. Thus if a *dump\_modify* option below is valid for the *atom* style, it is also valid for the *atom/mpiio* style, and similarly for the other styles which allow for use of MPI-IO.

---

Unless otherwise noted, the following keywords apply to all the various dump styles, including the *dump image* and *dump movie* styles.

---

The *append* keyword applies to all dump styles except *cfg* and *xtc* and *dcd*. It also applies only to text output files, not to binary or gzipped or image/movie files. If specified as *yes*, then dump snapshots are appended to the end of an existing dump file. If specified as *no*, then a new dump file will be created which will overwrite an existing file with the same name.

---

The *at* keyword only applies to the *netcdf* dump style. It can only be used if the *append yes* keyword is also used. The *N* argument is the index of which frame to append to. A negative value can be specified for *N*, which means a frame counted from the end of the file. The *at* keyword can only be used if the *dump\_modify* command is before the first command that causes dump snapshots to be output (e.g., a *run* or *minimize* command). Once the dump file has been opened, this keyword has no further effect.

---

The *buffer* keyword applies only to dump styles *atom*, *cfg*, *custom*, *local*, and *xyz*. It also applies only to text output files, not to binary or gzipped files. If specified as *yes*, which is the default, then each processor writes its output into an internal text buffer, which is then sent to the processor(s) which perform file writes, and written by those processor(s) as one large chunk of text. If specified as *no*, each processor sends its per-atom data in binary format to the processor(s) which perform file writes, and those processor(s) format and write it line by line into the output file.

The buffering mode is typically faster since each processor does the relatively expensive task of formatting the output for its own atoms. However it requires about twice the memory (per processor) for the extra buffering.

---

New in version 4May2022.

The *colname* keyword can be used to change the default header keyword for dump styles: *atom*, *custom*, and *cfg* and their compressed, ADIOS, and MPIIO variants. The setting for *ID string* replaces the default text with the provided string. *ID* can be a positive integer when it represents the column number counting from the left, a negative integer when it represents the column number from the right (i.e. -1 is the last column/keyword), or a custom dump keyword (or

compute, fix, property, or variable reference) and then it replaces the string for that specific keyword. For *atom* dump styles only the keywords “id”, “type”, “x”, “y”, “z”, “ix”, “iy”, “iz” can be accessed via string regardless of whether scaled or unwrapped coordinates were enabled or disabled, and it always assumes 8 columns for indexing regardless of whether image flags are enabled or not. For dump style *cfg* only changes to the “auxiliary” keywords (6th or later keyword) will become visible.

The *colname* keyword can be used multiple times. If multiple *colname* settings refer to the same keyword, the last setting has precedence. A setting of *default* clears all previous settings, reverting all values to their default names. Using the *scale* or *image* keyword will also reset all header keywords to their default values.

The *delay* keyword applies to all dump styles. No snapshots will be output until the specified *Dstep* timestep or later. Specifying *Dstep* < 0 is the same as turning off the delay setting. This is a way to turn off unwanted output early in a simulation, for example, during an equilibration phase.

The *element* keyword applies only to the dump *cfg*, *xyz*, and *image* styles. It associates element names (e.g., H, C, Fe) with LAMMPS atom types. See the list of element names at the bottom of this page.

In the case of dump *cfg*, this allows the [AtomEye](#) visualization package to read the dump file and render atoms with the appropriate size and color.

In the case of dump *image*, the output images will follow the same [AtomEye](#) convention. An element name is specified for each atom type (1 to Ntype) in the simulation. The same element name can be given to multiple atom types.

In the case of *xyz* format dumps, there are no restrictions to what label can be used as an element name. Any white-space separated text will be accepted.

The *every* keyword can be used with any dump style except the *dcd* and *xtc* styles. It specifies that the output of dump snapshots will now be performed on timesteps which are a multiple of a new *N* value. This overrides the dump frequency originally specified by the *dump* command.

The *every* keyword can be specified in one of two ways. It can be a numeric value in which case it must be > 0. Or it can be an *equal-style variable*, which should be specified as *v\_name*, where name is the variable name.

In this case, the variable is evaluated at the beginning of a run to determine the next timestep at which a dump snapshot will be written out. On that timestep the variable will be evaluated again to determine the next timestep, etc. Thus the variable should return timestep values. See the *stagger()* and *logfreq()* and *stride()* math functions for *equal-style variables*, as examples of useful functions to use in this context. Other similar math functions could easily be added as options for *equal-style variables*. Also see the *next()* function, which allows use of a file-style variable which reads successive values from a file, each time the variable is evaluated. Used with the *every* keyword, if the file contains a list of ascending timesteps, you can output snapshots whenever you wish.

Note that when using the variable option with the *every* keyword, you need to use the *first* option if you want an initial snapshot written to the dump file. The *every* keyword cannot be used with the dump *dcd* style.

For example, the following commands will write snapshots at timesteps 0,10,20,30,100,200,300,1000,2000,etc:

```
variable      s equal logfreq(10,3,10)
dump          1 all atom 100 tmp.dump
dump_modify  1 every v_s first yes
```

The following commands would write snapshots at the timesteps listed in file tmp.times:

```
variable      f file tmp.times
variable      s equal next(f)
dump          1 all atom 100 tmp.dump
dump_modify  1 every v_s
```

**Note:** When using a file-style variable with the *every* keyword, the file of timesteps must list a first timestep that is beyond the current timestep (e.g., it cannot be 0). And it must list one or more timesteps beyond the length of the run you perform. This is because the dump command will generate an error if the next timestep it reads from the file is not a value greater than the current timestep. Thus if you wanted output on steps 0,15,100 of a 100-timestep run, the file should contain the values 15,100,101 and you should also use the `dump_modify first` command. Any final value > 100 could be used in place of 101.

---

The *every/time* keyword can be used with any dump style except the *dcd* and *xtc* styles. It changes the frequency of dump snapshots from being based on the current timestep to being determined by elapsed simulation time, i.e. in time units of the *units* command, and specifies *Delta* for the interval between snapshots. This can be useful when the timestep size varies during a simulation run, e.g. by use of the *fix dt/reset* command. The default is to perform output on timesteps which are multiples of specified timestep value *N*; see the *every* keyword.

The *every/time* keyword can be used with any dump style except the *dcd* and *xtc* styles. It does two things. It specifies that the interval between dump snapshots will be set in simulation time (i.e. in time units of the *units* command). This can be useful when the timestep size varies during a simulation run (e.g., by use of the *fix dt/reset* command). The default is to specify the interval in timesteps; see the *every* keyword. The *every/time* command also sets the interval value.

**Note:** If you wish dump styles *atom*, *custom*, *local*, or *xyz* to include the simulation time as a field in the header portion of each snapshot, you also need to use the `dump_modify time` keyword with a setting of *yes*. See its documentation below.

---

Note that since snapshots are output on simulation steps, each snapshot will be written on the first timestep whose associated simulation time is  $\geq$  the exact snapshot time value.

As with the *every* option, the *Delta* value can be specified in one of two ways. It can be a numeric value in which case it must be > 0.0. Or it can be an *equal-style variable*, which should be specified as `v_name`, where name is the variable name.

In this case, the variable is evaluated at the beginning of a run to determine the next simulation time at which a dump snapshot will be written out. On that timestep the variable will be evaluated again to determine the next simulation time, etc. Thus the variable should return values in time units. Note the current timestep or simulation time can be used in an *equal-style variable* since they are both thermodynamic keywords. Also see the `next()` function, which allows use of a file-style variable which reads successive values from a file, each time the variable is evaluated. Used with the *every/time* keyword, if the file contains a list of ascending simulation times, you can output snapshots whenever you wish.

Note that when using the variable option with the *every/time* keyword, you need to use the *first* option if you want an initial snapshot written to the dump file. The *every/time* keyword cannot be used with the dump *dcd* style.

For example, the following commands will write snapshots at successive simulation times which grow by a factor of 1.5 with each interval. The `dt` value used in the variable is to avoid a zero result when the initial simulation time is 0.0.

```
variable      increase equal 1.5*(time+dt)
dump          1 all atom 100 tmp.dump
dump_modify   1 every/time v_increase first yes
```

The following commands would write snapshots at the times listed in file `tmp.times`:

```
variable      f file tmp.times
variable      s equal next(f)
```

(continues on next page)

(continued from previous page)

```

dump          1 all atom 100 tmp.dump
dump_modify  1 every/time v_s

```

**Note:** When using a file-style variable with the *every/time* keyword, the file of timesteps must list a first time that is beyond the time associated with the current timestep (e.g., it cannot be 0.0). And it must list one or more times beyond the length of the run you perform. This is because the dump command will generate an error if the next time it reads from the file is not a value greater than the current time. Thus if you wanted output at times 0,15,100 of a run of length 100 in simulation time, the file should contain the values 15,100,101 and you should also use the `dump_modify first` command. Any final value > 100 could be used in place of 101.

The *first* keyword determines whether a dump snapshot is written on the very first timestep after the dump command is invoked. This will always occur if the current timestep is a multiple of `$N$`, the frequency specified in the `dump` command or `dump_modify every` command, including timestep 0. It will also always occur if the current simulation time is a multiple of *Delta*, the time interval specified in the `dump_modify every/time` command.

But if this is not the case, a dump snapshot will only be written if the setting of this keyword is *yes*. If it is *no*, which is the default, then it will not be written.

Note that if the argument to the `dump_modify every dump_modify every/time` commands is a variable and not a numeric value, then specifying *first yes* is the only way to write a dump snapshot on the first timestep after the dump command is invoked.

The *flush* keyword determines whether a flush operation is invoked after a dump snapshot is written to the dump file. A flush insures the output in that file is current (no buffering by the OS), even if LAMMPS halts before the simulation completes. Flushes cannot be performed with dump style *xtc*.

The *format* keyword can be used to change the default numeric format output by the text-based dump styles: *atom*, *local*, *custom*, *cfg*, and *xyz* styles, and their MPIIO variants. Only the *line* or *none* options can be used with the *atom* and *xyz* styles.

All the specified format strings are C-style formats, such as used by the C/C++ `printf()` command. The *line* keyword takes a single argument which is the format string for an entire line of output for each atom (do not include a trailing “\n”), with *N* fields, which you must enclose in quotes if there is more than one field. The *int* and *float* keywords take a single format argument and are applied to all integer or floating-point quantities output. The setting for *M string* also takes a single format argument which is used for the *M*th value output in each line (e.g., the fifth column is output in high precision by “format 5 %20.15g”).

**Note:** When using the *line* keyword for the *cfg* style, the first two fields (atom ID and type) are not actually written into the CFG file, however you must include formats for them in the format string.

The *format* keyword can be used multiple times. The precedence is that for each value in a line of output, the *M* format (if specified) is used, else the *int* or *float* setting (if specified) is used, else the *line* setting (if specified) for that value is used, else the default setting is used. A setting of *none* clears all previous settings, reverting all values to their default format.

**Note:** Atom and molecule IDs are stored internally as 4-byte or 8-byte signed integers, depending on how LAMMPS was compiled. When specifying the *format int* option you can use a “%d”-style format identifier in the format string and LAMMPS will convert this to the corresponding 8-byte form if it is needed when outputting those values. However,

when specifying the *line* option or *format M string* option for those values, you should specify a format string appropriate for an 8-byte signed integer (e.g., one with “%ld”) if LAMMPS was compiled with the `-DLAMMPS_BIGBIG` option for 8-byte IDs.

---

**Note:** Any value written to a text-based dump file that is a per-atom quantity calculated by a *compute* or *fix* is stored internally as a floating-point value. If the value is actually an integer and you wish it to appear in the text dump file as a (large) integer, then you need to use an appropriate format. For example, these commands:

---

```
compute      1 all property/local batom1 batom2
dump         1 all local 100 tmp.bonds index c_1[1] c_1[2]
dump_modify  1 format line "%d %0.0f %0.0f"
```

---

will output the two atom IDs for atoms in each bond as integers. If the `dump_modify` command were omitted, they would appear as floating-point values, assuming they were large integers (more than six digits). The “index” keyword should use the “%d” format since it is not generated by a *compute* or *fix*, and is stored internally as an integer.

---

The *fileper* keyword is documented below with the *nfile* keyword.

---

The *header* keyword toggles whether the dump file will include a header. Excluding a header will reduce the size of the dump file for data produced by *pair tracker* or *bpm bond styles* which may not require the information typically written to the header.

---

The *image* keyword applies only to the dump *atom* style. If the *image* value is *yes*, three flags are appended to each atom’s coords which are the absolute box image of the atom in each dimension. For example, an *x* image flag of  $-2$  with a normalized coord of 0.5 means the atom is in the center of the box, but has passed through the box boundary twice and is really two box lengths to the left of its current coordinate. Note that for dump style *custom* these various values can be printed in the dump file by using the appropriate atom attributes in the dump command itself. Using this keyword will reset all custom header names set with *dump\_modify colname* to their respective default values.

---

The *label* keyword applies only to the dump *local* style. When it writes local information, such as bond or angle topology to a dump file, it will use the specified *label* to format the header. By default this includes two lines:

```
ITEM: NUMBER OF ENTRIES
ITEM: ENTRIES ...
```

---

The word “ENTRIES” will be replaced with the string specified (e.g., BONDS or ANGLES).

---

The *maxfiles* keyword can only be used when a “\*” wildcard is included in the dump file name (i.e., when writing a new file(s) for each snapshot). The specified *Fmax* is how many snapshots will be kept. Once this number is reached, the file(s) containing the oldest snapshot is deleted before a new dump file is written. If the specified  $Fmax \leq 0$ , then all files are retained.

This can be useful for debugging, especially if you do not know on what timestep something bad will happen (e.g., when LAMMPS will exit with an error). You can dump every time step and limit the number of dump files produced, even if you run for thousands of steps.

---



The *nfile* or *fileper* keywords can be used in conjunction with the “%” wildcard character in the specified dump file name, for all dump styles except the *dcd*, *image*, *movie*, *xtc*, and *xyz* styles (for which “%” is not allowed). As explained on the *dump* command doc page, the “%” character causes the dump file to be written in pieces, one piece for each of  $P$  processors. By default,  $P$  is the number of processors the simulation is running on. The *nfile* or *fileper* keyword can be used to set  $P$  to a smaller value, which can be more efficient when running on a large number of processors.

The *nfile* keyword sets  $P$  to the specified  $N_f$  value. For example, if  $N_f = 4$ , and the simulation is running on 100 processors, four files will be written by processors 0, 25, 50, and 75. Each will collect information from itself and the next 24 processors and write it to a dump file.

For the *fileper* keyword, the specified value of  $N_p$  means write one file for every  $N_p$  processors. For example, if  $N_p = 4$ , every fourth processor (0, 4, 8, 12, etc.) will collect information from itself and the next three processors and write it to a dump file.

---

The *pad* keyword only applies when the dump filename is specified with a wildcard “\*” character which becomes the timestep. If *pad* is 0, which is the default, the timestep is converted into a string of unpadded length (e.g., 100 or 12000 or 2000000). When *pad* is specified with  $Nchar > 0$ , the string is padded with leading zeroes so they are all the same length =  $Nchar$ . For example, *pad* 7 would yield 0000100, 0012000, 2000000. This can be useful so that post-processing programs can easily read the files in ascending timestep order.

---

The *pbz* keyword applies to all the dump styles. As explained on the *dump* doc page, atom coordinates in a dump file may be slightly outside the simulation box. This is because periodic boundary conditions are enforced only on timesteps when neighbor lists are rebuilt, which will not typically coincide with the timesteps dump snapshots are written. If the setting of this keyword is set to *yes*, then all atoms will be remapped to the periodic box before the snapshot is written, then restored to their original position. If it is set to *no* they will not be. The *no* setting is the default because it requires no extra computation.

---

The *precision* keyword only applies to the dump *xtc* style. A specified value of  $N$  means that coordinates are stored to  $1/N$  nanometer accuracy (e.g., for  $N = 1000$ , the coordinates are written to  $1/1000$  nanometer accuracy).

---

The *refresh* keyword only applies to the dump *custom*, *cfg*, *image*, and *movie* styles. It allows an “incremental” dump file to be written, by refreshing a compute that is used as a threshold for determining which atoms are included in a dump snapshot. The specified *c\_ID* gives the ID of the compute. It is prefixed by “c\_” to indicate a compute, which is the only current option. At some point, other options may be added (e.g., fixes or variables).

---

**Note:** This keyword can only be specified once for a dump. Refreshes of multiple computes cannot yet be performed.

---

The definition and motivation of an incremental dump file is as follows. Instead of outputting all atoms at each snapshot (with some associated values), you may only wish to output the subset of atoms with a value that has changed in some way compared to the value the last time that atom was output. In some scenarios this can result in a dramatically smaller dump file. If desired, by post-processing the sequence of snapshots, the values for all atoms at all timesteps can be inferred.

A concrete example is a simulation of atom diffusion in a solid, represented as atoms on a lattice. Diffusive hops are rare. Imagine that when a hop occurs an atom moves more than a distance  $Dhop$ . For any snapshot we only want to output atoms that have hopped since the last snapshot. This can be accomplished with something the following commands:

```
variable      Dhop equal 0.6
variable      check atom "c_dsp[4] > v_Dhop"
compute       dsp all displace/atom refresh check
dump          1 all custom 20 tmp.dump id type x y z
dump_modify   1 append yes thresh c_dsp[4] > ${Dhop} refresh c_dsp
```

The *compute displace/atom* command calculates the displacement of each atom from its reference position. The “4” index is the scalar displacement; 1, 2, and 3 are the *xyz* components of the displacement. The *dump\_modify thresh* command will cause only atoms that have displaced more than 0.6 Å to be output on a given snapshot (assuming metal units). However, note that when an atom is output, we also need to update the reference position for that atom to its new coordinates. So that it will not be output in every snapshot thereafter. That reference position is stored by *compute displace/atom*. So the *dump\_modify refresh* option triggers a call to *compute displace/atom* at the end of every dump to perform that update. The *refresh check* option shown as part of the *compute displace/atom* command enables the compute to respond to the call from the dump command, and update the appropriate reference positions. This is done by defining an *atom-style variable*, *check* in this example, which calculates a Boolean value (0 or 1) for each atom, based on the same criterion used by *dump\_modify thresh*.

See the *compute displace/atom* command for more details, including an example of how to produce output that includes an initial snapshot with the reference position of all atoms.

Note that only computes with a *refresh* option will work with *dump\_modify refresh*. See individual compute doc pages for details. Currently, only *compute displace/atom* supports this option. Others may be added at some point. If you use a compute that does not support refresh operations, LAMMPS will not complain; *dump\_modify refresh* will simply do nothing.

---

The *region* keyword only applies to the dump *custom*, *cfg*, *image*, and *movie* styles. If specified, only atoms in the region will be written to the dump file or included in the image/movie. Only one region can be applied as a filter (the last one specified). See the *region* command for more details. Note that a region can be defined as the “inside” or “outside” of a geometric shape, and it can be the “union” or “intersection” of a series of simpler regions.

---

The *scale* keyword applies only to the dump *atom* style. A scale value of *yes* means atom coords are written in normalized units from 0.0 to 1.0 in each box dimension. If the simulation box is triclinic (tilted), then all atom coords will still be between 0.0 and 1.0. A value of *no* means they are written in absolute distance units (e.g., Å or  $\sigma$ ). Using this keyword will reset all custom header names set with *dump\_modify colname* to their respective default values.

---

The *sfactor* and *tfactor* keywords only apply to the dump *xtc* style. They allow customization of the unit conversion factors used when writing to XTC files. By default, they are initialized for whatever *units* style is being used, to write out coordinates in nanometers and time in picoseconds. For example, for *real* units, LAMMPS defines *sfactor* = 0.1 and *tfactor* = 0.001, since the Å and fs used by *real* units are 0.1 nm and 0.001 ps, respectively. If you are using a units system with distance and time units far from nm and ps, you may wish to write XTC files with different units, since the compression algorithm used in XTC files is most effective when the typical magnitude of position data is between 10.0 and 0.1.

---

New in version 15Sep2022.

The *skip* keyword can be used with all dump styles. It allows a dump snapshot to be skipped (not written to the dump file), if a condition is met. The condition is computed by an *equal-style variable*, which should be specified as *v\_name*, where name is the variable name. If the variable evaluation returns a non-zero value, then the dump snapshot is skipped. If it returns zero, the dump proceeds as usual. Note that *equal-style variable* can contain Boolean operators which effectively evaluate as a true (non-zero) or false (zero) result.

The *skip* keyword can be useful for debugging purposes, e.g. to dump only on a particular timestep. Or to limit output to conditions of interest, e.g. only when the force on some atom exceeds a threshold value.

The *sort* keyword determines whether lines of per-atom output in a snapshot are sorted or not. A sort value of *off* means they will typically be written in indeterminate order, either in serial or parallel. This is the case even in serial if the *atom\_modify sort* option is turned on, which it is by default, to improve performance. A sort value of *id* means sort the output by atom ID. A sort value of *N* or  $-N$  means sort the output by the value in the *N*th column of per-atom info in either ascending or descending order.

The dump *local* style cannot be sorted by atom ID, since there are typically multiple lines of output per atom. Some dump styles, such as *dcd* and *xtc*, require sorting by atom ID to format the output file correctly. If multiple processors are writing the dump file, via the “%” wildcard in the dump filename and the *nfile* or *fileper* keywords are set to non-default values (i.e., the number of dump file pieces is not equal to the number of procs), then sorting cannot be performed.

In a parallel run, the per-processor dump file pieces can have significant imbalance in number of lines of per-atom info. The *balance* keyword determines whether the number of lines in each processor snapshot are balanced to be nearly the same. A balance value of *no* means no balancing will be done, while *yes* means balancing will be performed. This balancing preserves dump sorting order. For a serial run, this option is ignored since the output is already balanced.

**Note:** Unless it is required by the dump style, sorting dump file output requires extra overhead in terms of CPU and communication cost, as well as memory, versus unsorted output.

The *thermo* keyword only applies the dump styles *netcdf* and *yaml*. It triggers writing of *thermo* information to the dump file alongside per-atom data. The values included in the dump file are identical to the values specified by *thermo\_style*.

The *thresh* keyword only applies to the dump *custom*, *cfg*, *image*, and *movie* styles. Multiple thresholds can be specified. Specifying *none* turns off all threshold criteria. If thresholds are specified, only atoms whose attributes meet all the threshold criteria are written to the dump file or included in the image. The possible attributes that can be tested for are the same as those that can be specified in the *dump custom* command, with the exception of the *element* attribute, since it is not a numeric value. Note that a different attributes can be used than those output by the *dump custom* command. For example, you can output the coordinates and stress of atoms whose energy is above some threshold.

If an atom-style variable is used as the attribute, then it can produce continuous numeric values or effective Boolean 0/1 values, which may be useful for the comparison operator. Boolean values can be generated by variable formulas that use comparison or Boolean math operators or special functions like *gmask()* and *rmask()* and *grmask()*. See the *variable* command page for details.

The specified value must be a simple numeric value or the word LAST. If LAST is used, it refers to the value of the attribute the last time the dump command was invoked to produce a snapshot. This is a way to only dump atoms whose attribute has changed (or not changed). Three examples follow.

```
dump_modify ... thresh ix != LAST
```

This will dump atoms which have crossed the periodic *x* boundary of the simulation box since the last dump. (Note that atoms that crossed once and then crossed back between the two dump timesteps would not be included.)

```
region foo sphere 10 20 10 15
variable inregion atom rmask(foo)
dump_modify ... thresh v_inregion | ^ LAST
```

This will dump atoms which crossed the boundary of the spherical region since the last dump.

```
variable charge atom "(q > 0.5) || (q < -0.5)"
dump_modify ... thresh v_charge |^ LAST
```

This will dump atoms whose charge has changed from an absolute value less than  $\frac{1}{2}$  to greater than  $\frac{1}{2}$  (or vice versa) since the last dump (e.g., due to reactions and subsequent charge equilibration in a reactive force field).

The choice of operators listed above are the usual comparison operators. The XOR operation (exclusive or) is also included as “|^”. In this context, XOR means that if either the attribute or value is 0.0 and the other is non-zero, then the result is “true” and the threshold criterion is met. Otherwise it is not met.

---

The *time* keyword only applies to the dump *atom*, *custom*, *local*, and *xyz* styles (and their COMPRESS package versions *atom/gz*, *custom/gz* and *local/gz*). For the first three styles, if set to *yes*, each frame will contain two extra lines before the “ITEM: TIMESTEP” entry:

```
ITEM: TIME
<elapsed time>
```

For the *xyz* style, the simulation time is included on the same line as the timestep value.

This will output the current elapsed simulation time in current time units equivalent to the *thermo keyword time*. This is to simplify post-processing of trajectories using a variable time step (e.g., when using *fix dt/reset*). The default setting is *no*.

---

The *units* keyword only applies to the dump *atom*, *custom*, and *local* styles (and their COMPRESS package versions *atom/gz*, *custom/gz* and *local/gz*). If set to *yes*, each individual dump file will contain two extra lines at the very beginning with:

```
ITEM: UNITS
<units style>
```

This will output the current selected *units* style to the dump file and thus allows visualization and post-processing tools to determine the choice of units of the data in the dump file. The default setting is *no*.

---

The *unwrap* keyword only applies to the dump *dcd* and *xtc* styles. If set to *yes*, coordinates will be written “unwrapped” by the image flags for each atom. Unwrapped means that if the atom has passed through a periodic boundary one or more times, the value is printed for what the coordinate would be if it had not been wrapped back into the periodic box. Note that these coordinates may thus be far outside the box size stored with the snapshot.

---

The COMPRESS package offers both GZ and Zstd compression variants of styles *atom*, *custom*, *local*, *cfg*, and *xyz*. When using these styles the compression level can be controlled by the *compression\_level* keyword. File names with these styles have to end in either *.gz* or *.zst*.

GZ supports compression levels from  $-1$  (default),  $0$  (no compression), and  $1$  to  $9$ ,  $9$  being the best compression. The COMPRESS /gz styles use  $9$  as default compression level.

Zstd offers a wider range of compression levels, including negative levels that sacrifice compression for performance.  $0$  is the default, positive levels are  $1$  to  $22$ , with  $22$  being the most expensive compression. Zstd promises higher compression/decompression speeds for similar compression ratios. For more details see <http://facebook.github.io/zstd/>.

In addition, Zstd compressed files can include a checksum of the entire contents. The Zstd enabled dump styles enable this feature by default and it can be disabled with the *checksum* keyword.

---

## 16.45.4 Restrictions

Not all *dump\_modify* options can be applied to all dump styles. Details are in the discussions of the individual options.

## 16.45.5 Related commands

*dump*, *dump image*, *undump*

## 16.45.6 Default

The option defaults are

- append = no
- balance = no
- buffer = yes for dump styles *atom*, *custom*, *loca*, and *xyz*
- element = “C” for every atom type
- every = whatever it was set to via the *dump* command
- fileper = # of processors
- first = no
- flush = yes
- format = %d and %g for each integer or floating point value
- image = no
- label = ENTRIES
- maxfiles = -1
- nfile = 1
- pad = 0
- pbc = no
- precision = 1000
- region = none
- scale = yes
- sort = off for dump styles *atom*, *custom*, *cfg*, and *local*
- sort = id for dump styles *dcd*, *xtc*, and *xyz*
- thresh = none
- units = no
- unwrap = no
- compression\_level = 9 (gz variants)
- compression\_level = 0 (zstd variants)
- checksum = yes (zstd variants)

## 16.46 dump molfile command

### 16.46.1 Syntax

```
dump ID group-ID molfile N file format path
```

- ID = user-assigned name for the dump
- group-ID = ID of the group of atoms to be imaged
- molfile = style of dump command (other styles *atom* or *cfg* or *dcd* or *xtc* or *xyz* or *local* or *custom* are discussed on the *dump* doc page)
- N = dump every this many timesteps
- file = name of file to write to
- format = file format to be used
- path = file path with plugins (optional)

### 16.46.2 Examples

```
dump mf1 all molfile 10 melt1.xml hoond  
dump mf2 all molfile 10 melt2-*.pdb .  
dump mf3 all molfile 50 melt3.xyz xyz ./home/akohlmey/vmd/plugins/LINUX/molfile
```

### 16.46.3 Description

Dump a snapshot of atom coordinates and selected additional quantities to one or more files every N timesteps in one of several formats. Only information for atoms in the specified group is dumped. This specific dump style uses molfile plugins that are bundled with the **VMD** molecular visualization and analysis program.

Unless the filename contains a \* character, the output will be written to one single file with the specified format. Otherwise there will be one file per snapshot and the \* will be replaced by the time step number when the snapshot is written.

---

**Note:** Because periodic boundary conditions are enforced only on timesteps when neighbor lists are rebuilt, the coordinates of an atom written to a dump file may be slightly outside the simulation box.

---

The molfile plugin API has a few restrictions that have to be honored by this dump style: the number of atoms must not change, the atoms must be sorted, outside of the coordinates no change in atom properties (like type, mass, charge) will be recorded.

---

The *format* keyword determines what format is used to write out the dump. For this to work, LAMMPS must be able to find and load a compatible molfile plugin that supports this format. Settings made via the *dump\_modify* command can alter per atom properties like element names.

The *path* keyword determines which in directories. This is a “path” like other search paths, i.e. it can contain multiple directories separated by a colon (or semi-colon on windows). This keyword is optional and default to “.”, the current directory.

The *unwrap* option of the *dump\_modify* command allows coordinates to be written “unwrapped” by the image flags for each atom. Unwrapped means that if the atom has passed through a periodic boundary one or more times, the value is printed for what the coordinate would be if it had not been wrapped back into the periodic box. Note that these coordinates may thus be far outside the box size stored with the snapshot.

---

Dumps are performed on timesteps that are a multiple of N (including timestep 0) and on the last timestep of a minimization if the minimization converges. Note that this means a dump will not be performed on the initial timestep after the dump command is invoked, if the current timestep is not a multiple of N. This behavior can be changed via the *dump\_modify first* command, which can be useful if the dump command is invoked after a minimization ended on an arbitrary timestep. N can be changed between runs by using the *dump\_modify every* command. The *dump\_modify every* command also allows a variable to be used to determine the sequence of timesteps on which dump files are written.

---

### 16.46.4 Restrictions

The *molfile* dump style is part of the MOLFILE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Molfile plugins provide a consistent programming interface to read and write file formats commonly used in molecular simulations. The MOLFILE package only provides the interface code, not the plugins. These can be obtained from a VMD installation which has to match the platform that you are using to compile LAMMPS for. By adding plugins to VMD, support for new file formats can be added to LAMMPS (or VMD or other programs that use them) without having to re-compile the application itself. The plugins are installed in the directory: <VMD-HOME>/plugins/<VMDARCH>/molfile

---

**Note:** while the programming interface (API) to the plugins is backward compatible, the binary interface (ABI) has been changing over time, so it is necessary to compile this package with the plugin header files from VMD that match the binary plugins. These header files in the directory: <VMDHOME>/plugins/include For convenience, the package ships with a set of header files that are compatible with VMD 1.9 and 1.9.1 (June 2012)

---

### 16.46.5 Related commands

*dump*, *dump\_modify*, *undump*

### 16.46.6 Default

The default path is “.”. All other properties have to be specified.

## 16.47 dump netcdf command

## 16.48 dump netcdf/mpiio command

### 16.48.1 Syntax

```
dump ID group-ID netcdf N file args
dump ID group-ID netcdf/mpiio N file args
```

- ID = user-assigned name for the dump
- group-ID = ID of the group of atoms to be imaged
- *netcdf* or *netcdf/mpiio* = style of dump command (other styles *atom* or *cfg* or *dcd* or *xtc* or *xyz* or *local* or *custom* are discussed on the *dump* doc page)
- N = dump every this many timesteps
- file = name of file to write dump info to
- args = list of atom attributes, same as for *dump\_style custom*

### 16.48.2 Examples

```
dump 1 all netcdf 100 traj.nc type x y z vx vy vz
dump_modify 1 append yes at -1 thermo yes
dump 1 all netcdf/mpiio 1000 traj.nc id type x y z
dump 1 all netcdf 1000 traj.*.nc id type x y z
```

### 16.48.3 Description

Dump a snapshot of atom coordinates every N timesteps in Amber-style NetCDF file format. NetCDF files are binary, portable and self-describing. This dump style will write only one file on the root node. The dump style *netcdf* uses the [standard NetCDF library](#). All data is collected on one processor and then written to the dump file. Dump style *netcdf/mpiio* uses the [parallel NetCDF library](#) and MPI-IO to write to the dump file in parallel; it has better performance on a larger number of processors. Note that style *netcdf* outputs all atoms sorted by atom tag while style *netcdf/mpiio* outputs atoms in order of their MPI rank.

NetCDF files can be directly visualized via the following tools:

Ovito (<http://www.ovito.org/>). Ovito supports the AMBER convention and all extensions of this dump style.

- VMD (<http://www.ks.uiuc.edu/Research/vmd/>).
- AtomEye (<http://www.libatoms.org/>). The libAtoms version of AtomEye contains a NetCDF reader that is not present in the standard distribution of AtomEye.

In addition to per-atom data, *thermo* data can be included in the dump file. The data included in the dump file is identical to the data specified by *thermo\_style*.



## 16.48.4 Restrictions

The *netcdf* and *netcdf/mpiio* dump styles are part of the NETCDF package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

The *netcdf* and *netcdf/mpiio* dump styles currently cannot dump string properties or properties from variables.

## 16.48.5 Related commands

*dump*, *dump\_modify*, *undump*

## 16.49 dump vtk command

### 16.49.1 Syntax

```
dump ID group-ID vtk N file args
```

- ID = user-assigned name for the dump
- group-ID = ID of the group of atoms to be dumped
- vtk = style of dump command (other styles such as *atom* or *cfg* or *dcd* or *xtc* or *xyz* or *local* or *custom* are discussed on the *dump* doc page)
- N = dump every this many timesteps
- file = name of file to write dump info to
- args = same as arguments for *dump\_style custom*

### 16.49.2 Examples

```
dump dmpvtk all vtk 100 dump*.myforce.vtk id type vx fx
dump dmpvtp flow vtk 100 dump*.*.displace.vtp id type c_myD[1] c_myD[2] c_myD[3] v_ke
```

### 16.49.3 Description

Dump a snapshot of atom quantities to one or more files every *N* timesteps in a format readable by the [VTK visualization toolkit](#) or other visualization tools that use it, such as [ParaView](#). The time steps on which dump output is written can also be controlled by a variable; see the *dump\_modify every* command for details.

This dump style is similar to *dump\_style custom* but uses the VTK library to write data to VTK simple legacy or XML format, depending on the filename extension specified for the dump file. This can be either *\*.vtk* for the legacy format or *\*.vtp* and *\*.vtu*, respectively, for XML format; see the [VTK homepage](#) for a detailed description of these formats. Since this naming convention conflicts with the way binary output is usually specified (see below), the *dump\_modify binary* command allows setting of a binary option for this dump style explicitly.

Only information for atoms in the specified group is dumped. The *dump\_modify thresh and region* commands can also alter what atoms are included; see details below.

As described below, special characters (“\*”, “%”) in the filename determine the kind of output.

**Warning:** Because periodic boundary conditions are enforced only on timesteps when neighbor lists are rebuilt, the coordinates of an atom written to a dump file may be slightly outside the simulation box.

**Warning:** Unless the *dump\_modify sort* option is invoked, the lines of atom information written to dump files will be in an indeterminate order for each snapshot. This is even true when running on a single processor, if the *atom\_modify sort* option is on, which it is by default. In this case atoms are re-ordered periodically during a simulation, due to spatial sorting. It is also true when running in parallel, because data for a single snapshot is collected from multiple processors, each of which owns a subset of the atoms.

For the *vtk* style, sorting is off by default. See the *dump\_modify* page for details.

---

The dimensions of the simulation box are written to a separate file for each snapshot (either in legacy VTK or XML format depending on the format of the main dump file) with the suffix *\_boundingBox* appended to the given dump filename.

For an orthogonal simulation box this information is saved as a rectilinear grid (legacy *.vtk* or *.vtr* XML format).

Triclinic simulation boxes (non-orthogonal) are saved as hexahedrons in either legacy *.vtk* or *.vtu* XML format.

Style *vtk* allows you to specify a list of atom attributes to be written to the dump file for each atom. The list of possible attributes is the same as for the *dump\_style custom* command; see its page for a listing and an explanation of each attribute.

---

**Note:** Since position data is required to write VTK files the atom attributes “x y z” do not have to be specified explicitly; they will be included in the dump file regardless. Also, in contrast to the *custom* style, the specified *vtk* attributes are rearranged to ensure correct ordering of vector components (except for *computes* and *fixes* - these have to be given in the right order) and duplicate entries are removed.

---

The VTK format uses a single snapshot of the system per file, thus a wildcard “\*” must be included in the filename, as discussed below. Otherwise the dump files will get overwritten with the new snapshot each time.

---

Dumps are performed on timesteps that are a multiple of N (including timestep 0) and on the last timestep of a minimization if the minimization converges. Note that this means a dump will not be performed on the initial timestep after the dump command is invoked, if the current timestep is not a multiple of N. This behavior can be changed via the *dump\_modify first* command, which can also be useful if the dump command is invoked after a minimization ended on an arbitrary timestep. N can be changed between runs by using the *dump\_modify every* command. The *dump\_modify every* command also allows a variable to be used to determine the sequence of timesteps on which dump files are written. In this mode a dump on the first timestep of a run will also not be written unless the *dump\_modify first* command is used.

Dump filenames can contain two wildcard characters. If a “\*” character appears in the filename, then one file per snapshot is written and the “\*” character is replaced with the timestep value. For example, *tmp.dump\*.vtk* becomes *tmp.dump0.vtk*, *tmp.dump10000.vtk*, *tmp.dump20000.vtk*, etc. Note that the *dump\_modify pad* command can be used to insure all timestep numbers are the same length (e.g. 00010), which can make it easier to read a series of dump files in order with some post-processing tools.

If a “%” character appears in the filename, then each of P processors writes a portion of the dump file, and the “%” character is replaced with the processor ID from 0 to P-1 preceded by an underscore character. For example, *tmp.dump%.vtp* becomes *tmp.dump\_0.vtp*, *tmp.dump\_1.vtp*, ... *tmp.dump\_P-1.vtp*, etc. This creates smaller files and can be a fast mode of output on parallel machines that support parallel I/O for output.

---

By default,  $P$  = the number of processors meaning one file per processor, but  $P$  can be set to a smaller value via the *nfile* or *fileper* keywords of the *dump\_modify* command. These options can be the most efficient way of writing out dump files when running on large numbers of processors.

For the legacy VTK format “%” is ignored and  $P = 1$ , i.e., only processor 0 does write files.

Note that using the “\*” and “%” characters together can produce a large number of small dump files!

If *dump\_modify binary* is used, the dump file (or files, if “\*” or “%” is also used) is written in binary format. A binary dump file will be about the same size as a text version, but will typically write out much faster.

## 16.49.4 Restrictions

The *vtk* style does not support writing of gzipped dump files.

The *vtk* dump style is part of the VTK package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

To use this dump style, you also must link to the VTK library. See the info in *lib/vtk/README* and insure the *Makefile.lammps* file in that directory is appropriate for your machine.

The *vtk* dump style supports neither buffering or custom format strings.

## 16.49.5 Related commands

*dump*, *dump image*, *dump\_modify*, *undump*

## 16.49.6 Default

By default, files are written in ASCII format. If the file extension is not one of *.vtk*, *.vtp* or *.vtu*, the legacy VTK file format is used.

## 16.50 dynamical\_matrix command

Accelerator Variants: *dynamical\_matrix/kk*

### 16.50.1 Syntax

```
dynamical_matrix group-ID style gamma args keyword value ...
```

- group-ID = ID of group of atoms to displace
- style = *regular* or *eskm*
- gamma = finite different displacement length (distance units)
- one or more keyword/arg pairs may be appended

keyword = *file* or *binary*

*file* name = name of output file for the dynamical matrix

*binary* arg = *yes* or *no* or *gzip*

## 16.50.2 Examples

```
dynamical_matrix 1 regular 0.000001
dynamical_matrix 1 eskm 0.000001
dynamical_matrix 3 regular 0.00004 file dynmat.dat
dynamical_matrix 5 eskm 0.00000001 file dynamical.dat binary yes
```

## 16.50.3 Description

Calculate the dynamical matrix by finite difference of the selected group,

$$D = \frac{\Phi_{ij}^{\alpha\beta}}{\sqrt{M_i M_j}}$$

where D is the dynamical matrix and  $\Phi$  is the force constant matrix defined by

$$\Phi_{ij}^{\alpha\beta} = \frac{\partial^2 U}{\partial x_{i,\alpha} \partial x_{j,\beta}}$$

The output for the dynamical matrix is printed three elements at a time. The three elements are the three  $\beta$  elements for a respective  $i/\alpha/j$  combination. Each line is printed in order of  $j$  increasing first,  $\alpha$  second, and  $i$  last.

If the style eskm is selected, the dynamical matrix will be in units of inverse squared femtoseconds. These units will then conveniently leave frequencies in THz.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 16.50.4 Restrictions

The command collects an array of nine times the number of atoms in a group on every single MPI rank, so the memory requirements can be very significant for large systems.

This command is part of the PHONON package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 16.50.5 Related commands

*fix phonon*, *fix numdiff*,

*compute hma* uses an analytic formulation of the Hessian provided by a pair\_style's Pair::single\_hessian() function, if implemented.

## 16.50.6 Default

The default settings are file = "dynmat.dyn", binary = no

## 16.51 echo command

### 16.51.1 Syntax

```
echo style
```

- style = *none* or *screen* or *log* or *both*

### 16.51.2 Examples

```
echo both  
echo log
```

### 16.51.3 Description

This command determines whether LAMMPS echoes each input script command to the screen and/or log file as it is read and processed. If an input script has errors, it can be useful to look at echoed output to see the last command processed.

The *command-line switch* -echo can be used in place of this command.

### 16.51.4 Restrictions

none

### 16.51.5 Related commands

none

## 16.51.6 Default

```
echo log
```

## 16.52 fix command

### 16.52.1 Syntax

```
fix ID group-ID style args
```

- ID = user-assigned name for the fix
- group-ID = ID of the group of atoms to apply the fix to
- style = one of a long list of possible style names (see below)
- args = arguments used by a particular style

### 16.52.2 Examples

```
fix 1 all nve  
fix 3 all nvt temp 300.0 300.0 0.01  
fix mine top setforce 0.0 NULL 0.0
```

### 16.52.3 Description

Set a fix that will be applied to a group of atoms. In LAMMPS, a “fix” is any operation that is applied to the system during timestepping or minimization. Examples include updating of atom positions and velocities due to time integration, controlling temperature, applying constraint forces to atoms, enforcing boundary conditions, computing diagnostics, etc. There are hundreds of fixes defined in LAMMPS and new ones can be added; see the *Modify* page for details.

Fixes perform their operations at different stages of the timestep. If two or more fixes operate at the same stage of the timestep, they are invoked in the order they were specified in the input script.

The ID of a fix can only contain alphanumeric characters and underscores.

Fixes can be deleted with the *unfix* command.

---

**Note:** The *unfix* command is the only way to turn off a fix; simply specifying a new fix with a similar style will not turn off the first one. This is especially important to realize for integration fixes. For example, using a *fix nve* command for a second run after using a *fix nvt* command for the first run will not cancel out the NVT time integration invoked by the “fix nvt” command. Thus, two time integrators would be in place!

---

If you specify a new fix with the same ID and style as an existing fix, the old fix is deleted and the new one is created (presumably with new settings). This is the same as if an “unfix” command were first performed on the old fix, except that the new fix is kept in the same order relative to the existing fixes as the old one originally was. Note that this operation also wipes out any additional changes made to the old fix via the *fix\_modify* command.

The *fix\_modify* command allows settings for some fixes to be reset. See the page for individual fixes for details.

Some fixes store an internal “state” which is written to binary restart files via the *restart* or *write\_restart* commands. This allows the fix to continue on with its calculations in a restarted simulation. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file. See the doc pages for individual fixes for info on which ones can be restarted.

Some fixes calculate one of three styles of quantities: global, per-atom, or local, which can be used by other commands or output as described below. A global quantity is one or more system-wide values (e.g., the energy of a wall interacting with particles). A per-atom quantity is one or more values per atom (e.g., the displacement vector for each atom since time 0). Per-atom values are set to 0.0 for atoms not in the specified fix group. Local quantities are calculated by each processor based on the atoms it owns, but there may be zero or more per atoms.

Note that a single fix can produce either global or per-atom or local quantities (or none at all), but not both global and per-atom. It can produce local quantities in tandem with global or per-atom quantities. The fix page will explain.

Global, per-atom, and local quantities each come in three kinds: a single scalar value, a vector of values, or a 2d array of values. The doc page for each fix describes the style and kind of values it produces (e.g., a per-atom vector). Some fixes produce more than one kind of a single style (e.g., a global scalar and a global vector).

When a fix quantity is accessed, as in many of the output commands discussed below, it can be referenced via the following bracket notation, where ID is the ID of the fix:

f_ID	entire scalar, vector, or array
f_ID[I]	one element of vector, one column of array
f_ID[I][J]	one element of array

In other words, using one bracket reduces the dimension of the quantity once (vector → scalar, array → vector). Using two brackets reduces the dimension twice (array → scalar). Thus, a command that uses scalar fix values as input can also process elements of a vector or array.

Note that commands and *variables* that use fix quantities typically do not allow for all kinds (e.g., a command may require a vector of values, not a scalar), and even if they do, the context in which they are called can be used to resolve which output is being requested. This means there is no ambiguity about referring to a fix quantity as f\_ID even if it produces, for example, both a scalar and vector. The doc pages for various commands explain the details.

In LAMMPS, the values generated by a fix can be used in several ways:

- Global values can be output via the *thermo\_style custom* or *fix ave/time* command. Alternatively, the values can be referenced in an *equal-style variable* command.
- Per-atom values can be output via the *dump custom* command, or they can be time-averaged via the *fix ave/atom* command or reduced by the *compute reduce* command. Alternatively, per-atom values can be referenced in an *atom-style variable*.
- Local values can be reduced by the *compute reduce* command or histogrammed by the *fix ave/histo* command. They can also be output by the *dump local* command.

See the *Howto output* page for a summary of various LAMMPS output options, many of which involve fixes.

The results of fixes that calculate global quantities can be either “intensive” or “extensive” values. Intensive means the value is independent of the number of atoms in the simulation (e.g., temperature). Extensive means the value scales with the number of atoms in the simulation (e.g., total rotational kinetic energy). *Thermodynamic output* will normalize extensive values by the number of atoms in the system, depending on the “thermo\_modify norm” setting. It will not normalize intensive values. If a fix value is accessed in another way (e.g., by a *variable*), you may want to know whether it is an intensive or extensive value. See the page for individual fix styles for further info.

Each fix style has its own page that describes its arguments and what it does, as listed below. Here is an alphabetical list of fix styles available in LAMMPS. They are also listed in more compact form on the [Commands fix](#) doc page.

There are also additional accelerated fix styles included in the LAMMPS distribution for faster performance on CPUs, GPUs, and KNLs. The individual style names on the [Commands fix](#) doc page are followed by one or more of (g,i,k,o,t) to indicate which accelerated styles exist.

- *accelerate/cos* - apply cosine-shaped acceleration to atoms
- *acks2/reaxff* - apply ACKS2 charge equilibration
- *adapt* - change a simulation parameter over time
- *adapt/fep* - enhanced version of fix adapt
- *addforce* - add a force to each atom
- *addtorque* - add a torque to a group of atoms
- *amoeba/bitorsion* - torsion/torsion terms in AMOEBA force field
- *amoeba/pitorsion* - 6-body terms in AMOEBA force field
- *append/atoms* - append atoms to a running simulation
- *atc* - initiates a coupled MD/FE simulation
- *atom/swap* - Monte Carlo atom type swapping
- *ave/atom* - compute per-atom time-averaged quantities
- *ave/chunk* - compute per-chunk time-averaged quantities
- *ave/correlate* - compute/output time correlations
- *ave/correlate/long* - alternative to *ave/correlate* that allows efficient calculation over long time windows
- *ave/histo* - compute/output time-averaged histograms
- *ave/histo/weight* - weighted version of fix ave/histo
- *ave/time* - compute/output global time-averaged quantities
- *aveforce* - add an averaged force to each atom
- *balance* - perform dynamic load-balancing
- *brownian* - overdamped translational brownian motion
- *brownian/asphere* - overdamped translational and rotational brownian motion for ellipsoids
- *brownian/sphere* - overdamped translational and rotational brownian motion for spheres
- *bocs* - NPT style time integration with pressure correction
- *bond/break* - break bonds on the fly
- *bond/create* - create bonds on the fly
- *bond/create/angle* - create bonds on the fly with angle constraints
- *bond/react* - apply topology changes to model reactions
- *bond/swap* - Monte Carlo bond swapping
- *box/relax* - relax box size during energy minimization
- *charge/regulation* - Monte Carlo sampling of charge regulation
- *cmap* - CMAP torsion/torsion terms in CHARMM force field



- *colvars* - interface to the collective variables “Colvars” library
- *controller* - apply control loop feedback mechanism
- *damping/cundall* - Cundall non-viscous damping for granular simulations
- *deform* - change the simulation box size/shape
- *deposit* - add new atoms above a surface
- *dpd/energy* - constant energy dissipative particle dynamics
- *drag* - drag atoms towards a defined coordinate
- *drude* - part of Drude oscillator polarization model
- *drude/transform/direct* - part of Drude oscillator polarization model
- *drude/transform/inverse* - part of Drude oscillator polarization model
- *dt/reset* - reset the timestep based on velocity, forces
- *edpd/source* - add heat source to eDPD simulations
- *efield* - impose electric field on system
- *ehex* - enhanced heat exchange algorithm
- *electrode/conp* - impose electric potential
- *electrode/conq* - impose total electric charge
- *electrode/thermo* - apply thermo-potentiostat
- *electron/stopping* - electronic stopping power as a friction force
- *electron/stopping/fit* - electronic stopping power as a friction force
- *enforce2d* - zero out z-dimension velocity and force
- *eos/cv* - applies a mesoparticle equation of state to relate the particle internal energy to the particle internal temperature
- *eos/table* - applies a tabulated mesoparticle equation of state to relate the particle internal energy to the particle internal temperature
- *eos/table/rx* - applies a tabulated mesoparticle equation of state to relate the concentration-dependent particle internal energy to the particle internal temperature
- *evaporate* - remove atoms from simulation periodically
- *external* - callback to an external driver program
- *ffl* - apply a Fast-Forward Langevin equation thermostat
- *filter/corotate* - implement corotation filter to allow larger timesteps with r-RESPA
- *flow/gauss* - Gaussian dynamics for constant mass flux
- *freeze* - freeze atoms in a granular simulation
- *gcmc* - grand canonical insertions/deletions
- *gld* - generalized Langevin dynamics integrator
- *gle* - generalized Langevin equation thermostat
- *gravity* - add gravity to atoms in a granular simulation
- *grem* - implements the generalized replica exchange method

- *halt* - terminate a dynamics run or minimization
- *heat* - add/subtract momentum-conserving heat
- *hyper/global* - global hyperdynamics
- *hyper/local* - local hyperdynamics
- *imd* - implements the “Interactive MD” (IMD) protocol
- *indent* - impose force due to an indenter
- *ipi* - enable LAMMPS to run as a client for i-PI path-integral simulations
- *langevin* - Langevin temperature control
- *langevin/drude* - Langevin temperature control of Drude oscillators
- *langevin/eff* - Langevin temperature control for the electron force field model
- *langevin/spin* - Langevin temperature control for a spin or spin-lattice system
- *latte* - wrapper on LATTE density-functional tight-binding code
- *lb/fluid* - lattice-Boltzmann fluid on a uniform mesh
- *lb/momentum* - *fix momentum* replacement for use with a lattice-Boltzmann fluid
- *lb/viscous* - *fix viscous* replacement for use with a lattice-Boltzmann fluid
- *lineforce* - constrain atoms to move in a line
- *manifoldforce* - restrain atoms to a manifold during minimization
- *mdi/qm* - LAMMPS operates as driver for a quantum code via the MolSSI Driver Interface (MDI)
- *meso/move* - move mesoscopic SPH/SDPD particles in a prescribed fashion
- *mol/swap* - Monte Carlo atom type swapping with a molecule
- *momentum* - zero the linear and/or angular momentum of a group of atoms
- *momentum/chunk* - zero the linear and/or angular momentum of a chunk of atoms
- *move* - move atoms in a prescribed fashion
- *mscg* - apply MSCG method for force-matching to generate coarse grain models
- *msst* - multi-scale shock technique (MSST) integration
- *mvv/dpd* - DPD using the modified velocity-Verlet integration algorithm
- *mvv/edpd* - constant energy DPD using the modified velocity-Verlet algorithm
- *mvv/tdpd* - constant temperature DPD using the modified velocity-Verlet algorithm
- *neb* - nudged elastic band (NEB) spring forces
- *neb/spin* - nudged elastic band (NEB) spring forces for spins
- *nph* - constant NPH time integration via Nose/Hoover
- *nph/asphere* - NPH for aspherical particles
- *nph/body* - NPH for body particles
- *nph/eff* - NPH for nuclei and electrons in the electron force field model
- *nph/sphere* - NPH for spherical particles
- *nphug* - constant-stress Hugoniotat integration

- *npt* - constant NPT time integration via Nose/Hoover
- *npt/asphere* - NPT for aspherical particles
- *npt/body* - NPT for body particles
- *npt/cauchy* - NPT with Cauchy stress
- *npt/eff* - NPT for nuclei and electrons in the electron force field model
- *npt/sphere* - NPT for spherical particles
- *npt/uef* - NPT style time integration with diagonal flow
- *numdiff* - numerically approximate atomic forces using finite energy differences
- *numdiff/virial* - numerically approximate virial stress tensor using finite energy differences
- *nve* - constant NVE time integration
- *nve/asphere* - NVE for aspherical particles
- *nve/asphere/noforce* - NVE for aspherical particles without forces
- *nve/awpmd* - NVE for the Antisymmetrized Wave Packet Molecular Dynamics model
- *nve/body* - NVE for body particles
- *nve/dot* - rigid body constant energy time integrator for coarse grain models
- *nve/dotc/langevin* - Langevin style rigid body time integrator for coarse grain models
- *nve/eff* - NVE for nuclei and electrons in the electron force field model
- *nve/limit* - NVE with limited step length
- *nve/line* - NVE for line segments
- *nve/manifold/rattle* - NVE time integration for atoms constrained to a curved surface (manifold)
- *nve/noforce* - NVE without forces (update positions only)
- *nve/sphere* - NVE for spherical particles
- *nve/bpm/sphere* - NVE for spherical particles used in the BPM package
- *nve/spin* - NVE for a spin or spin-lattice system
- *nve/tri* - NVE for triangles
- *nvk* - constant kinetic energy time integration
- *nvt* - NVT time integration via Nose/Hoover
- *nvt/asphere* - NVT for aspherical particles
- *nvt/body* - NVT for body particles
- *nvt/eff* - NVE for nuclei and electrons in the electron force field model
- *nvt/manifold/rattle* - NVT time integration for atoms constrained to a curved surface (manifold)
- *nvt/sllod* - NVT for NEMD with SLLD equations
- *nvt/sllod/eff* - NVT for NEMD with SLLD equations for the electron force field model
- *nvt/sphere* - NVT for spherical particles
- *nvt/uef* - NVT style time integration with diagonal flow
- *oneway* - constrain particles on move in one direction

- *orient/bcc* - add grain boundary migration force for BCC
- *orient/fcc* - add grain boundary migration force for FCC
- *orient/eco* - add generalized grain boundary migration force
- *pafi* - constrained force averages on hyper-planes to compute free energies (PAFI)
- *pair* - access per-atom info from pair styles
- *phonon* - calculate dynamical matrix from MD simulations
- *pimd* - Feynman path integral molecular dynamics
- *planeforce* - constrain atoms to move in a plane
- *plumed* - wrapper on PLUMED free energy library
- *poems* - constrain clusters of atoms to move as coupled rigid bodies
- *polarize/bem/gmres* - compute induced charges at the interface between impermeable media with different dielectric constants with generalized minimum residual (GMRES)
- *polarize/bem/icc* - compute induced charges at the interface between impermeable media with different dielectric constants with the successive over-relaxation algorithm
- *polarize/functional* - compute induced charges at the interface between impermeable media with different dielectric constants with the energy variational approach
- *pour* - pour new atoms/molecules into a granular simulation domain
- *precession/spin* - apply a precession torque to each magnetic spin
- *press/berendsen* - pressure control by Berendsen barostat
- *print* - print text and variables during a simulation
- *propel/self* - model self-propelled particles
- *property/atom* - add customized per-atom values
- *python/invoke* - call a Python function during a simulation
- *python/move* - move particles using a Python function during a simulation run
- *qbmsst* - quantum bath multi-scale shock technique time integrator
- *qeql/comb* - charge equilibration for COMB potential
- *qeql/dynamic* - charge equilibration via dynamic method
- *qeql/fire* - charge equilibration via FIRE minimizer
- *qeql/point* - charge equilibration via point method
- *qeql/reaxff* - charge equilibration for ReaxFF potential
- *qeql/shielded* - charge equilibration via shielded method
- *qeql/slater* - charge equilibration via Slater method
- *qmmm* - functionality to enable a quantum mechanics/molecular mechanics coupling
- *qtb* - implement quantum thermal bath scheme
- *rattle* - RATTLE constraints on bonds and/or angles
- *reaxff/bonds* - write out ReaxFF bond information
- *reaxff/species* - write out ReaxFF molecule information

- *recenter* - constrain the center-of-mass position of a group of atoms
- *restrain* - constrain a bond, angle, dihedral
- *rhok* - add bias potential for long-range ordered systems
- *rigid* - constrain one or more clusters of atoms to move as a rigid body with NVE integration
- *rigid/meso* - constrain clusters of mesoscopic SPH/SDPD particles to move as a rigid body
- *rigid/nph* - constrain one or more clusters of atoms to move as a rigid body with NPH integration
- *rigid/nph/small* - constrain many small clusters of atoms to move as a rigid body with NPH integration
- *rigid/npt* - constrain one or more clusters of atoms to move as a rigid body with NPT integration
- *rigid/npt/small* - constrain many small clusters of atoms to move as a rigid body with NPT integration
- *rigid/nve* - constrain one or more clusters of atoms to move as a rigid body with alternate NVE integration
- *rigid/nve/small* - constrain many small clusters of atoms to move as a rigid body with alternate NVE integration
- *rigid/nvt* - constrain one or more clusters of atoms to move as a rigid body with NVT integration
- *rigid/nvt/small* - constrain many small clusters of atoms to move as a rigid body with NVT integration
- *rigid/small* - constrain many small clusters of atoms to move as a rigid body with NVE integration
- *rx* - solve reaction kinetic ODEs for a defined reaction set
- *saed/vtk* - time-average the intensities from *compute saed*
- *setforce* - set the force on each atom
- *setforce/spin* - set magnetic precession vectors on each atom
- *shake* - SHAKE constraints on bonds and/or angles
- *shardlow* - integration of DPD equations of motion using the Shardlow splitting
- *smd* - applied a steered MD force to a group
- *smd/adjust\_dt* - calculate a new stable time increment for use with SMD integrators
- *smd/integrate\_tlsph* - explicit time integration with total Lagrangian SPH pair style
- *smd/integrate\_ulsph* - explicit time integration with updated Lagrangian SPH pair style
- *smd/move\_tri\_surf* - update position and velocity near rigid surfaces using SPH integrators
- *smd/setvel* - sets each velocity component, ignoring forces, for Smooth Mach Dynamics
- *smd/wall\_surface* - create a rigid wall with a triangulated surface for use in Smooth Mach Dynamics
- *sph* - time integration for SPH/DPDE particles
- *sph/stationary* - update energy and density but not position or velocity in Smooth Particle Hydrodynamics
- *spring* - apply harmonic spring force to group of atoms
- *spring/chunk* - apply harmonic spring force to each chunk of atoms
- *spring/rg* - spring on radius of gyration of group of atoms
- *spring/self* - spring from each atom to its origin
- *srd* - stochastic rotation dynamics (SRD)
- *store/force* - store force on each atom
- *store/state* - store attributes for each atom

- *tdpd/source* - add external concentration source
- *temp/berendsen* - temperature control by Berendsen thermostat
- *temp/csld* - canonical sampling thermostat with Langevin dynamics
- *temp/csvr* - canonical sampling thermostat with Hamiltonian dynamics
- *temp/rescale* - temperature control by velocity rescaling
- *temp/rescale/eff* - temperature control by velocity rescaling in the electron force field model
- *tfmc* - perform force-bias Monte Carlo with time-stamped method
- *tgnt/drude* - NVT time integration for Drude polarizable model via temperature-grouped Nose-Hoover
- *tgnt/drude* - NPT time integration for Drude polarizable model via temperature-grouped Nose-Hoover
- *thermal/conductivity* - Mueller-Plathe kinetic energy exchange for thermal conductivity calculation
- *ti/spring* - perform thermodynamic integration between a solid and an Einstein crystal
- *tmd* - guide a group of atoms to a new configuration
- *tmm* - two-temperature model for electronic/atomic coupling (replicated grid)
- *tmm/grid* - two-temperature model for electronic/atomic coupling (distributed grid)
- *tmm/mod* - enhanced two-temperature model with additional options
- *tune/kpace* - auto-tune *k*-space parameters
- *vector* - accumulate a global vector every *N* timesteps
- *viscosity* - Mueller-Plathe momentum exchange for viscosity calculation
- *viscous* - viscous damping for granular simulations
- *viscous/sphere* - viscous damping on angular velocity for granular simulations
- *wall/body/polygon* - time integration for body particles of style *rounded/polygon*
- *wall/body/polyhedron* - time integration for body particles of style *rounded/polyhedron*
- *wall/colloid* - Lennard-Jones wall interacting with finite-size particles
- *wall/ees* - wall for ellipsoidal particles
- *wall/gran* - frictional wall(s) for granular simulations
- *wall/gran/region* - *fix wall/region* equivalent for use with granular particles
- *wall/harmonic* - harmonic spring wall
- *wall/lj1043* - Lennard-Jones 10–4–3 wall
- *wall/lj126* - Lennard-Jones 12–6 wall
- *wall/lj93* - Lennard-Jones 9–3 wall
- *wall/morse* - Morse potential wall
- *wall/piston* - moving reflective piston wall
- *wall/reflect* - reflecting wall(s)
- *wall/reflect/stochastic* - reflecting wall(s) with finite temperature
- *wall/region* - use region surface as wall
- *wall/region/ees* - use region surface as wall for ellipsoidal particles

- *wall/srd* - slip/no-slip wall for SRD particles
- *widom* - Widom insertions of atoms or molecules

## 16.52.4 Restrictions

Some fix styles are part of specific packages. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info. The doc pages for individual fixes tell if it is part of a package.

## 16.52.5 Related commands

*unfix*, *fix\_modify*

## 16.52.6 Default

none

# 16.53 *fix\_modify* command

## 16.53.1 Syntax

```
fix_modify fix-ID keyword value ...
```

- fix-ID = ID of the fix to modify
- one or more keyword/value pairs may be appended
- keyword = *bodyforces* or *colname* or *dynamic/dof* or *energy* or *press* or *respa* or *temp* or *virial*
  - bodyforces* value = *early* or *late*  
early/late = compute rigid-body forces/torques early or late in the timestep
  - colname* values = ID string  
string = new column header name  
ID = integer from 1 to N, or integer from -1 to -N, where N = # of quantities.  
→being output  
or a fix output property keyword or reference to compute, fix, property or  
→variable.
  - dynamic/dof* value = *yes* or *no*  
yes/no = do or do not re-compute the number of degrees of freedom (DOF).  
→contributing to the temperature
  - energy* value = *yes* or *no*
  - press* value = compute ID that calculates a pressure
  - respa* value = 1 to *max respa level* or 0 (for outermost level)
  - temp* value = compute ID that calculates a temperature
  - virial* value = *yes* or *no*

## 16.53.2 Examples

```
fix_modify 3 temp myTemp press myPress
fix_modify 1 energy yes
fix_modify tether respa 2
fix_modify ave colname c_thermo_press Pressure colname 1 Temperature
```

## 16.53.3 Description

Modify one or more parameters of a previously defined fix. Only specific fix styles support specific parameters. See the doc pages for individual fix commands for info on which ones support which `fix_modify` parameters.

The `temp` keyword is used to determine how a fix computes temperature. The specified compute ID must have been previously defined by the user via the `compute` command and it must be a style of compute that calculates a temperature. All fixes that compute temperatures define their own compute by default, as described in their documentation. Thus this option allows the user to override the default method for computing T.

The `press` keyword is used to determine how a fix computes pressure. The specified compute ID must have been previously defined by the user via the `compute` command and it must be a style of compute that calculates a pressure. All fixes that compute pressures define their own compute by default, as described in their documentation. Thus this option allows the user to override the default method for computing P.

The `energy` keyword can be used with fixes that support it, which is explained at the bottom of their doc page. *Energy yes* will add a contribution to the potential energy of the system. More specifically, the fix's global or per-atom energy is included in the calculation performed by the `compute pe` or `compute pe/atom` commands. The former is what is used the `thermo_style` command for output of any quantity that includes the global potential energy of the system. Note that the `compute pe` and `compute pe/atom` commands also have an option to include or exclude the contribution from fixes. For fixes that tally a global energy, it can also be printed with thermodynamic output by using the keyword `f_ID` in the `thermo_style` custom command, where ID is the fix-ID of the appropriate fix.

---

**Note:** If you are performing an *energy minimization* with one of these fixes and want the energy and forces it produces to be part of the optimization criteria, you must specify the *energy yes* setting.

---

---

**Note:** For most fixes that support the *energy* keyword, the default setting is *no*. For a few it is *yes*, when a user would expect that to be the case. The page of each fix gives the default.

---

The *virial* keyword can be used with fixes that support it, which is explained at the bottom of their doc page. *Virial yes* will add a contribution to the virial of the system. More specifically, the fix's global or per-atom virial is included in the calculation performed by the `compute pressure` or `compute stress/atom` commands. The former is what is used the `thermo_style` command for output of any quantity that includes the global pressure of the system. Note that the `compute pressure` and `compute stress/atom` commands also have an option to include or exclude the contribution from fixes.

---

**Note:** If you are performing an *energy minimization* with *box relaxation* and one of these fixes and want the virial contribution of the fix to be part of the optimization criteria, you must specify the *virial yes* setting.

---

---

**Note:** For most fixes that support the *virial* keyword, the default setting is *no*. For a few it is *yes*, when a user would expect that to be the case. The page of each fix gives the default.

---



For fixes that set or modify forces, it may be possible to select at which *r-RESPA* level the fix operates via the *respa* keyword. The RESPA level at which the fix is active can be selected. This is a number ranging from 1 to the number of levels. If the RESPA level is larger than the current maximum, the outermost level will be used, which is also the default setting. This default can be restored using a value of 0 for the RESPA level. The affected fix has to be enabled to support this feature; if not, *fix\_modify* will report an error. Active fixes with a custom RESPA level setting are reported with their specified level at the beginning of a r-RESPA run.

The *dynamic/dof* keyword determines whether the number of atoms *N* in the fix group and their associated degrees of freedom are re-computed each time a temperature is computed. Only fix styles that calculate their own internal temperature use this option. Currently this is only the *fix rigid/nvt/small* and *fix rigid/npt/small* commands for the purpose of thermostating rigid body translation and rotation. By default, *N* and their DOF are assumed to be constant. If you are adding atoms or molecules to the system (see the *fix pour*, *fix deposit*, and *fix gcmc* commands) or expect atoms or molecules to be lost (e.g. due to exiting the simulation box or via *fix evaporate*), then this option should be used to insure the temperature is correctly normalized.

---

**Note:** Other thermostating fixes, such as *fix nvt*, do not use the *dynamic/dof* keyword because they use a temperature compute to calculate temperature. See the *compute\_modify dynamic/dof* command for a similar way to insure correct temperature normalization for those thermostats.

---

The *bodyforces* keyword determines whether the forces and torques acting on rigid bodies are computed *early* at the post-force stage of each timestep (right after per-atom forces have been computed and communicated among processors), or *late* at the final-integrate stage of each timestep (after any other fixes have finished their post-force tasks). Only the rigid-body integration fixes use this option, which includes *fix rigid* and *fix rigid/small*, and their variants, and also *fix poems*.

The default is *late*. If there are other fixes that add forces to individual atoms, then the rigid-body constraints will include these forces when time-integrating the rigid bodies. If *early* is specified, then new fixes can be written that use or modify the per-body force and torque, before time-integration of the rigid bodies occurs. Note however this has the side effect, that fixes such as *fix addforce*, *fix setforce*, *fix spring*, which add forces to individual atoms will have no effect on the motion of the rigid bodies if they are specified in the input script after the *fix rigid* command. LAMMPS will give a warning if that is the case.

The *colname* keyword can be used to change the default header keywords in output files of fix styles that support it: currently only *fix ave/time* is supported. The setting for *ID string* replaces the default text with the provided string. *ID* can be a positive integer when it represents the column number counting from the left, a negative integer when it represents the column number from the right (i.e. -1 is the last column/keyword), or a custom fix output keyword (or compute, fix, property, or variable reference) and then it replaces the string for that specific keyword. The *colname* keyword can be used multiple times. If multiple *colname* settings refer to the same keyword, the last setting has precedence.

## 16.53.4 Restrictions

none

## 16.53.5 Related commands

*fix*, *compute temp*, *compute pressure*, *thermo\_style*

## 16.53.6 Default

The option defaults are temp = ID defined by fix, press = ID defined by fix, energy = no, virial = different for each fix style, respa = 0, bodyforce = late.

# 16.54 group command

## 16.54.1 Syntax

```
group ID style args
```

- ID = user-defined name of the group
- style = *delete* or *clear* or *empty* or *region* or *type* or *id* or *molecule* or *variable* or *include* or *subtract* or *union* or *intersect* or *dynamic* or *static*

*delete* = no args

*clear* = no args

*empty* = no args

*region* args = region-ID

*type* or *id* or *molecule*

args = list of one or more atom types, atom IDs, or molecule IDs

any entry in list can be a sequence formatted as A:B or A:B:C where

A = starting index, B = ending index,

C = increment between indices, 1 if not specified

args = logical value

logical = "<" or "<=" or ">" or ">=" or "==" or "!="

value = an atom type or atom ID or molecule ID (depending on style)

args = logical value1 value2

logical = "<>"

value1,value2 = atom types or atom IDs or molecule IDs (depending on style)

*variable* args = variable-name

*include* args = molecule

molecule = add atoms to group with same molecule ID as atoms already in group

*subtract* args = two or more group IDs

*union* args = one or more group IDs

*intersect* args = two or more group IDs

*dynamic* args = parent-ID keyword value ...

one or more keyword/value pairs may be appended

keyword = *region* or *var* or *property* or *every*

*region* value = region-ID

*var* value = name of variable

*property* value = name of custom integer or floating point vector

every value = N = update group every this many timesteps  
 static = no args

## 16.54.2 Examples

```
group edge region regstrip
group water type 3 4
group sub id 10 25 50
group sub id 10 25 50 500:1000
group sub id 100:10000:10
group sub id <= 150
group polyA molecule <> 50 250
group hienergy variable eng
group hienergy include molecule
group boundary subtract all a2 a3
group boundary union lower upper
group boundary intersect upper flow
group boundary delete
group mine dynamic all region myRegion every 100
```

## 16.54.3 Description

Identify a collection of atoms as belonging to a group. The group ID can then be used in other commands such as *fix*, *compute*, *dump*, or *velocity* to act on those atoms together.

If the group ID already exists, the group command adds the specified atoms to the group.

---

**Note:** By default groups are static, meaning the atoms are permanently assigned to the group. For example, if the *region* style is used to assign atoms to a group, the atoms will remain in the group even if they later move out of the region. As explained below, the *dynamic* style can be used to make a group dynamic so that a periodic determination is made as to which atoms are in the group. Since many LAMMPS commands operate on groups of atoms, you should think carefully about whether making a group dynamic makes sense for your model.

---

A group with the ID *all* is predefined. All atoms belong to this group. This group cannot be deleted, or made dynamic.

The *delete* style removes the named group and un-assigns all atoms that were assigned to that group. Since there is a restriction (see below) that no more than 32 groups can be defined at any time, the *delete* style allows you to remove groups that are no longer needed, so that more can be specified. You cannot delete a group if it has been used to define a current *fix* or *compute* or *dump*.

The *clear* style un-assigns all atoms that were assigned to that group. This may be dangerous to do during a simulation run (e.g., using the *run every* command if a *fix* or *compute* or other operation expects the atoms in the group to remain constant), but LAMMPS does not check for this.

The *empty* style creates an empty group, which is useful for commands like *fix gcmc* or with complex scripts that add atoms to a group.

The *region* style puts all atoms in the region volume into the group. Note that this is a static one-time assignment. The atoms remain assigned (or not assigned) to the group even in they later move out of the region volume.

The *type*, *id*, and *molecule* styles put all atoms with the specified atom types, atom IDs, or molecule IDs into the group. These three styles can use arguments specified in one of two formats.

The first format is a list of values (types or IDs). For example, the second command in the examples above puts all atoms of type 3 or 4 into the group named *water*. Each entry in the list can be a colon-separated sequence A:B or A:B:C, as in two of the examples above. A “sequence” generates a sequence of values (types or IDs), with an optional increment. The first example with 500:1000 has the default increment of 1 and would add all atom IDs from 500 to 1000 (inclusive) to the group *sub*, along with 10, 25, and 50 since they also appear in the list of values. The second example with 100:10000:10 uses an increment of 10 and would thus would add atoms IDs 100, 110, 120, . . . , 9990, 10000 to the group *sub*.

The second format is a *logical* followed by one or two values (type or ID). The 7 valid logicals are listed above. All the logicals except <> take a single argument. The third example above adds all atoms with IDs from 1 to 150 to the group named *sub*. The logical <> means “between” and takes 2 arguments. The fourth example above adds all atoms belonging to molecules with IDs from 50 to 250 (inclusive) to the group named *polyA*.

The *variable* style evaluates a variable to determine which atoms to add to the group. It must be an *atom-style variable* previously defined in the input script. If the variable evaluates to a non-zero value for a particular atom, then that atom is added to the specified group.

Atom-style variables can specify formulas that include thermodynamic quantities, per-atom values such as atom coordinates, or per-atom quantities calculated by computes, fixes, or other variables. They can also include Boolean logic where two numeric values are compared to yield a 1 or 0 (effectively a true or false). Thus, using the *variable* style is a general way to flag specific atoms to include or exclude from a group.

For example, these lines define a variable “*eatom*” that calculates the potential energy of each atom and includes it in the group if its potential energy is above the threshold value  $-3.0$ .

```
compute      1 all pe/atom
compute      2 all reduce sum c_1
thermo_style custom step temp pe c_2
run          0

variable     eatom atom "c_1 > -3.0"
group        hienergy variable eatom
```

Note that these lines

```
compute      2 all reduce sum c_1
thermo_style custom step temp pe c_2
run          0
```

are necessary to insure that the “*eatom*” variable is current when the group command invokes it. Because the *eatom* variable computes the per-atom energy via the *pe/atom* compute, it will only be current if a run has been performed which evaluated pairwise energies, and the *pe/atom* compute was actually invoked during the run. Printing the thermodynamic info for compute 2 insures that this is the case, since it sums the *pe/atom* compute values (in the reduce compute) to output them to the screen. See the “Variable Accuracy” section of the *variable* page for more details on insuring that variables are current when they are evaluated between runs.

The *include* style with its arg *molecule* adds atoms to a group that have the same molecule ID as atoms already in the group. The molecule ID = 0 is ignored in this operation, since it is assumed to flag isolated atoms that are not part of molecules. An example of where this operation is useful is if the *region* style has been used previously to add atoms to a group that are within a geometric region. If molecules straddle the region boundary, then atoms outside the region that are part of molecules with atoms inside the region will not be in the group. Using the group command a second time with *include molecule* will add those atoms that are outside the region to the group.

**Note:** The *include molecule* operation is relatively expensive in a parallel sense. This is because it requires communication of relevant molecule IDs between all the processors and each processor to loop over its atoms once per processor,

to compare its atoms to the list of molecule IDs from every other processor. Hence it scales as  $N$ , rather than  $N/P$  as most of the group operations do, where  $N$  is the number of atoms, and  $P$  is the number of processors.

---

The *subtract* style takes a list of two or more existing group names as arguments. All atoms that belong to the first group, but not to any of the other groups are added to the specified group.

The *union* style takes a list of one or more existing group names as arguments. All atoms that belong to any of the listed groups are added to the specified group.

The *intersect* style takes a list of two or more existing group names as arguments. Atoms that belong to every one of the listed groups are added to the specified group.

---

The *dynamic* style flags an existing or new group as dynamic. This means atoms will be (re)assigned to the group periodically as a simulation runs. This is in contrast to static groups where atoms are permanently assigned to the group. The way the assignment occurs is as follows. Only atoms in the group specified as the parent group via the parent-ID are assigned to the dynamic group before the following conditions are applied.

If the *region* keyword is used, atoms not in the specified region are removed from the dynamic group.

If the *var* keyword is used, the variable name must be an atom-style or atomfile-style variable. The variable is evaluated and atoms whose per-atom values are 0.0, are removed from the dynamic group.

If the *property* keyword is used, the name refers to a custom integer or floating point per-atom vector defined via the *fix property/atom* command. This means the values in the vector can be read as part of a data file with the *read\_data* command or specified with the *set* command. Or accessed and changed via the *library interface to LAMMPS*, or by styles you add to LAMMPS (pair, fix, compute, etc) which access the custom vector and modify its values. Which means the values can be modified between or during simulations. Atoms whose values in the custom vector are zero are removed from the dynamic group. Note that the name of the custom per-atom vector is specified just as *name*, not as *i\_name* or *d\_name* as it is for other commands that use different kinds of custom atom vectors or arrays as arguments.

The assignment of atoms to a dynamic group is done at the beginning of each run and on every timestep that is a multiple of  $N$ , which is the argument for the *every* keyword ( $N = 1$  is the default). For an energy minimization, via the *minimize* command, an assignment is made at the beginning of the minimization, but not during the iterations of the minimizer.

The point in the timestep at which atoms are assigned to a dynamic group is after interatomic forces have been computed, but before any fixes which alter forces or otherwise update the system have been invoked. This means that atom positions have been updated, neighbor lists and ghost atoms are current, and both intermolecular and intramolecular forces have been calculated based on the new coordinates. Thus the region criterion, if applied, should be accurate. Also, any computes invoked by an atom-style variable should use updated information for that timestep (e.g., potential energy/atom or coordination number/atom). Similarly, fixes or computes which are invoked after that point in the timestep, should operate on the new group of atoms.

---

**Note:** If the *region* keyword is used to determine what atoms are in the dynamic group, atoms can move outside of the simulation box between reneighboring events. Thus if you want to include all atoms on the left side of the simulation box, you probably want to set the left boundary of the region to be outside the simulation box by some reasonable amount (e.g., up to the cutoff of the potential), else they may be excluded from the dynamic region.

---

Here is an example of using a dynamic group to shrink the set of atoms being integrated by using a spherical region with a variable radius (shrinking from 18 to 5 over the course of the run). This could be used to model a quench of the system, freezing atoms outside the shrinking sphere, then converting the remaining atoms to a static group and running further.

```
variable      nsteps equal 5000
variable      rad equal 18-(step/v_nsteps)*(18-5)
region        ss sphere 20 20 0 v_rad
group         mobile dynamic all region ss
fix           1 mobile nve
run           ${nsteps}
group         mobile static
run           ${nsteps}
```

---

**Note:** All fixes and computes take a group ID as an argument, but they do not all allow for use of a dynamic group. If you get an error message that this is not allowed, but feel that it should be for the fix or compute in question, then please post your reasoning to the [LAMMPS forum at MatSci](#) and we can look into changing it. The same applies if you come across inconsistent behavior when dynamic groups are allowed.

---

The *static* style removes the setting for a dynamic group, converting it to a static group (the default). The atoms in the static group are those currently in the dynamic group.

---

#### 16.54.4 Restrictions

There can be no more than 32 groups defined at one time, including “all”.

The parent group of a dynamic group cannot itself be a dynamic group.

#### 16.54.5 Related commands

*dump, fix, region, velocity*

#### 16.54.6 Default

All atoms belong to the “all” group.

### 16.55 group2ndx command

### 16.56 ndx2group command

#### 16.56.1 Syntax

```
group2ndx file group-ID ...
ndx2group file group-ID ...
```

- file = name of index file to write out or read in
- zero or more group IDs may be appended

## 16.56.2 Examples

```
group2ndx allindex.ndx
group2ndx someindex.ndx upper lower mobile
ndx2group someindex.ndx
ndx2group someindex.ndx mobile
```

## 16.56.3 Description

Write or read a Gromacs style index file in text format that associates atom IDs with the corresponding group definitions. This index file can be used in combination with Gromacs analysis tools or to import group definitions into the *fix colvars* input file. It can also be used to save and restore group definitions for static groups.

The *group2ndx* command will write group definitions to an index file. Without specifying any group IDs, all groups will be written to the index file. When specifying group IDs, only those groups will be written to the index file. In order to follow the Gromacs conventions, the group *all* will be renamed to *System* in the index file.

The *ndx2group* command will create or update group definitions from those stored in an index file. Without specifying any group IDs, all groups except *System* will be read from the index file and the corresponding groups recreated. If a group of the same name already exists, it will be completely reset. When specifying group IDs, those groups, if present, will be read from the index file and restored.

## 16.56.4 Restrictions

This command requires that atoms have atom IDs, since this is the information that is written to the index file.

These commands are part of the COLVARS package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 16.56.5 Related commands

*group, dump, fix colvars*

## 16.56.6 Default

none

## 16.57 hyper command

### 16.57.1 Syntax

```
hyper N Nevent fix-ID compute-ID keyword values ...
```

- N = # of timesteps to run
- Nevent = check for events every this many steps
- fix-ID = ID of a fix that applies a global or local bias potential, can be NULL

- compute-ID = ID of a compute that identifies when an event has occurred
- zero or more keyword/value pairs may be appended
- keyword = *min* or *dump* or *rebond*

```

min values = etol ftol maxiter maxeval
  etol = stopping tolerance for energy, used in quenching
  ftol = stopping tolerance for force, used in quenching
  maxiter = max iterations of minimize, used in quenching
  maxeval = max number of force/energy evaluations, used in quenching
dump value = dump-ID
  dump-ID = ID of dump to trigger whenever an event takes place
rebond value = Nrebond
  Nrebond = frequency at which to reset bonds, even if no event has occurred

```

## 16.57.2 Examples

```

compute event all event/displace 1.0
fix HG mobile hyper/global 3.0 0.3 0.4 800.0
hyper 5000 100 HG event min 1.0e-6 1.0e-6 100 100 dump 1 dump 5

```

## 16.57.3 Description

Run a bond-boost hyperdynamics (HD) simulation where time is accelerated by application of a bias potential to one or more pairs of nearby atoms in the system. This command can be used to run both global and local hyperdynamics. In global HD a single bond within the system is biased on each timestep. In local HD multiple bonds (separated by a sufficient distance) can be biased simultaneously at each timestep. In the bond-boost hyperdynamics context, a “bond” is not a covalent bond between a pair of atoms in a molecule. Rather it is simply a pair of nearby atoms as discussed below.

Both global and local HD are described in ([Voter2013](#)) by Art Voter and collaborators. Similar to parallel replica dynamics (PRD), global and local HD are methods for performing accelerated dynamics that are suitable for infrequent-event systems that obey first-order kinetics. A good overview of accelerated dynamics methods (AMD) for such systems is given in ([Voter2002](#)) from the same group. To quote from the review paper: “The dynamical evolution is characterized by vibrational excursions within a potential basin, punctuated by occasional transitions between basins. The transition probability is characterized by  $p(t) = k \cdot \exp(-kt)$  where  $k$  is the rate constant.”

Both HD and PRD produce a time-accurate trajectory that effectively extends the timescale over which a system can be simulated, but they do it differently. HD uses a single replica of the system and accelerates time by biasing the interaction potential in a manner such that each timestep is effectively longer. PRD creates  $N_r$  replicas of the system and runs dynamics on each independently with a normal unbiased potential until an event occurs in one of the replicas. The time between events is reduced by a factor of  $N_r$  replicas. For both methods, per CPU second, more physical time elapses and more events occur. See the [prd](#) page for more info about PRD.

An HD run has several stages, which are repeated each time an event occurs, as explained below. The logic for an HD run is as follows:

```

quench
create initial list of bonds

while (time remains):
  run dynamics for Nevent steps
  quench

```

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```

check for an event
if event occurred: reset list of bonds
restore pre-quench state

```

The list of bonds is the list of atom pairs of atoms that are within a short cutoff distance of each other after the system energy is minimized (quenched). This list is created and reset by a *fix hyper/global* or *fix hyper/local* command specified as *fix-ID*. At every dynamics timestep, the same *fix* selects one of more bonds to apply a bias potential to.

---

**Note:** The style of *fix* associated with the specified *fix-ID* determines whether you are running the global versus local hyperdynamics algorithm.

---

Dynamics (with the bias potential) is run continuously, stopping every *Nevent* steps to check if a transition event has occurred. The specified *N* for total steps must be a multiple of *Nevent*. check is performed by quenching the system and comparing the resulting atom coordinates to the coordinates from the previous basin.

A quench is an energy minimization and is performed by whichever algorithm has been defined by the *min\_style* command. Minimization parameters may be set via the *min\_modify* command and by the *min* keyword of the hyper command. The latter are the settings that would be used with the *minimize* command. Note that typically, you do not need to perform a highly-converged minimization to detect a transition event, though you may need to in order to prevent a set of atoms in the system from relaxing to a saddle point.

The event check is performed by a compute with the specified *compute-ID*. Currently there is only one compute that works with the hyper command, which is the *compute event/displace* command. Other event-checking computes may be added. *Compute event/displace* checks whether any atom in the compute group has moved further than a specified threshold distance. If so, an event has occurred.

If this happens, the list of bonds is reset, since some bond pairs are likely now too far apart, and new pairs are likely close enough to be considered a bond. The pre-quenched state of the system (coordinates and velocities) is restored, and dynamics continue.

At the end of the hyper run, a variety of statistics are output to the screen and logfile. These include info relevant to both global and local hyperdynamics, such as the number of events and the elapsed hyper time (accelerated time), And it includes info specific to one or the other, depending on which style of *fix* was specified by *fix-ID*.

---

The optional keywords operate as follows.

As explained above, the *min* keyword can be used to specify parameters for the quench. Their meaning is the same as for the *minimize* command

The *dump* keyword can be used to trigger a specific dump command with the specified *dump-ID* to output a snapshot each time an event is detected. It can be specified multiple times with different *dump-ID* values, as in the example above. These snapshots will be for the quenched state of the system on a timestep that is a multiple of *Nevent*, i.e. a timestep after the event has occurred. Note that any dump command in the input script will also output snapshots at whatever timestep interval it defines via its *N* argument; see the *dump* command for details. This means if you only want a particular dump to output snapshots when events are detected, you should specify its *N* as a value larger than the length of the hyperdynamics run.

As in the code logic above, the bond list is normally only reset when an event occurs. The *rebond* keyword will force a reset of the bond list every *Nrebond* steps, even if an event has not occurred. *Nrebond* must be a multiple of *Nevent*. This can be useful to check if more frequent resets alter event statistics, perhaps because the parameters chosen for defining what is a bond and what is an event are producing bad dynamics in the presence of the bias potential.

---

## 16.57.4 Restrictions

This command can only be used if LAMMPS was built with the REPLICA package. See the *Build package* doc page for more info.

## 16.57.5 Related commands

*fix hyper/global, fix hyper/local, compute event/displace, prd*

## 16.57.6 Default

The option defaults are min = 0.1 0.1 40 50 and time = steps.

---

(Voter2013) S. Y. Kim, D. Perez, A. F. Voter, J Chem Phys, 139, 144110 (2013).

(Voter2002) Voter, Montalenti, Germann, Annual Review of Materials Research 32, 321 (2002).

## 16.58 if command

### 16.58.1 Syntax

```
if boolean then t1 t2 ... elif boolean f1 f2 ... elif boolean f1 f2 ... else e1 e2 ...
```

- boolean = a Boolean expression evaluated as TRUE or FALSE (see below)
- then = required word
- t1,t2,...,tN = one or more LAMMPS commands to execute if condition is met, each enclosed in quotes
- elif = optional word, can appear multiple times
- f1,f2,...,fN = one or more LAMMPS commands to execute if elif condition is met, each enclosed in quotes (optional arguments)
- else = optional argument
- e1,e2,...,eN = one or more LAMMPS commands to execute if no condition is met, each enclosed in quotes (optional arguments)

### 16.58.2 Examples

```
if "${steps} > 1000" then quit
if "${myString} == a10" then quit
if "$x <= $y" then "print 'X is smaller = $x'" else "print 'Y is smaller = $y'"
if "(${eng} > 0.0) || ($n < 1000)" then &
  "timestep 0.005" &
elif $n<10000 &
  "timestep 0.01" &
else &
  "timestep 0.02" &
```

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```
"print 'Max step reached'"
if "${eng} > ${eng_previous}" then "jump file1" else "jump file2"
```

### 16.58.3 Description

This command provides an if-then-else capability within an input script. A Boolean expression is evaluated and the result is TRUE or FALSE. Note that as in the examples above, the expression can contain variables, as defined by the *variable* command, which will be evaluated as part of the expression. Thus a user-defined formula that reflects the current state of the simulation can be used to issue one or more new commands.

If the result of the Boolean expression is TRUE, then one or more commands (t1, t2, ..., tN) are executed. If it is FALSE, then Boolean expressions associated with successive elif keywords are evaluated until one is found to be true, in which case its commands (f1, f2, ..., fN) are executed. If no Boolean expression is TRUE, then the commands associated with the else keyword, namely (e1, e2, ..., eN), are executed. The elif and else keywords and their associated commands are optional. If they are not specified and the initial Boolean expression is FALSE, then no commands are executed.

The syntax for Boolean expressions is described below.

Each command (t1, f1, e1, etc.) can be any valid LAMMPS input script command. If the command is more than one word, it must be enclosed in quotes, so it will be treated as a single argument, as in the examples above.

---

**Note:** If a command itself requires a quoted argument (e.g., a *print* command), then double and single quotes can be used and nested in the usual manner, as in the examples above and below. The *Commands parse* page has more details on using quotes in arguments. Only one level of nesting is allowed, but that should be sufficient for most use cases.

---

Note that by using the line continuation character “&”, the if command can be spread across many lines, though it is still a single command:

```
if "$a < $b" then &
  "print 'Minimum value = $a'" &
  "run 1000" &
else &
  'print "Minimum value = $b"' &
  "minimize 0.001 0.001 1000 10000"
```

Note that if one of the commands to execute is *quit*, as in the first example above, then executing the command will cause LAMMPS to halt.

Note that by jumping to a label in the same input script, the if command can be used to break out of a loop. See the *variable delete* command for info on how to delete the associated loop variable, so that it can be re-used later in the input script.

Here is an example of a loop which checks every 1000 steps if the system temperature has reached a certain value, and if so, breaks out of the loop to finish the run. Note that any variable could be checked, so long as it is current on the timestep when the run completes. As explained on the *variable* doc page, this can be insured by including the variable in thermodynamic output.

```
variable myTemp equal temp
label loop
variable a loop 1000
run 1000
if "${myTemp} < 300.0" then "jump SELF break"
```

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```

next a
jump SELF loop
label break
print "ALL DONE"

```

Here is an example of a double loop which uses the `if` and `jump` commands to break out of the inner loop when a condition is met, then continues iterating through the outer loop.

```

label      loopa
variable  a loop 5
  label   loopb
  variable b loop 5
  print   "A,B = $a,$b"
  run     10000
  if      "$b > 2" then "jump SELF break"
next      b
jump     in.script loopb
label    break
variable b delete
next     a
jump    SELF loopa

```

The Boolean expressions for the `if` and `elif` keywords have a C-like syntax. Note that each expression is a single argument within the `if` command. Thus if you want to include spaces in the expression for clarity, you must enclose the entire expression in quotes.

An expression is built out of numbers (which start with a digit or period or minus sign) or strings (which start with a letter and can contain alphanumeric characters, underscores, or forward slashes):

```

0.2, 100, 1.0e20, -15.4, ...
InP, myString, a123, ab_23_cd, lj/cut, ...

```

and Boolean operators:

```
A == B, A != B, A < B, A <= B, A > B, A >= B, A && B, A || B, A |^ B, !A
```

Each A and B is a number or string or a variable reference like `$a` or `${abc}`, or A or B can be another Boolean expression.

Note that all variables used will be substituted for before the Boolean expression is evaluated. A variable can produce a number, like an *equal-style variable*, or it can produce a string, like an *index-style variable*.

The Boolean operators `==` and `!=` can operate on a pair of strings or numbers. They cannot compare a number to a string. All the other Boolean operations can only operate on numbers.

Expressions are evaluated left to right and have the usual C-style precedence: the unary logical NOT operator `!` has the highest precedence, the 4 relational operators `<`, `<=`, `>`, and `>=` are next; the two remaining relational operators `==` and `!=` are next; then the logical AND operator `&&`; and finally the logical OR operator `||` and logical XOR (exclusive or) operator `|^` have the lowest precedence. Parenthesis can be used to group one or more portions of an expression and/or enforce a different order of evaluation than what would occur with the default precedence.

When the six relational operators (first six in list above) compare two numbers, they return either a 1.0 or 0.0 depending on whether the relationship between A and B is TRUE or FALSE.

When the three logical operators (last three in list above) compare two numbers, they also return either a 1.0 or 0.0 depending on whether the relationship between A and B is TRUE or FALSE (or just A). The logical AND operator will

return 1.0 if both its arguments are non-zero, else it returns 0.0. The logical OR operator will return 1.0 if either of its arguments is non-zero, else it returns 0.0. The logical XOR operator will return 1.0 if one of its arguments is zero and the other non-zero, else it returns 0.0. The logical NOT operator returns 1.0 if its argument is 0.0, else it returns 0.0. The 3 logical operators can only be used to operate on numbers, not on strings.

The overall Boolean expression produces a TRUE result if the numeric result is non-zero. If the result is zero, the expression result is FALSE.

---

**Note:** If the Boolean expression is a single numeric value with no Boolean operators, it will be FALSE if the value = 0.0, otherwise TRUE. If the Boolean expression is a single string, an error message will be issued.

---

## 16.58.4 Restrictions

none

## 16.58.5 Related commands

*variable, print*

## 16.58.6 Default

none

# 16.59 improper\_coeff command

## 16.59.1 Syntax

```
improper_coeff N args
```

- N = numeric improper type (see asterisk form below), or type label
- args = coefficients for one or more improper types

## 16.59.2 Examples

```
improper_coeff 1 300.0 0.0
improper_coeff * 80.2 -1 2
improper_coeff *4 80.2 -1 2

labelmap improper 1 benzene
improper_coeff benzene 300.0 0.0
```

### 16.59.3 Description

Specify the improper force field coefficients for one or more improper types. The number and meaning of the coefficients depends on the improper style. Improper coefficients can also be set in the data file read by the *read\_data* command or in a restart file.

$N$  can be specified in one of two ways. An explicit numeric value can be used, as in the first example above. Or  $N$  can be a type label, which is an alphanumeric string defined by the *labelmap* command or in a section of a data file read by the *read\_data* command.

For numeric values only, a wild-card asterisk can be used to set the coefficients for multiple improper types. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If  $N$  = the number of improper types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to n (inclusive). A trailing asterisk means all types from n to  $N$  (inclusive). A middle asterisk means all types from m to n (inclusive).

Note that using an *improper\_coeff* command can override a previous setting for the same improper type. For example, these commands set the coeffs for all improper types, then overwrite the coeffs for just improper type 2:

```
improper_coeff * 300.0 0.0
improper_coeff 2 50.0 0.0
```

A line in a data file that specifies improper coefficients uses the exact same format as the arguments of the *improper\_coeff* command in an input script, except that wild-card asterisks should not be used since coefficients for all  $N$  types must be listed in the file. For example, under the “Improper Coeffs” section of a data file, the line that corresponds to the first example above would be listed as

```
1 300.0 0.0
```

The *improper\_style class2* is an exception to this rule, in that an additional argument is used in the input script to allow specification of the cross-term coefficients. See its doc page for details.

---

The list of all improper styles defined in LAMMPS is given on the *improper\_style* doc page. They are also listed in more compact form on the *Commands improper* doc page.

On either of those pages, click on the style to display the formula it computes and its coefficients as specified by the associated *improper\_coeff* command.

---

### 16.59.4 Restrictions

This command must come after the simulation box is defined by a *read\_data*, *read\_restart*, or *create\_box* command. An improper style must be defined before any improper coefficients are set, either in the input script or in a data file.

### 16.59.5 Related commands

*improper\_style*

## 16.59.6 Default

none

## 16.60 improper\_style command

### 16.60.1 Syntax

```
improper_style style
```

- style = *none* or *hybrid* or *class2* or *cvff* or *harmonic*

### 16.60.2 Examples

```
improper_style harmonic
improper_style cvff
improper_style hybrid cvff harmonic
```

### 16.60.3 Description

Set the formula(s) LAMMPS uses to compute improper interactions between quadruplets of atoms, which remain in force for the duration of the simulation. The list of improper quadruplets is read in by a *read\_data* or *read\_restart* command from a data or restart file. Note that the ordering of the 4 atoms in an improper quadruplet determines the definition of the improper angle used in the formula for each style. See the doc pages of individual styles for details.

Hybrid models where impropers are computed using different improper potentials can be setup using the *hybrid* improper style.

The coefficients associated with an improper style can be specified in a data or restart file or via the *improper\_coeff* command.

All improper potentials store their coefficient data in binary restart files which means *improper\_style* and *improper\_coeff* commands do not need to be re-specified in an input script that restarts a simulation. See the *read\_restart* command for details on how to do this. The one exception is that *improper\_style hybrid* only stores the list of sub-styles in the restart file; improper coefficients need to be re-specified.

---

**Note:** When both an improper and pair style is defined, the *special\_bonds* command often needs to be used to turn off (or weight) the pairwise interaction that would otherwise exist between a group of 4 bonded atoms.

---

Here is an alphabetic list of improper styles defined in LAMMPS. Click on the style to display the formula it computes and coefficients specified by the associated *improper\_coeff* command.

Click on the style to display the formula it computes, any additional arguments specified in the *improper\_style* command, and coefficients specified by the associated *improper\_coeff* command.

There are also additional accelerated pair styles included in the LAMMPS distribution for faster performance on CPUs, GPUs, and KNLs. The individual style names on the *Commands improper* page are followed by one or more of (g,i,k,o,t) to indicate which accelerated styles exist.

- *none* - turn off improper interactions

- *zero* - topology but no interactions
- *hybrid* - define multiple styles of improper interactions
- *amoeba* - AMOEBA out-of-plane improper
- *class2* - COMPASS (class 2) improper
- *cosq* - improper with a cosine squared term
- *cvff* - CVFF improper
- *distance* - improper based on distance between atom planes
- *distharm* - improper that is harmonic in the out-of-plane distance
- *fourier* - improper with multiple cosine terms
- *harmonic* - harmonic improper
- *inversion/harmonic* - harmonic improper with Wilson-Decius out-of-plane definition
- *ring* - improper which prevents planar conformations
- *umbrella* - DREIDING improper

*sqdistharm* - improper that is harmonic in the square of the out-of-plane distance

---

## 16.60.4 Restrictions

Improper styles can only be set for atom\_style choices that allow improper to be defined.

Most improper styles are part of the MOLECULE package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. The doc pages for individual improper potentials tell if it is part of a package.

## 16.60.5 Related commands

*improper\_coeff*

## 16.60.6 Default

```
improper_style none
```

## 16.61 include command

### 16.61.1 Syntax

```
include file
```

- file = filename of new input script to switch to



## 16.61.2 Examples

```
include newfile
include in.run2
```

## 16.61.3 Description

This command opens a new input script file and begins reading LAMMPS commands from that file. When the new file is finished, the original file is returned to. Include files can be nested as deeply as desired. If input script A includes script B, and B includes A, then LAMMPS could run for a long time.

If the filename is a variable (see the *variable* command), different processor partitions can run different input scripts.

## 16.61.4 Restrictions

none

## 16.61.5 Related commands

*variable, jump*

## 16.61.6 Default

none

# 16.62 info command

## 16.62.1 Syntax

```
info args
```

- args = one or more of the following keywords: *out, all, system, memory, communication, computes, dumps, fixes, groups, regions, variables, coeffs, styles, time, accelerator, or configuration*
- out values = *screen, log, append filename, overwrite filename*
- styles values = *all, angle, atom, bond, compute, command, dump, dihedral, fix, improper, integrate, kspace, minimize, pair, region*

## 16.62.2 Examples

```
info system
info groups computes variables
info all out log
info all out append info.txt
info styles all
info styles atom
```

## 16.62.3 Description

Print out information about the current internal state of the running LAMMPS process. This can be helpful when debugging or validating complex input scripts. Several output categories are available and one or more output category may be requested.

The *out* flag controls where the output is sent. It can only be sent to one target. By default this is the screen, if it is active. The *log* argument selects the log file instead. With the *append* and *overwrite* option, followed by a filename, the output is written to that file, which is either appended to or overwritten, respectively.

The *all* flag activates printing all categories listed below.

The *configuration* category prints some information about the LAMMPS version as well as architecture and OS it is run on.

The *memory* category prints some information about the current memory allocation of MPI rank 0 (this is the amount of dynamically allocated memory reported by LAMMPS classes). Where supported, also some OS specific information about the size of the reserved memory pool size (this is where `malloc()` and the new operator request memory from) and the maximum resident set size is reported (this is the maximum amount of physical memory occupied so far).

The *system* category prints a general system overview listing. This includes the unit style, atom style, number of atoms, bonds, angles, dihedrals, and impropers and the number of the respective types, box dimensions and properties, force computing styles and more.

The *communication* category prints a variety of information about communication and parallelization: the MPI library version level, the number of MPI ranks and OpenMP threads, the communication style and layout, the processor grid dimensions, ghost atom communication mode, cutoff, and related settings.

The *computes* category prints a list of all currently defined computes, their IDs and styles and groups they operate on.

The *dumps* category prints a list of all currently active dumps, their IDs, styles, filenames, groups, and dump frequencies.

The *fixes* category prints a list of all currently defined fixes, their IDs and styles and groups they operate on.

The *groups* category prints a list of all currently defined groups.

The *regions* category prints a list of all currently defined regions, their IDs and styles and whether “inside” or “outside” atoms are selected.

The *variables* category prints a list of all currently defined variables, their names, styles, definition and last computed value, if available.

The *coeffs* category prints a list for each defined force style (pair, bond, angle, dihedral, improper) indicating which of the corresponding coefficients have been set. This can be very helpful to debug error messages like “All pair coeffs are not set”.

The *accelerator* category prints out information about compile time settings of included accelerator support for the GPU, KOKKOS, INTEL, and OPENMP packages.

The *styles* category prints the list of styles available in the current LAMMPS binary. It supports one of the following options to control which category of styles is printed out:

- all
- angle
- atom
- bond
- compute
- command
- dump
- dihedral
- fix
- improper
- integrate
- kspace
- minimize
- pair
- region

The *time* category prints the accumulated CPU and wall time for the process that writes output (usually MPI rank 0).

## 16.62.4 Restrictions

none

## 16.62.5 Related commands

*print*

## 16.62.6 Default

The *out* option has the default *screen*.

The *styles* option has the default *all*.

## 16.63 jump command

### 16.63.1 Syntax

```
jump file label
```

- file = filename of new input script to switch to
- label = optional label within file to jump to

## 16.63.2 Examples

```
jump newfile
jump in.run2 runloop
jump SELF runloop
```

## 16.63.3 Description

This command closes the current input script file, opens the file with the specified name, and begins reading LAMMPS commands from that file. Unlike the *include* command, the original file is not returned to, although by using multiple jump commands it is possible to chain from file to file or back to the original file.

If the word “SELF” is used for the filename, then the current input script is re-opened and read again.

---

**Note:** The SELF option is not guaranteed to work when the current input script is being read through stdin (standard input), e.g.

---

```
lmp_g++ < in.script
```

since the SELF option invokes the C-library `rewind()` call, which may not be supported for stdin on some systems or by some MPI implementations. This can be worked around by using the *-in command-line switch*, e.g.

```
lmp_g++ -in in.script
```

or by using the *-var command-line switch* to pass the script name as a variable to the input script. In the latter case, a *variable* called “fname” could be used in place of SELF, e.g.

```
lmp_g++ -var fname in.script < in.script
```

The second argument to the jump command is optional. If specified, it is treated as a label and the new file is scanned (without executing commands) until the label is found, and commands are executed from that point forward. This can be used to loop over a portion of the input script, as in this example. These commands perform 10 runs, each of 10000 steps, and create 10 dump files named file.1, file.2, etc. The *next* command is used to exit the loop after 10 iterations. When the “a” variable has been incremented for the tenth time, it will cause the next jump command to be skipped.

```
variable a loop 10
label loop
dump 1 all atom 100 file.$a
run 10000
undump 1
next a
jump in.lj loop
```

If the jump *file* argument is a variable, the jump command can be used to cause different processor partitions to run different input scripts. In this example, LAMMPS is run on 40 processors, with 4 partitions of 10 procs each. An in.file containing the example variable and jump command will cause each partition to run a different simulation.

```
mpirun -np 40 lmp_ibm -partition 4x10 -in in.file

variable f world script.1 script.2 script.3 script.4
jump $f
```

Here is an example of a loop which checks every 1000 steps if the system temperature has reached a certain value, and if so, breaks out of the loop to finish the run. Note that any variable could be checked, so long as it is current on the timestep when the run completes. As explained on the *variable* doc page, this can be insured by including the variable in thermodynamic output.

```
variable myTemp equal temp
label loop
variable a loop 1000
run 1000
if "${myTemp} < 300.0" then "jump SELF break"
next a
jump SELF loop
label break
print "ALL DONE"
```

Here is an example of a double loop which uses the *if* and *jump* commands to break out of the inner loop when a condition is met, then continues iterating through the outer loop.

```
label loopa
variable a loop 5
  label loopb
  variable b loop 5
  print "A,B = $a,$b"
  run 10000
  if "$b > 2" then "jump SELF break"
next b
jump in.script loopb
label break
variable b delete
next a
jump SELF loopa
```

### 16.63.4 Restrictions

If you jump to a file and it does not contain the specified label, LAMMPS will come to the end of the file and exit.

### 16.63.5 Related commands

*variable, include, label, next*

### 16.63.6 Default

none

## 16.64 kim command

### 16.64.1 Syntax

```
kim sub-command args
```

- sub-command = *init* or *interactions* or *query* or *param* or *property*
- args = arguments used by a particular sub-command

### 16.64.2 Examples

```
kim init args
kim interactions args
kim query args
kim param args
kim property args
```

### 16.64.3 Description

The *kim command* includes a set of sub-commands that allow LAMMPS users to use interatomic models (IM) (potentials and force fields) and their predictions for various physical properties archived in the [Open Knowledgebase of Interatomic Models \(OpenKIM\)](#) repository.

Using OpenKIM provides LAMMPS users with immediate access to a large number of verified IMs and their predictions. OpenKIM IMs have multiple benefits including [reliability](#), [reproducibility](#) and [convenience](#).

There are two types of IMs archived in OpenKIM:

1. The first type is called a *KIM Portable Model (PM)*. A KIM PM is an independent computer implementation of an IM written in one of the languages supported by KIM (C, C++, Fortran) that conforms to the KIM Application Programming Interface ([KIM API](#)) Portable Model Interface (PMI) standard. A KIM PM will work seamlessly with any simulation code that supports the KIM API/PMI standard (including LAMMPS; see [complete list of supported codes](#)).
2. The second type is called a *KIM Simulator Model (SM)*. A KIM SM is an IM that is implemented natively within a simulation code (*simulator*) that supports the KIM API Simulator Model Interface (SMI); in this case LAMMPS. A separate SM package is archived in OpenKIM for each parameterization of the IM, which includes all of the necessary parameter files, LAMMPS commands, and metadata (supported species, units, etc.) needed to run the IM in LAMMPS.

With these two IM types, OpenKIM can archive and test almost all IMs that can be used by LAMMPS. (It is easy to contribute new IMs to OpenKIM, see the [upload instructions](#).)

OpenKIM IMs are uniquely identified by a [KIM ID](#). The extended KIM ID consists of a human-readable prefix identifying the type of IM, authors, publication year, and supported species, separated by two underscores from the KIM ID itself, which begins with an IM code (*MO* for a KIM Portable Model, and *SM* for a KIM Simulator Model) followed by a unique 12-digit code and a 3-digit version identifier. By convention SM prefixes begin with *Sim\_* to readily identify them.

```
SW_StillingerWeber_1985_Si__MO_405512056662_005
Sim_LAMMPS_ReaxFF_StrachanVanDuinChakraborty_2003_CHNO__SM_107643900657_001
```

Each OpenKIM IM has a dedicated “Model Page” on [OpenKIM](#) providing all the information on the IM including a title, description, authorship and citation information, test and verification check results, visualizations of results, a wiki with documentation and user comments, and access to raw files, and other information. The URL for the Model Page is constructed from the [extended KIM ID](#) of the IM:

```
https://openkim.org/id/extended_KIM_ID
```

For example, for the Stillinger-Weber potential listed above the Model Page is located at:

```
https://openkim.org/id/SW_StillingerWeber_1985_Si__MO_405512056662_005
```

See the [current list of KIM PMs and SMs archived in OpenKIM](#). This list is sorted by species and can be filtered to display only IMs for certain species combinations.

See [Obtaining KIM Models](#) to learn how to install a pre-built binary of the OpenKIM Repository of Models.

---

**Note:** It is also possible to locally install IMs not archived in OpenKIM, in which case their names do not have to conform to the KIM ID format.

---

#### 16.64.4 Using OpenKIM IMs with LAMMPS (*kim init*, *kim interactions*)

Two sub-commands are employed when using OpenKIM IMs in LAMMPS, one to select the IM and perform necessary initialization (*kim init*), and the second to set up the IM for use by executing any necessary LAMMPS commands (*kim interactions*). Both are required.

##### Syntax

```
kim init model user_units unitarg
kim interactions typeargs
```

- model = name of the KIM interatomic model (the KIM ID for models archived in OpenKIM)
- user\_units = the LAMMPS *units* style assumed in the LAMMPS input script
- unitarg = *unit\_conversion\_mode* (optional)
- typeargs = atom type to species mapping (one entry per atom type) or *fixed\_types* for models with a preset fixed mapping

##### Examples

```
kim init SW_StillingerWeber_1985_Si__MO_405512056662_005 metal
kim interactions Si

kim init Sim_LAMMPS_ReaxFF_StrachanVanDuinChakraborty_2003_CHNO__SM_107643900657_001 real
kim init Sim_LAMMPS_ReaxFF_StrachanVanDuinChakraborty_2003_CHNO__SM_107643900657_001_
↪metal unit_conversion_mode
kim interactions C H O

kim init Sim_LAMMPS_IFF_PCFF_HeinzMishraLinEmami_2015Ver1v5_
↪FccmetalsMineralsSolventsPolymers__SM_039297821658_000 real
kim interactions fixed_types
```

See the *examples/kim* directory for example input scripts that use KIM PMs and KIM SMs.

### OpenKIM IM Initialization (*kim init*)

The *kim* command followed by *init* sub-command must be issued **before** the simulation box is created (normally at the top of the file). This command sets the OpenKIM IM that will be used and may issue additional commands changing LAMMPS default settings that are required for using the selected IM (such as *units* or *atom\_style*). If needed, those settings can be overridden, however, typically a script containing a *kim init* command would not include *units* and *atom\_style* commands.

The required arguments of *kim init* are the *model* name of the IM to be used in the simulation (for an IM archived in OpenKIM this is its *extended KIM ID*, and the *user\_units*, which are the LAMMPS *units style* used in the input script. (Any dimensioned numerical values in the input script and values read in from files are expected to be in the *user\_units* system.)

The selected IM can be either a *KIM PM* or a *KIM SM*. For a KIM SM, the *kim init* command verifies that the SM is designed to work with LAMMPS (and not another simulation code). In addition, the LAMMPS version used for defining the SM and the LAMMPS version being currently run are printed to help diagnose any incompatible changes to input script or command syntax between the two LAMMPS versions.

Based on the selected model *kim init* may modify the *atom\_style*. Some SMs have requirements for this setting. If this is the case, then *atom\_style* will be set to the required style. Otherwise, the value is left unchanged (which in the absence of an *atom\_style* command in the input script is the *default atom\_style value*).

Regarding units, the *kim init* behaves in different ways depending on whether or not *unit conversion mode* is activated as indicated by the optional *unitarg* argument. If unit conversion mode is **not** active, then *user\_units* must either match the required units of the IM or the IM must be able to adjust its units to match. (The latter is only possible with some KIM PMs; SMs can never adjust their units.) If a match is possible, the LAMMPS *units* command is called to set the units to *user\_units*. If the match fails, the simulation is terminated with an error. The *kim init* command also sets the default value for the *skin* (extra distance beyond force cutoff) as 2.0 Angstroms and sets the default value for the *timestep* size as 1.0 femtosecond.

Here is an example of a LAMMPS script to compute the cohesive energy of a face-centered cubic (fcc) lattice for the MEAM potential by Pascuet and Fernandez (2015) for Al.

```
kim      init Sim_LAMMPS_MEAM_PascuetFernandez_2015_Al__SM_811588957187_000 metal
boundary  p p p
lattice  fcc 4.049
region   simbox block 0 1 0 1 0 1 units lattice
create_box 1 simbox
create_atoms 1 box
mass     1 26.981539
kim      interactions Al
run      0
variable Ec equal (pe/count(all))
print    "Cohesive Energy = ${Ec} eV"
```

The above script will end with an error in the *kim init* line if the IM is changed to another potential for Al that does not work with *metal* units. To address this, *kim init* offers the *unit\_conversion\_mode* as shown below.

If unit conversion mode *is* active, then *kim init* calls the LAMMPS *units* command to set the units to the IM's required or preferred units. Conversion factors between the IM's units and the *user\_units* are defined for all *physical quantities* (mass, distance, etc.). (Note that converting to or from the "l" unit style is not supported.) These factors are stored as *internal style variables* with the following standard names:



```

_u_mass
_u_distance
_u_time
_u_energy
_u_velocity
_u_force
_u_torque
_u_temperature
_u_pressure
_u_viscosity
_u_charge
_u_dipole
_u_efield
_u_density

```

If desired, the input script can be designed to work with these conversion factors so that the script will work without change with any OpenKIM IM. (This approach is used in the [OpenKIM Testing Framework](#).)

For example, the script given above for the cohesive energy of fcc Al can be rewritten to work with any IM regardless of units. The following script constructs an fcc lattice with a lattice parameter defined in meters, computes the total energy, and prints the cohesive energy in Joules regardless of the units of the IM.

```

kim          init Sim_LAMMPS_MEAM_PascuetFernandez_2015_Al__SM_811588957187_000 si unit_
↳conversion_mode
boundary     p p p
lattice      fcc $(4.049e-10*v__u_distance)
region       simbox block 0 1 0 1 0 1 units lattice
create_box   1 simbox
create_atoms 1 box
mass         1 $(4.480134e-26*v__u_mass)
kim          interactions Al
neighbor     $(2e-10*v__u_distance) bin
run          0
variable     Ec_in_J equal (pe/count(all))/v__u_energy
print       "Cohesive Energy = ${Ec_in_J} J"

```

Note the multiplication by `v__u_distance` and `v__u_mass` to convert from SI units (specified in the `kim init` command) to whatever units the IM uses (metal in this case), and the division by `v__u_energy` to convert from the IM's energy units to SI units (Joule). This script will work correctly for any IM for Al (KIM PM or SM) selected by the `kim init` command.

Care must be taken to apply unit conversion to dimensional variables read in from a file. For example, if a configuration of atoms is read in from a dump file using the `read_dump` command, the following can be done to convert the box and all atomic positions to the correct units:

```

change_box all x scale ${_u_distance} &
              y scale ${_u_distance} &
              z scale ${_u_distance} &
              xy final $(xy*v__u_distance) &
              xz final $(xz*v__u_distance) &
              yz final $(yz*v__u_distance) &
              remap

```

**Note:** Unit conversion will only work if the conversion factors are placed in all appropriate places in the input script.

It is up to the user to do this correctly.

---

### OpenKIM IM Execution (*kim interactions*)

The second and final step in using an OpenKIM IM is to execute the *kim interactions* command. This command must be preceded by a *kim init* command and a command that defines the number of atom types  $N$  (such as *create\_box*). The *kim interactions* command has one argument *typeargs*. This argument contains either a list of  $N$  chemical species, which defines a mapping between atom types in LAMMPS to the available species in the OpenKIM IM, or the keyword *fixed\_types* for models that have a preset fixed mapping (i.e. the mapping between LAMMPS atom types and chemical species is defined by the model and cannot be changed). In the latter case, the user must consult the model documentation to see how many atom types there are and how they map to the chemical species.

For example, consider an OpenKIM IM that supports Si and C species. If the LAMMPS simulation has four atom types, where the first three are Si, and the fourth is C, the following *kim interactions* command would be used:

```
kim interactions Si Si Si C
```

Alternatively, for a model with a fixed mapping the command would be:

```
kim interactions fixed_types
```

The *kim interactions* command performs all the necessary steps to set up the OpenKIM IM selected in the *kim init* command. The specific actions depend on whether the IM is a KIM PM or a KIM SM. For a KIM PM, a *pair\_style kim* command is executed followed by the appropriate *pair\_coeff* command. For example, for the Ercolessi and Adams (1994) KIM PM for Al set by the following commands:

```
kim init EAM_Dynamo_ErcolessiAdams_1994_Al__MO_123629422045_005 metal
...
... box specification lines skipped
...
kim interactions Al
```

the *kim interactions* command executes the following LAMMPS input commands:

```
pair_style kim EAM_Dynamo_ErcolessiAdams_1994_Al__MO_123629422045_005
pair_coeff * * Al
```

For a KIM SM, the generated input commands may be more complex and require that LAMMPS is built with the required packages included for the type of potential being used. The set of commands to be executed is defined in the SM specification file, which is part of the SM package. For example, for the Strachan et al. (2003) ReaxFF SM set by the following commands:

```
kim init Sim_LAMMPS_ReaxFF_StrachanVanDuinChakraborty_2003_CHNO__SM_107643900657_000 real
...
... box specification lines skipped
...
kim interactions C H N O
```

the *kim interactions* command executes the following LAMMPS input commands:

```
pair_style reaxff lmp_control safezone 2.0 mincap 100
pair_coeff * *ffield.reax.rdx C H N O
fix reaxqeq all qeq/reaxff 1 0.0 10.0 1.0e-6 param.qeq
```

---

**Note:** The files *lmp\_control*, *ffield.reax.rdx* and *param.qeq* are specific to the Strachan et al. (2003) ReaxFF parameterization and are archived as part of the SM package in OpenKIM.

---

**Note:** Parameters like cutoff radii and charge tolerances, which have an effect on IM predictions, are also included in the SM definition ensuring reproducibility.

---

**Note:** When using *kim init* and *kim interactions* to select and set up an OpenKIM IM, other LAMMPS commands for the same functions (such as *pair\_style*, *pair\_coeff*, *bond\_style*, *bond\_coeff*, fixes related to charge equilibration, etc.) should normally not appear in the input script.

---

**Note:** *kim interactions* must be called each time after the *change\_box* command to provide the correct settings (it should be called with the same *typeargs* as the first call.) The reason is that changing a periodic boundary to a non-periodic one, or in general, using the *change\_box* command after the interactions are set via *kim interactions* or *pair\_coeff* commands might affect some of the settings. For example, SM models containing Coulombic terms in the interactions require different settings if a periodic boundary changes to a non-periodic one. In other cases, the second call to *kim interactions* does not affect any other settings.

---

## 16.64.5 Using OpenKIM Web Queries in LAMMPS (*kim query*)

The *kim query* command performs a web query to retrieve the predictions of an IM set by *kim init* for material properties archived in OpenKIM.

### Syntax

```
kim query variable formatarg query_function queryargs
```

- variable(s) = single name or list of names of (string style) LAMMPS variable(s) where a query result or parameter get result is stored. Variables that do not exist will be created by the command
- formatarg = *list*, *split*, or *index* (optional):
  - list* = returns a single string with a list of space separated values (e.g. "1.0 2.0 3.0"), which is placed in a LAMMPS variable as defined by the *variable* argument. [default]
  - split* = returns the values separately in new variables with names based on the prefix specified in *variable* and a number appended to indicate which element in the list of values is in the variable
  - index* = returns a variable style index that can be incremented via the next command. This enables the construction of simple loops
- query\_function = name of the OpenKIM web API query function to be used
- queryargs = a series of *keyword=value* pairs that represent the web query; supported keywords depend on the query function

## Examples

```
kim query a0 get_lattice_constant_cubic crystal=[fcc] species=[Al] units=[angstrom]
kim query model index get_available_models species=[Al] potential_type=[eam]
```

The result of the query is stored in one or more *string style variables* as determined by the optional *formatarg* argument. For the “list” setting of *formatarg* (or if *formatarg* is not specified), the result is returned as a space-separated list of values in *variable*. The *formatarg* keyword “split” separates the result values into individual variables of the form *prefix\_I*, where *prefix* is set to the *kim query variable* argument and *I* ranges from 1 to the number of returned values. The number and order of the returned values is determined by the type of query performed. The *formatarg* keyword “index” returns a *variable style index* that can be incremented via the *next* command. This enables the construction of simple loops over the returned values by the type of query performed.

---

**Note:** *kim query* only supports queries that return a single result or an array of values. More complex queries that return a JSON structure are not currently supported. An attempt to use *kim query* in such cases will generate an error.

---

The second required argument *query\_function* is the name of the query function to be called (e.g. *get\_lattice\_constant\_cubic*). All following *arguments* are parameters handed over to the web query in the format *keyword=value*, where *value* is always an array of one or more comma-separated items in brackets. The list of supported keywords and the type and format of their values depend on the query function used. The current list of query functions is available on the OpenKIM webpage at <https://openkim.org/doc/usage/kim-query>.

---

**Note:** All query functions, except *get\_available\_models*, require the *model* keyword, which identifies the IM whose predictions are being queried. *kim query* automatically generates the *model* keyword based on the IM set in by *kim init*, and it can be overwritten if specified as an argument to the *kim query*. Where *kim init* is not specified, the *model* keyword must be provided as an argument to the *kim query*.

---

---

**Note:** Each *query\_function* is associated with a default method (implemented as a **KIM Test**) used to compute this property. In cases where there are multiple methods in OpenKIM for computing a property, a *method* keyword can be provided to select the method of choice. See the [query documentation](#) to see which methods are available for a given *query\_function*.

---

## *kim query* Usage Examples and Further Clarifications

The data obtained by *kim query* commands can be used as part of the setup or analysis phases of LAMMPS simulations. Some examples are given below.

### Define an equilibrium fcc crystal

```
kim      init EAM_Dynamo_ErcolessiAdams_1994_Al__MO_123629422045_005 metal
boundary p p p
kim      query a0 get_lattice_constant_cubic crystal=[fcc] species=[Al] units=[angstrom]
lattice  fcc ${a0}
...
```

```
units    metal
kim      query a0 get_lattice_constant_cubic crystal=[fcc] species=[Al] units=[angstrom]
↳model=[EAM_Dynamo_ErcolessiAdams_1994_Al__MO_123629422045_005]
```

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```
lattice fcc ${a0}
...
```

The *kim query* command retrieves from [OpenKIM](#) the equilibrium lattice constant predicted by the Ercolessi and Adams (1994) potential for the fcc structure and places it in variable *a0*. This variable is then used on the next line to set up the crystal. By using *kim query*, the user is saved the trouble and possible error of tracking this value down, or of having to perform an energy minimization to find the equilibrium lattice constant.

**Note:** In *unit\_conversion\_mode* the results obtained from a *kim query* would need to be converted to the appropriate units system. For example, in the above script, the lattice command would need to be changed to: “lattice fcc  $(v\_a0*v\_u\_distance)$ ”.

### Define an equilibrium hcp crystal

```
kim      init EAM_Dynamo_MendeleevAckland_2007v3_Zr__MO_004835508849_000 metal
boundary p p p
kim      query latconst split get_lattice_constant_hexagonal crystal=[hcp] species=[Zr]
↳units=[angstrom]
lattice  custom ${latconst_1} a1 0.5 -0.866025 0 a2 0.5 0.866025 0 a3 0 0 $(latconst_2/
↳latconst_1) &
        basis 0.333333 0.666666 0.25 basis 0.666666 0.333333 0.75
...
```

In this case the *kim query* returns two arguments (since the hexagonal close packed (hcp) structure has two independent lattice constants). The *formatarg* keyword “split” places the two values into the variables *latconst\_1* and *latconst\_2*. (These variables are created if they do not already exist.)

### Define a crystal at finite temperature accounting for thermal expansion

```
kim      init EAM_Dynamo_ErcolessiAdams_1994_Al__MO_123629422045_005 metal
boundary p p p
kim      query a0 get_lattice_constant_cubic crystal=[fcc] species=[Al] units=[angstrom]
kim      query alpha get_linear_thermal_expansion_coefficient_cubic crystal=[fcc]
↳species=[Al] units=[1/K] temperature=[293.15] temperature_units=[K]
variable DeltaT equal 300
lattice  fcc $(v_a0*v_alpha*v_DeltaT)
...
```

As in the previous example, the equilibrium lattice constant is obtained for the Ercolessi and Adams (1994) potential. However, in this case the crystal is scaled to the appropriate lattice constant at room temperature (293.15 K) by using the linear thermal expansion constant predicted by the potential.

**Note:** When passing numerical values as arguments (as in the case of the temperature in the above example) it is also possible to pass a tolerance indicating how close to the value is considered a match. If no tolerance is passed a default value is used. If multiple results are returned (indicating that the tolerance is too large), *kim query* will return an error. See the [query documentation](#) to see which numerical arguments and tolerances are available for a given *query function*.

### Compute defect formation energy

```
kim      init EAM_Dynamo_ErcolessiAdams_1994_Al__MO_123629422045_005 metal
...
```

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```

... Build fcc crystal containing some defect and compute the total energy
... which is stored in the variable *Etot*
...
kim      query Ec get_cohesive_energy_cubic crystal=[fcc] species=[Al] units=[eV]
variable Eform equal ${Etot} - count(all)*${Ec}
...

```

The defect formation energy  $E_{form}$  is computed by subtracting the ideal fcc cohesive energy of the atoms in the system from  $E_{tot}$ . The ideal fcc cohesive energy of the atoms is obtained from OpenKIM for the Ercolessi and Adams (1994) potential.

#### Retrieve equilibrium fcc crystal of all EAM potentials that support a specific species

```

kim      query model index get_available_models species=[Al] potential_type=[eam]
label   model_loop
kim      query latconst get_lattice_constant_cubic crystal=[fcc] species=[Al]
↳units=[angstrom] model=[${model}]
print   "FCC lattice constant (${model} potential) = ${latconst}"
...
...    do something with current value of latconst
...
next    model
jump    SELF model_loop

```

In this example, the *index* mode of *formatarg* is used. The first *kim query* returns the list of all available EAM potentials that support the *Al* species and archived in OpenKIM. The result of the query operation is stored in the LAMMPS variable *model* as an index *variable*. This variable is used later to access the values one at a time within a loop as shown in the example. The second *kim query* command retrieves from OpenKIM the equilibrium lattice constant predicted by each potential for the fcc structure and places it in variable *latconst*.

---

**Note:** *kim query* commands return results archived in OpenKIM. These results are obtained using programs for computing material properties (KIM Tests and KIM Test Drivers) that were contributed to OpenKIM. In order to give credit to Test developers, the number of times results from these programs are queried is tracked. No other information about the nature of the query or its source is recorded.

---

### 16.64.6 Accessing KIM Model Parameters from LAMMPS (*kim param*)

All IMs are functional forms containing a set of parameters. These parameters' values are typically selected to best reproduce a training set of quantum mechanical calculations or available experimental data. For example, a Lennard-Jones potential intended to model argon might have the values of its two parameters, epsilon, and sigma, fit to the dimer dissociation energy or thermodynamic properties at a critical point of the phase diagram.

Normally a user employing an IM should not modify its parameters since, as noted above, these are selected to reproduce material properties. However, there are cases where accessing and modifying IM parameters is desired, such as for assessing uncertainty, fitting an IM, or working with an ensemble of IMs. As explained *above*, IMs archived in OpenKIM are either Portable Models (PMs) or Simulator Models (SMs). KIM PMs are complete independent implementations of an IM, whereas KIM SMs are wrappers to an IM implemented within LAMMPS. Two different mechanisms are provided for accessing IM parameters in these two cases:

- For a KIM PM, the *kim param* command can be used to *get* and *set* the values of the PM's parameters as explained below.

- For a KIM SM, the user should consult the documentation page for the specific IM and follow instructions there for how to modify its parameters (if possible).

The `kim param get` and `kim param set` commands provide an interface to access and change the parameters of a KIM PM that “publishes” its parameters and makes them publicly available (see the [KIM API documentation](#) for details).

---

**Note:** The `kim param set/get` command must be preceded by a `kim interactions` command (or alternatively by a `pair_style kim` and `pair_coeff` commands). The `kim param set` command may be used wherever a `pair_coeff` command may occur.

---

## Syntax

```
kim param get param_name index_range variable formatarg
kim param set param_name index_range values
```

- `param_name` = name of a KIM portable model parameter (which is published by the PM and available for access). The specific string used to identify a parameter is defined by the PM. For example, for the [Stillinger-Weber \(SW\) potential in OpenKIM](#), the parameter names are *A*, *B*, *p*, *q*, *sigma*, *gamma*, *cutoff*, *lambda*, *costheta0*
- `index_range` = KIM portable model parameter index range (an integer for a single element, or pair of integers separated by a colon for a range of elements)
- `variable(s)` = single name or list of names of (string style) LAMMPS variable(s) where a query result or parameter get result is stored. Variables that do not exist will be created by the command
- `formatarg` = *list*, *split*, or *explicit* (optional): .. parsed-literal:

```
*list* = returns a single string with a list of space separated values
        (e.g. "1.0 2.0 3.0"), which is placed in a LAMMPS variable as
        defined by the *variable* argument.
*split* = returns the values separately in new variables with names based
        on the prefix specified in *variable* and a number appended to
        indicate which element in the list of values is in the variable
*explicit* = returns the values separately in one more more variable names
        provided as arguments that precede *formatarg*\ . [default]
```

- `values` = new value(s) to replace the current value(s) of a KIM portable model parameter

---

**Note:** The list of all the parameters that a PM exposes for access/mutation are automatically written to the lammps log file when `kim init` is called.

---

Each published parameter of a KIM PM takes the form of an array of numerical values. The array can contain one element for a single-valued parameter, or a set of values. For example, the [multispecies SW potential for the Zn-Cd-Hg-S-Se-Te system](#) has the same parameter names as the [single-species SW potential](#), but each parameter array contains 21 entries that correspond to the parameter values used for each pairwise combination of the model’s six supported species (this model does not have parameters specific to individual ternary combinations of its supported species).

The `index_range` argument may either be an integer referring to a specific element within the array associated with the parameter specified by `param_name`, or a pair of integers separated by a colon that refer to a slice of this array. In both cases, one-based indexing is used to refer to the entries of the array.

The result of a `get` operation for a specific `index_range` is stored in one or more [LAMMPS string style variables](#) as determined by the optional `formatarg` argument [documented above](#). If not specified, the default for `formatarg` is “explicit” for the `kim param` command.

For the case where the result is an array with multiple values (i.e. *index\_range* contains a range), the optional “split” or “explicit” *formatarg* keywords can be used to separate the results into multiple variables; see the examples below. Multiple parameters can be retrieved with a single call to *kim param get* by repeating the argument list following *get*.

For a *set* operation, the *values* argument contains the new value(s) for the element(s) of the parameter specified by *index\_range*. For the case where multiple values are being set, *values* contains a set of values separated by spaces. Multiple parameters can be set with a single call to *kim param set* by repeating the argument list following *set*.

### ***kim param* Usage Examples and Further Clarifications**

Examples of getting and setting KIM PM parameters with further clarifications are provided below.

#### **Getting a scalar parameter**

```
kim init SW_StillingerWeber_1985_Si__MO_405512056662_005 metal
...
kim interactions Si
kim param get A 1 VARA
```

or

```
...
pair_style kim SW_StillingerWeber_1985_Si__MO_405512056662_005
pair_coeff * * Si
kim param get A 1 VARA
```

In these cases, the value of the SW *A* parameter is retrieved and placed in the LAMMPS variable *VARA*. The variable *VARA* can be used in the remainder of the input script in the same manner as any other LAMMPS variable.

#### **Getting multiple scalar parameters with a single call**

```
...
kim interactions Si
kim param get A 1 VARA B 1 VARB
```

In this example, it is shown how to retrieve the *A* and *B* parameters of the SW potential and store them in the LAMMPS variables *VARA* and *VARB*.

#### **Getting a range of values from a parameter**

There are several options when getting a range of values from a parameter determined by the *formatarg* argument.

```
kim init SW_ZhouWardMartin_2013_CdTeZnSeHgS__MO_503261197030_002 metal
...
kim interactions Te Zn Se
kim param get lambda 7:9 LAM_TeTe LAM_TeZn LAM_TeSe
```

In this case, *formatarg* is not specified and therefore the default “explicit” mode is used. (The behavior would be the same if the word *explicit* were added after *LAM\_TeSe*.) Elements 7, 8 and 9 of parameter *lambda* retrieved by the *get* operation are placed in the LAMMPS variables *LAM\_TeTe*, *LAM\_TeZn* and *LAM\_TeSe*, respectively.

---

**Note:** In the above example, elements 7-9 of the *lambda* parameter correspond to Te-Te, Te-Zn and Te-Se interactions. This can be determined by visiting the [model page for the specified potential](#) and looking at its parameter file linked to at the bottom of the page (file with *.param* ending) and consulting the README documentation provided with the driver for the PM being used. A link to the driver is provided at the top of the model page.

---



```

...
kim interactions Te Zn Se
kim param get lambda 15:17 LAMS list
variable LAM_VALUE index ${LAMS}
label loop_on_lambda
...
... do something with the current value of lambda
...
next LAM_VALUE
jump SELF loop_on_lambda

```

In this case, the “list” mode of *formatarg* is used. The result of the *get* operation is stored in the LAMMPS variable *LAMS* as a string containing the three retrieved values separated by spaces, e.g “1.0 2.0 3.0”. This can be used in LAMMPS with an *index* variable to access the values one at a time within a loop as shown in the example. At each iteration of the loop *LAM\_VALUE* contains the current value of lambda.

```

...
kim interactions Te Zn Se
kim param get lambda 15:17 LAM split

```

In this case, the “split” mode of *formatarg* is used. The three values retrieved by the *get* operation are stored in the three LAMMPS variables *LAM\_15*, *LAM\_16* and *LAM\_17*. The provided name “LAM” is used as prefix and the location in the lambda array is appended to create the variable names.

### Setting a scalar parameter

```

kim init SW_StillingerWeber_1985_Si__MO_405512056662_005 metal
...
kim interactions Si
kim param set gamma 1 2.6

```

Here, the SW potential’s gamma parameter is set to 2.6. Note that the *get* and *set* commands work together, so that a *get* following a *set* operation will return the new value that was set. For example,

```

...
kim interactions Si
kim param get gamma 1 ORIG_GAMMA
kim param set gamma 1 2.6
kim param get gamma 1 NEW_GAMMA
...
print "original gamma = ${ORIG_GAMMA}, new gamma = ${NEW_GAMMA}"

```

Here, *ORIG\_GAMMA* will contain the original gamma value for the SW potential, while *NEW\_GAMMA* will contain the value 2.6.

### Setting multiple scalar parameters with a single call

```

kim init SW_ZhouWardMartin_2013_CdTeZnSeHgS__MO_503261197030_002 metal
...
kim interactions Cd Te
variable VARG equal 2.6
variable VARS equal 2.0951
kim param set gamma 1 ${VARG} sigma 3 ${VARS}

```

In this case, the first element of the *gamma* parameter and third element of the *sigma* parameter are set to 2.6 and 2.0951, respectively. This example also shows how LAMMPS variables can be used when setting parameters.

#### Setting a range of values of a parameter

```
kim init SW_ZhouWardMartin_2013_CdTeZnSeHgS__M0_503261197030_002 metal
...
kim interactions Cd Te Zn Se Hg S
kim param set sigma 2:6 2.35214 2.23869 2.04516 2.43269 1.80415
```

In this case, elements 2 through 6 of the parameter *sigma* are set to the values 2.35214, 2.23869, 2.04516, 2.43269 and 1.80415 in order.

### 16.64.7 Writing material properties in standard KIM Property Instance format (*kim property*)

The OpenKIM system includes a collection of Tests (material property calculation codes), Models (interatomic potentials), Predictions, and Reference Data (DFT or experiments). Specifically, a KIM Test is a computation that when coupled with a KIM Model generates the prediction of that model for a specific material property rigorously defined by a KIM Property Definition (see the [KIM Properties Framework](#) for further details). A prediction of a material property for a given model is a specific numerical realization of a property definition, referred to as a “Property Instance.” The objective of the *kim property* command is to make it easy to output material properties in a standardized, machine readable, format that can be easily ingested by other programs. Additionally, it aims to make it as easy as possible to convert a LAMMPS script that computes a material property into a KIM Test that can then be uploaded to [openkim.org](http://openkim.org)

A developer interested in creating a KIM Test using a LAMMPS script should first determine whether a property definition that applies to their calculation already exists in OpenKIM by searching the [properties page](#). If none exists, it is possible to use a locally defined property definition contained in a file until it can be uploaded to the official repository (see below). Once one or more applicable property definitions have been identified, the *kim property create*, *kim property modify*, *kim property remove*, and *kim property destroy*, commands provide an interface to create, set, modify, remove, and destroy instances of them within a LAMMPS script.

#### Syntax

```
kim property create instance_id property_id
kim property modify instance_id key key_name key_name_key key_name_value
kim property remove instance_id key key_name
kim property destroy instance_id
kim property dump file
```

- *instance\_id* = a positive integer identifying the KIM property instance; (note that the results file can contain multiple property instances)
- *property\_id* = identifier of a [KIM Property Definition](#), which can be (1) a property short name, (2) the full unique ID of the property (including the contributor and date), (3) a file name corresponding to a local property definition file
- *key\_name* = one of the keys belonging to the specified KIM property definition
- *key\_name\_key* = a key belonging to a key-value pair (standardized in the [KIM Properties Framework](#))
- *key\_name\_value* = value to be associated with a *key\_name\_key* in a key-value pair
- *file* = name of a file to write the currently defined set of KIM property instances to

Examples of each of the three *property\_id* cases are shown below,

```
kim property create 1 atomic-mass
kim property create 2 cohesive-energy-relation-cubic-crystal
```

```
kim property create 1 tag:brunnels@noreply.openkim.org,2016-05-11:property/atomic-mass
kim property create 2 tag:staff@noreply.openkim.org,2014-04-15:property/cohesive-energy-
↪relation-cubic-crystal
```

```
kim property create 1 new-property.edn
kim property create 2 /home/mary/marys-kim-properties/dissociation-energy.edn
```

In the last example, “new-property.edn” and “/home/mary/marys-kim-properties/dissociation-energy.edn” are the names of files that contain user-defined (local) property definitions.

A KIM property instance takes the form of a “map”, i.e. a set of key-value pairs akin to Perl’s hash, Python’s dictionary, or Java’s Hashtable. It consists of a set of property key names, each of which is referred to here by the *key\_name* argument, that are defined as part of the relevant KIM Property Definition and include only lowercase alphanumeric characters and dashes. The value paired with each property key is itself a map whose possible keys are defined as part of the [KIM Properties Framework](#); these keys are referred to by the *key\_name\_key* argument and their associated values by the *key\_name\_value* argument. These values may either be scalars or arrays, as stipulated in the property definition.

**Note:** Each map assigned to a *key\_name* must contain the *key\_name\_key* “source-value” and an associated *key\_name\_value* of the appropriate type (as defined in the relevant KIM Property Definition). For keys that are defined as having physical units, the “source-unit” *key\_name\_key* must also be given a string value recognized by [GNU units](#).

Once a *kim property create* command has been given to instantiate a property instance, maps associated with the property’s keys can be edited using the *kim property modify* command. In using this command, the special keyword “key” should be given, followed by the property key name and the key-value pair in the map associated with the key that is to be set. For example, the *atomic-mass* property definition consists of two property keys named “mass” and “species.” An instance of this property could be created like so:

```
kim property create 1 atomic-mass
kim property modify 1 key species source-value Al
kim property modify 1 key mass source-value 26.98154
kim property modify 1 key mass source-unit amu
```

or, equivalently,

```
kim property create 1 atomic-mass
kim property modify 1 key species source-value Al &
                    key mass source-value 26.98154 &
                    source-unit amu
```

## *kim property* Usage Examples and Further Clarifications

### Create

```
kim property create instance_id property_id
```

The *kim property create* command takes as input a property instance ID and the property definition name, and creates an initial empty property instance data structure. For example,

```
kim property create 1 atomic-mass
kim property create 2 cohesive-energy-relation-cubic-crystal
```

creates an empty property instance of the “atomic-mass” property definition with instance ID 1 and an empty instance of the “cohesive-energy-relation-cubic-crystal” property with ID 2. A list of published property definitions in OpenKIM can be found on the [properties page](#).

One can also provide the name of a file in the current working directory or the path of a file containing a valid property definition. For example,

```
kim property create 1 new-property.edn
```

where “new-property.edn” refers to a file name containing a new property definition that does not exist in OpenKIM.

If the *property\_id* given cannot be found in OpenKIM and no file of this name containing a valid property definition can be found, this command will produce an error with an appropriate message. Calling *kim property create* with the same instance ID multiple times will also produce an error.

### Modify

```
kim property modify instance_id key key_name key_name_key key_name_value
```

The *kim property modify* command incrementally builds the property instance by receiving property definition keys along with associated arguments. Each *key\_name* is associated with a map containing one or more key-value pairs (in the form of *key\_name\_key-key\_name\_value* pairs). For example,

```
kim property modify 1 key species source-value Al
kim property modify 1 key mass source-value 26.98154
kim property modify 1 key mass source-unit amu
```

where the special keyword “key” is followed by a *key\_name* (“species” or “mass” in the above) and one or more key-value pairs. These key-value pairs may continue until either another “key” keyword is given or the end of the command line is reached. Thus, the above could equivalently be written as

```
kim property modify 1 key species source-value Al      &
                    key mass source-value 26.98154 &
                    key mass source-unit amu
```

As an example of modifying multiple key-value pairs belonging to the map of a single property key, the following command modifies the map of the “cohesive-potential-energy” property key to contain the key “source-unit” which is assigned a value of “eV” and the key “digits” which is assigned a value of 5,

```
kim property modify 2 key cohesive-potential-energy source-unit eV digits 5
```

---

**Note:** The relevant data types of the values in the map are handled automatically based on the specification of the key in the KIM Property Definition. In the example above, this means that the value “eV” will automatically be interpreted

as a string while the value 5 will be interpreted as an integer.

The values contained in maps can either be scalars, as in all of the examples above, or arrays depending on which is stipulated in the corresponding Property Definition. For one-dimensional arrays, a single one-based index must be supplied that indicates which element of the array is to be modified. For multidimensional arrays, multiple indices must be given depending on the dimensionality of the array.

**Note:** All array indexing used by *kim property modify* is one-based, i.e. the indices are enumerated 1, 2, 3, ...

**Note:** The dimensionality of arrays are defined in the the corresponding Property Definition. The extent of each dimension of an array can either be a specific finite number or indefinite and determined at run time. If an array has a fixed extent, attempting to modify an out-of-range index will fail with an error message.

For example, the “species” property key of the `cohesive-energy-relation-cubic-crystal` property is a one-dimensional array that can contain any number of entries based on the number of atoms in the unit cell of a given cubic crystal. To assign an array containing the string “Al” four times to the “source-value” key of the “species” property key, we can do so by issuing:

```
kim property modify 2 key species source-value 1 Al
kim property modify 2 key species source-value 2 Al
kim property modify 2 key species source-value 3 Al
kim property modify 2 key species source-value 4 Al
```

**Note:** No declaration of the number of elements in this array was given; *kim property modify* will automatically handle memory management to allow an arbitrary number of elements to be added to the array.

**Note:** In the event that *kim property modify* is used to set the value of an array index without having set the values of all lesser indices, they will be assigned default values based on the data type associated with the key in the map:

Data type	Default value
int	0
float	0.0
string	""
file	""

For example, doing the following:

```
kim property create 2 cohesive-energy-relation-cubic-crystal
kim property modify 2 key species source-value 4 Al
```

will result in the “source-value” key in the map for the property key “species” being assigned the array [“”, “”, “”, “Al”].

For convenience, the index argument provided may refer to an inclusive range of indices by specifying two integers separated by a colon (the first integer must be less than or equal to the second integer, and no whitespace should be included). Thus, the snippet above could equivalently be written:

```
kim property modify 2 key species source-value 1:4 Al Al Al Al
```

Calling this command with a non-positive index, e.g. `kim property modify 2 key species source-value 0 Al`, or an incorrect number of input arguments, e.g. `kim property modify 2 key species source-value 1:4 Al Al`, will result in an error.

As an example of modifying multidimensional arrays, consider the “basis-atoms” key in the `cohesive-energy-relation-cubic-crystal` property definition. This is a two-dimensional array containing the fractional coordinates of atoms in the unit cell of the cubic crystal. In the case of, e.g. a conventional fcc unit cell, the “source-value” key in the map associated with this key should be assigned the following value:

```
[[0.0, 0.0, 0.0],
 [0.5, 0.5, 0.0],
 [0.5, 0.0, 0.5],
 [0.0, 0.5, 0.5]]
```

While each of the twelve components could be set individually, we can instead set each row at a time using colon notation:

```
kim property modify 2 key basis-atom-coordinates source-value 1 1:3 0.0 0.0 0.0
kim property modify 2 key basis-atom-coordinates source-value 2 1:3 0.5 0.5 0.0
kim property modify 2 key basis-atom-coordinates source-value 3 1:3 0.5 0.0 0.5
kim property modify 2 key basis-atom-coordinates source-value 4 1:3 0.0 0.5 0.5
```

Where the first index given refers to a row and the second index refers to a column. We could, instead, choose to set each column at a time like so:

```
kim property modify 2 key basis-atom-coordinates source-value 1:4 1 0.0 0.5 0.5 0.0 &
                    key basis-atom-coordinates source-value 1:4 2 0.0 0.5 0.0 0.5 &
                    key basis-atom-coordinates source-value 1:4 3 0.0 0.0 0.5 0.5
```

**Note:** Multiple calls of `kim property modify` made for the same instance ID can be combined into a single invocation, meaning the following are both valid:

```
kim property modify 2 key basis-atom-coordinates source-value 1 1:3 0.0 0.0 0.0 &
                    key basis-atom-coordinates source-value 2 1:3 0.5 0.5 0.0 &
                    key basis-atom-coordinates source-value 3 1:3 0.5 0.0 0.5 &
                    key basis-atom-coordinates source-value 4 1:3 0.0 0.5 0.5
```

```
kim property modify 2 key short-name source-value 1 fcc &
                    key species source-value 1:4 Al Al Al Al &
                    key a source-value 1:5 3.9149 4.0000 4.032 4.0817 4.1602 &
                    source-unit angstrom &
                    digits 5 &
                    key basis-atom-coordinates source-value 1 1:3 0.0 0.0 0.0 &
                    key basis-atom-coordinates source-value 2 1:3 0.5 0.5 0.0 &
                    key basis-atom-coordinates source-value 3 1:3 0.5 0.0 0.5 &
                    key basis-atom-coordinates source-value 4 1:3 0.0 0.5 0.5
```

**Note:** For multidimensional arrays, only one colon-separated range is allowed in the index listing. Therefore,

```
kim property modify 2 key basis-atom-coordinates 1 1:3 0.0 0.0 0.0
```

is valid but

```
kim property modify 2 key basis-atom-coordinates 1:2 1:3 0.0 0.0 0.0 0.0 0.0 0.0
```

is not.

---

**Note:** After one sets a value in a map with the *kim property modify* command, additional calls will overwrite the previous value.

---

### Remove

```
kim property remove instance_id key key_name
```

The *kim property remove* command can be used to remove a property key from a property instance. For example,

```
kim property remove 2 key basis-atom-coordinates
```

### Destroy

```
kim property destroy instance_id
```

The *kim property destroy* command deletes a previously created property instance ID. For example,

```
kim property destroy 2
```

**Note:** If this command is called with an instance ID that does not exist, no error is raised.

---

### Dump

The *kim property dump* command can be used to write the content of all currently defined property instances to a file:

```
kim property dump file
```

For example,

```
kim property dump results.edn
```

**Note:** Issuing the *kim property dump* command clears all existing property instances from memory.

---

## 16.64.8 Citation of OpenKIM IMs

When publishing results obtained using OpenKIM IMs researchers are requested to cite the OpenKIM project (*Tadmor*), KIM API (*Elliott*), and the specific IM codes used in the simulations, in addition to the relevant scientific references for the IM. The citation format for an IM is displayed on its page on [OpenKIM](#) along with the corresponding BibTex file, and is automatically added to the LAMMPS citation reminder.

Citing the IM software (KIM infrastructure and specific PM or SM codes) used in the simulation gives credit to the researchers who developed them and enables open source efforts like OpenKIM to function.

## 16.64.9 Restrictions

The *kim* command is part of the KIM package. It is only enabled if LAMMPS is built with that package. A requirement for the KIM package, is the KIM API library that must be downloaded from the [OpenKIM website](#) and installed before LAMMPS is compiled. When installing LAMMPS from binary, the kim-api package is a dependency that is automatically downloaded and installed. The *kim query* command requires the *libcurl* library to be installed. The *kim property* command requires *Python* 3.6 or later and the *kim-property* python package to be installed. See the KIM section of the [Packages details](#) for details.

Furthermore, when using *kim* command to run KIM SMs, any packages required by the native potential being used or other commands or fixes that it invokes must be installed.

## 16.64.10 Related commands

*pair\_style kim*

---

**(Tadmor)** Tadmor, Elliott, Sethna, Miller and Becker, JOM, 63, 17 (2011). doi: <https://doi.org/10.1007/s11837-011-0102-6>

**(Elliott)** Elliott, Tadmor and Bernstein, <https://openkim.org/kim-api> (2011) doi: <https://doi.org/10.25950/FF8F563A>

## 16.65 kspace\_modify command

### 16.65.1 Syntax

`kspace_modify keyword value ...`

- one or more keyword/value pairs may be listed
- keyword = *collective* or *compute* or *cutoff/adjust* or *diff* or *disp/auto* or *fftbench* or *force/disp/kspace* or *force/disp/real* or *force* or *gewald/disp* or *gewald* or *kmax/ewald* or *mesh* or *minorder* or *mix/disp* or *order/disp* or *order* or *overlap* or *scafacos* or *slab* or *splittol*

*collective* value = *yes* or *no*

*compute* value = *yes* or *no*

*cutoff/adjust* value = *yes* or *no*

*diff* value = *ad* or *ik* = 2 or 4 FFTs for PPPM in smoothed or non-smoothed mode

*disp/auto* value = *yes* or *no*

*fftbench* value = *yes* or *no*

*force/disp/real* value = accuracy (force units)

*force/disp/kspace* value = accuracy (force units)



*force* value = accuracy (force units)  
*gewald* value = rinv (1/distance units)  
     rinv = G-ewald parameter for Coulombics  
*gewald/disp* value = rinv (1/distance units)  
     rinv = G-ewald parameter for dispersion  
*kmax/ewald* value = kx ky kz  
     kx,ky,kz = number of Ewald sum kspace vectors in each dimension  
*mesh* value = x y z  
     x,y,z = grid size in each dimension for long-range Coulombics  
*mesh/disp* value = x y z  
     x,y,z = grid size in each dimension for 1/r<sup>6</sup> dispersion  
*minororder* value = M  
     M = min allowed extent of Gaussian when auto-adjusting to minimize grid\_  
     ↪communication  
*mix/disp* value = *pair* or *geom* or *none*  
*order* value = N  
     N = extent of Gaussian for PPPM or MSM mapping of charge to grid  
*order/disp* value = N  
     N = extent of Gaussian for PPPM mapping of dispersion term to grid  
*overlap* = *yes* or *no* = whether the grid stencil for PPPM is allowed to overlap into\_  
     ↪more than the nearest-neighbor processor  
*pressure/scalar* value = *yes* or *no*  
*scafacos* values = option value1 value2 ...  
     option = *tolerance*  
         value = *energy* or *energy\_rel* or *field* or *field\_rel* or *potential* or *potential\_rel*  
     option = *fmm\_tuning*  
         value = 0 or 1  
*slab* value = *volfactor* or *nozforce*  
     volfactor = ratio of the total extended volume used in the  
         2d approximation compared with the volume of the simulation domain  
     ew2d EW2D correction (available with ELECTRODE package)  
     nozforce turns off kspace forces in the z direction  
*splittol* value = tol  
     tol = relative size of two eigenvalues (see discussion below)  
*wire* value = volfactor (available with ELECTRODE package)  
     volfactor = ratio of the total extended dimension used in the 1d  
         approximation compared with the dimension of the simulation domain

## 16.65.2 Examples

```

kspace_modify mesh 24 24 30 order 6
kspace_modify slab 3.0
kspace_modify scafacos tolerance energy
  
```

### 16.65.3 Description

Set parameters used by the *k*space solvers defined by the *k*space\_style command. Not all parameters are relevant to all *k*space styles.

---

The *collective* keyword applies only to PPPM. It is set to *no* by default, except on IBM BlueGene machines. If this option is set to *yes*, LAMMPS will use MPI collective operations to remap data for 3d-FFT operations instead of the default point-to-point communication. This is faster on IBM BlueGene machines, and may also be faster on other machines if they have an efficient implementation of MPI collective operations and adequate hardware.

---

The *compute* keyword allows Kspace computations to be turned off, even though a *k*space\_style is defined. This is not useful for running a real simulation, but can be useful for debugging purposes or for computing only partial forces that do not include the Kspace contribution. You can also do this by simply not defining a *k*space\_style, but a Kspace-compatible *pair\_style* requires a *k*space style to be defined. This keyword gives you that option.

---

The *cutoff/adjust* keyword applies only to MSM. If this option is turned on, the Coulombic cutoff will be automatically adjusted at the beginning of the run to give the desired estimated error. Other cutoffs such as LJ will not be affected. If the grid is not set using the *mesh* command, this command will also attempt to use the optimal grid that minimizes cost using an estimate given by (*Hardy*). Note that this cost estimate is not exact, somewhat experimental, and still may not yield the optimal parameters.

---

The *diff* keyword specifies the differentiation scheme used by the PPPM method to compute forces on particles given electrostatic potentials on the PPPM mesh. The *ik* approach is the default for PPPM and is the original formulation used in (*Hockney*). It performs differentiation in Kspace, and uses 3 FFTs to transfer each component of the computed fields back to real space for total of 4 FFTs per timestep.

The analytic differentiation *ad* approach uses only 1 FFT to transfer information back to real space for a total of 2 FFTs per timestep. It then performs analytic differentiation on the single quantity to generate the 3 components of the electric field at each grid point. This is sometimes referred to as “smoothed” PPPM. This approach requires a somewhat larger PPPM mesh to achieve the same accuracy as the *ik* method. Currently, only the *ik* method (default) can be used for a triclinic simulation cell with PPPM. The *ad* method is always used for MSM.

---

**Note:** Currently, not all PPPM styles support the *ad* option. Support for those PPPM variants will be added later.

---

The *disp/auto* option controls whether the *pppm/disp* is allowed to generate PPPM parameters automatically. If set to *no*, parameters have to be specified using the *gewald/disp*, *mesh/disp*, *force/disp/real* or *force/disp/k*space keywords, or the code will stop with an error message. When this option is set to *yes*, the error message will not appear and the simulation will start. For a typical application, using the automatic parameter generation will provide simulations that are either inaccurate or slow. Using this option is thus not recommended. For guidelines on how to obtain good parameters, see the *How-To* discussion.

---

The *ffibench* keyword applies only to PPPM. It is off by default. If this option is turned on, LAMMPS will perform a short FFT benchmark computation and report its timings, and will thus finish some seconds later than it would if this option were off.

---

The *force/disp/real* and *force/disp/kSPACE* keywords set the force accuracy for the real and reciprocal space computations for the dispersion part of *pppm/disp*. As shown in (*Isele-Holder*), optimal performance and accuracy in the results is obtained when these values are different.

---

The *force* keyword overrides the relative accuracy parameter set by the *kSPACE\_style* command with an absolute accuracy. The accuracy determines the RMS error in per-atom forces calculated by the long-range solver and is thus specified in force units. A negative value for the accuracy setting means to use the relative accuracy parameter. The accuracy setting is used in conjunction with the pairwise cutoff to determine the number of K-space vectors for style *ewald*, the FFT grid size for style *pppm*, or the real space grid size for style *msm*.

---

The *gewald* keyword sets the value of the Ewald or PPPM G-ewald parameter for charge as *rinv* in reciprocal distance units. Without this setting, LAMMPS chooses the parameter automatically as a function of cutoff, precision, grid spacing, etc. This means it can vary from one simulation to the next which may not be desirable for matching a KSpace solver to a pre-tabulated pairwise potential. This setting can also be useful if Ewald or PPPM fails to choose a good grid spacing and G-ewald parameter automatically. If the value is set to 0.0, LAMMPS will choose the G-ewald parameter automatically. MSM does not use the *gewald* parameter.

---

The *gewald/disp* keyword sets the value of the Ewald or PPPM G-ewald parameter for dispersion as *rinv* in reciprocal distance units. It has the same meaning as the *gewald* setting for Coulombics.

---

The *kmax/ewald* keyword sets the number of kspace vectors in each dimension for kspace style *ewald*. The three values must be positive integers, or else (0,0,0), which unsets the option. When this option is not set, the Ewald sum scheme chooses its own kspace vectors, consistent with the user-specified accuracy and pairwise cutoff. In any case, if kspace style *ewald* is invoked, the values used are printed to the screen and the log file at the start of the run.

---

The *mesh* keyword sets the grid size for kspace style *pppm* or *msm*. In the case of PPPM, this is the FFT mesh, and each dimension must be factorizable into powers of 2, 3, and 5. In the case of MSM, this is the finest scale real-space mesh, and each dimension must be factorizable into powers of 2. When this option is not set, the PPPM or MSM solver chooses its own grid size, consistent with the user-specified accuracy and pairwise cutoff. Values for x,y,z of 0,0,0 unset the option.

---

The *mesh/disp* keyword sets the grid size for kspace style *pppm/disp*. This is the FFT mesh for long-range dispersion and each dimension must be factorizable into powers of 2, 3, and 5. When this option is not set, the PPPM solver chooses its own grid size, consistent with the user-specified accuracy and pairwise cutoff. Values for x,y,z of 0,0,0 unset the option.

---

The *minorder* keyword allows LAMMPS to reduce the *order* setting if necessary to keep the communication of ghost grid point limited to exchanges between nearest-neighbor processors. See the discussion of the *overlap* keyword for details. If the *overlap* keyword is set to *yes*, which is the default, this is never needed. If it set to *no* and overlap occurs, then LAMMPS will reduce the order setting, one step at a time, until the ghost grid overlap only extends to nearest neighbor processors. The *minorder* keyword limits how small the *order* setting can become. The minimum allowed value for PPPM is 2, which is the default. If *minorder* is set to the same value as *order* then no reduction is allowed, and LAMMPS will generate an error if the grid communication is non-nearest-neighbor and *overlap* is set to *no*. The *minorder* keyword is not currently supported in MSM.

---

The *mix/disp* keyword selects the mixing rule for the dispersion coefficients. With *pair*, the dispersion coefficients of unlike types are computed as indicated with *pair\_modify*. With *geom*, geometric mixing is enforced on the dispersion coefficients in the *k*space coefficients. When using the arithmetic mixing rule, this will speed-up the simulations but introduces some error in the force computations, as shown in (Wennberg). With *none*, it is assumed that no mixing rule is applicable. Splitting of the dispersion coefficients will be performed as described in (Isele-Holder).

This splitting can be influenced with the *splittol* keywords. Only the eigenvalues that are larger than *tol* compared to the largest eigenvalues are included. Using this keywords the original matrix of dispersion coefficients is approximated. This leads to faster computations, but the accuracy in the reciprocal space computations of the dispersion part is decreased.

---

The *order* keyword determines how many grid spacings an atom's charge extends when it is mapped to the grid in *k*space style *pppm* or *msm*. The default for this parameter is 5 for PPPM and 8 for MSM, which means each charge spans 5 or 8 grid cells in each dimension, respectively. For the LAMMPS implementation of MSM, the order can range from 4 to 10 and must be even. For PPPM, the minimum allowed setting is 2 and the maximum allowed setting is 7. The larger the value of this parameter, the smaller that LAMMPS will set the grid size, to achieve the requested accuracy. Conversely, the smaller the order value, the larger the grid size will be. Note that there is an inherent trade-off involved: a small grid will lower the cost of FFTs or MSM direct sum, but a larger order parameter will increase the cost of interpolating charge/fields to/from the grid.

The PPPM order parameter may be reset by LAMMPS when it sets up the FFT grid if the implied grid stencil extends beyond the grid cells owned by neighboring processors. Typically this will only occur when small problems are run on large numbers of processors. A warning will be generated indicating the order parameter is being reduced to allow LAMMPS to run the problem. Automatic adjustment of the order parameter is not supported in MSM.

---

The *order/disp* keyword determines how many grid spacings an atom's dispersion term extends when it is mapped to the grid in *k*space style *pppm/disp*. It has the same meaning as the *order* setting for Coulombics.

---

The *overlap* keyword can be used in conjunction with the *minorder* keyword with the PPPM styles to adjust the amount of communication that occurs when values on the FFT grid are exchanged between processors. This communication is distinct from the communication inherent in the parallel FFTs themselves, and is required because processors interpolate charge and field values using grid point values owned by neighboring processors (i.e. ghost point communication). If the *overlap* keyword is set to *yes* then this communication is allowed to extend beyond nearest-neighbor processors, e.g. when using lots of processors on a small problem. If it is set to *no* then the communication will be limited to nearest-neighbor processors and the *order* setting will be reduced if necessary, as explained by the *minorder* keyword discussion. The *overlap* keyword is always set to *yes* in MSM.

---

The *pressure/scalar* keyword applies only to MSM. If this option is turned on, only the scalar pressure (i.e.  $(P_{xx} + P_{yy} + P_{zz})/3.0$ ) will be computed, which can be used, for example, to run an isotropic barostat. Computing the full pressure tensor with MSM is expensive, and this option provides a faster alternative. The scalar pressure is computed using a relationship between the Coulombic energy and pressure (Hummer) instead of using the virial equation. This option cannot be used to access individual components of the pressure tensor, to compute per-atom virial, or with suffix *k*space/*pair* styles of MSM, like OMP or GPU.

---

The *scafacos* keyword is used for settings that are passed to the ScaFaCoS library when using *k*space\_style *scafacos*.

The *tolerance* option affects how the *accuracy* specified with the *k*space\_style command is interpreted by ScaFaCoS. The following values may be used:

- energy = absolute accuracy in total Coulombic energy

- `energy_rel` = relative accuracy in total Coulombic energy
- `potential` = absolute accuracy in total Coulombic potential
- `potential_rel` = relative accuracy in total Coulombic potential
- `field` = absolute accuracy in electric field
- `field_rel` = relative accuracy in electric field

The values with suffix `_rel` indicate the tolerance is a relative tolerance; the other values impose an absolute tolerance on the given quantity. Absolute tolerance in this case means, that for a given quantity  $q$  and a given absolute tolerance of  $t_a$  the result should be between  $q-t_a$  and  $q+t_a$ . For a relative tolerance  $t_r$  the relative error should not be greater than  $t_r$ , i.e.  $\text{abs}(1 - (\text{result}/q)) < t_r$ . As a consequence of this, the tolerance type should be checked, when performing computations with a high absolute field / energy. E.g. if the total energy in the system is 1000000.0 an absolute tolerance of  $1e-3$  would mean that the result has to be between 999999.999 and 1000000.001, which would be equivalent to a relative tolerance of  $1e-9$ .

The `energy` and `energy_rel` values, set a tolerance based on the total Coulombic energy of the system. The `potential` and `potential_rel` set a tolerance based on the per-atom Coulombic energy. The `field` and `field_rel` tolerance types set a tolerance based on the electric field values computed by ScaFaCoS. Since per-atom forces are derived from the per-atom electric field, this effectively sets a tolerance on the forces, similar to other LAMMPS KSpace styles, as explained on the [kpace\\_style](#) doc page.

Note that not all ScaFaCoS solvers support all tolerance types. These are the allowed values for each method:

- `fmm` = energy and `energy_rel`
- `p2nfft` = field (1d-,2d-,3d-periodic systems) or potential (0d-periodic)
- `p3m` = field
- `ewald` = field
- `direct` = has no tolerance tuning

If the tolerance type is not changed, the default values for the tolerance type are the first values in the above list, e.g. `energy` is the default tolerance type for the `fmm` solver.

The `fmm_tuning` option is only relevant when using the FMM method. It activates (value=1) or deactivates (value=0) an internal tuning mechanism for the FMM solver. The tuning operation runs sequentially and can be very time-consuming. Usually it is not needed for systems with a homogeneous charge distribution. The default for this option is therefore 0. The FMM internal tuning is performed once, when the solver is set up.

---

The `slab` keyword allows an Ewald or PPPM solver to be used for a systems that are periodic in x,y but non-periodic in z - a [boundary](#) setting of “boundary p p f”. This is done by treating the system as if it were periodic in z, but inserting empty volume between atom slabs and removing dipole inter-slab interactions so that slab-slab interactions are effectively turned off. The `volfactor` value sets the ratio of the extended dimension in z divided by the actual dimension in z. The recommended value is 3.0. A larger value is inefficient; a smaller value introduces unwanted slab-slab interactions. The use of fixed boundaries in z means that the user must prevent particle migration beyond the initial z-bounds, typically by providing a wall-style fix. The methodology behind the `slab` option is explained in the paper by (Yeh). The `slab` option is also extended to non-neutral systems ([Ballenegger](#)).

An alternative slab option can be invoked with the `nozforce` keyword in lieu of the `volfactor`. This turns off all kspace forces in the z direction. The `nozforce` option is not supported by MSM. For MSM, any combination of periodic, non-periodic, or shrink-wrapped boundaries can be set using [boundary](#) (the slab approximation in not needed). The `slab` keyword is not currently supported by Ewald or PPPM when using a triclinic simulation cell. The slab correction has also been extended to point dipole interactions ([Klapp](#)) in [kpace\\_style ewald/disp](#), [ewald/dipole](#), and [pppm/dipole](#).

**Note:** If you wish to apply an electric field in the Z-direction, in conjunction with the *slab* keyword, you should do it by adding explicit charged particles to the +/- Z surfaces. If you do it via the *fix efield* command, it will not give the correct dielectric constant due to the Yeh/Berkowitz (*Yeh*) correction not being compatible with how *fix efield* works.

---

The *force/disp/real* and *force/disp/kspace* keywords set the force accuracy for the real and reciprocal space computations for the dispersion part of pppm/disp. As shown in (*Isele-Holder*), optimal performance and accuracy in the results is obtained when these values are different.

The *disp/auto* option controls whether the pppm/disp is allowed to generate PPPM parameters automatically. If set to *no*, parameters have to be specified using the *gewald/disp*, *mesh/disp*, *force/disp/real* or *force/disp/kspace* keywords, or the code will stop with an error message. When this option is set to *yes*, the error message will not appear and the simulation will start. For a typical application, using the automatic parameter generation will provide simulations that are either inaccurate or slow. Using this option is thus not recommended. For guidelines on how to obtain good parameters, see the *Howto dispersion* doc page.

---

## 16.65.4 Restrictions

none

## 16.65.5 Related commands

*kspace\_style*, *boundary*

## 16.65.6 Default

The option defaults are *mesh = mesh/disp = 0 0 0*, *order = order/disp = 5* (PPPM), *order = 10* (MSM), *minorder = 2*, *overlap = yes*, *force = -1.0*, *gewald = gewald/disp = 0.0*, *slab = 1.0*, *compute = yes*, *cutoff/adjust = yes* (MSM), *pressure/scalar = yes* (MSM), *fftbench = no* (PPPM), *diff = ik* (PPPM), *mix/disp = pair*, *force/disp/real = -1.0*, *force/disp/kspace = -1.0*, *split = 0*, *tol = 1.0e-6*, and *disp/auto = no*. For *pppm/intel*, *order = order/disp = 7*. For *scafacos* settings, the *scafacos* tolerance option depends on the method chosen, as documented above. The *scafacos fmm\_tuning* default = 0.

---

**(Hockney)** Hockney and Eastwood, Computer Simulation Using Particles, Adam Hilger, NY (1989).

**(Yeh)** Yeh and Berkowitz, J Chem Phys, 111, 3155 (1999).

**(Ballenegger)** Ballenegger, Arnold, Cerda, J Chem Phys, 131, 094107 (2009).

**(Klapp)** Klapp, Schoen, J Chem Phys, 117, 8050 (2002).

**(Hardy)** David Hardy thesis: Multilevel Summation for the Fast Evaluation of Forces for the Simulation of Biomolecules, University of Illinois at Urbana-Champaign, (2006).

**(Hummer)** Hummer, Gronbeck-Jensen, Neumann, J Chem Phys, 109, 2791 (1998)

**(Isele-Holder)** Isele-Holder, Mitchell, Hammond, Kohlmeyer, Ismail, J Chem Theory Comput, 9, 5412 (2013).

**(Wennberg)** Wennberg, Murtola, Hess, Lindahl, J Chem Theory Comput, 9, 3527 (2013).

## 16.66 `kspace_style` command

### 16.66.1 Syntax

```
kspace_style style value
```

- `style` = *none* or *ewald* or *ewald/dipole* or *ewald/dipole/spin* or *ewald/disp* or *ewald/disp/dipole* or *ewald/omp* or *ewald/electrode* or *pppm* or *pppm/cg* or *pppm/disp* or *pppm/tip4p* or *pppm/stagger* or *pppm/disp/tip4p* or *pppm/gpu* or *pppm/intel* or *pppm/disp/intel* or *pppm/kk* or *pppm/omp* or *pppm/cg/omp* or *pppm/disp/tip4p/omp* or *pppm/tip4p/omp* or *pppm/dielectric* or *pppm/disp/dielectric* or *pppm/electrode* or *pppm/electrode/intel* or *msm* or *msm/cg* or *msm/omp* or *msm/cg/omp* or *msm/dielectric* or *scafacos*

*none* value = none

*ewald* value = accuracy

accuracy = desired relative error in forces

*ewald/dipole* value = accuracy

accuracy = desired relative error in forces

*ewald/dipole/spin* value = accuracy

accuracy = desired relative error in forces

*ewald/disp* value = accuracy

accuracy = desired relative error in forces

*ewald/disp/dipole* value = accuracy

accuracy = desired relative error in forces

*ewald/omp* value = accuracy

accuracy = desired relative error in forces

*ewald/electrode* value = accuracy

accuracy = desired relative error in forces

*pppm* value = accuracy

accuracy = desired relative error in forces

*pppm/cg* values = accuracy (smallq)

accuracy = desired relative error in forces

smallq = cutoff for charges to be considered (optional) (charge units)

*pppm/dipole* value = accuracy

accuracy = desired relative error in forces

*pppm/dipole/spin* value = accuracy

accuracy = desired relative error in forces

*pppm/disp* value = accuracy

accuracy = desired relative error in forces

*pppm/tip4p* value = accuracy

accuracy = desired relative error in forces

*pppm/disp/tip4p* value = accuracy

accuracy = desired relative error in forces

*pppm/gpu* value = accuracy

accuracy = desired relative error in forces

*pppm/intel* value = accuracy

accuracy = desired relative error in forces

*pppm/disp/intel* value = accuracy

accuracy = desired relative error in forces

*pppm/kk* value = accuracy

accuracy = desired relative error in forces

*pppm/omp* value = accuracy

accuracy = desired relative error in forces

*pppm/cg/omp* values = accuracy (smallq)

accuracy = desired relative error in forces  
smallq = cutoff for charges to be considered (optional) (charge units)  
*pppm/disp/omp* value = accuracy  
accuracy = desired relative error in forces  
*pppm/tip4p/omp* value = accuracy  
accuracy = desired relative error in forces  
*pppm/disp/tip4p/omp* value = accuracy  
accuracy = desired relative error in forces  
*pppm/stagger* value = accuracy  
accuracy = desired relative error in forces  
*pppm/dielectric* value = accuracy  
accuracy = desired relative error in forces  
*pppm/disp/dielectric* value = accuracy  
accuracy = desired relative error in forces  
*pppm/electrode* value = accuracy  
accuracy = desired relative error in forces  
*pppm/electrode/intel* value = accuracy  
accuracy = desired relative error in forces  
*msm* value = accuracy  
accuracy = desired relative error in forces  
*msm/cg* value = accuracy (smallq)  
accuracy = desired relative error in forces  
smallq = cutoff for charges to be considered (optional) (charge units)  
*msm/omp* value = accuracy  
accuracy = desired relative error in forces  
*msm/cg/omp* value = accuracy (smallq)  
accuracy = desired relative error in forces  
smallq = cutoff for charges to be considered (optional) (charge units)  
*msm/dielectric* value = accuracy  
accuracy = desired relative error in forces  
*scafacos* values = method accuracy  
method = fmm or p2nfft or p3m or ewald or direct  
accuracy = desired relative error in forces

## 16.66.2 Examples

```
kpspace_style pppm 1.0e-4  
kpspace_style pppm/cg 1.0e-5 1.0e-6  
kpspace_style msm 1.0e-4  
kpspace_style scafacos fmm 1.0e-4  
kpspace_style none
```

Used in input scripts:

```
examples/peptide/in.peptide
```



### 16.66.3 Description

Define a long-range solver for LAMMPS to use each timestep to compute long-range Coulombic interactions or long-range  $1/r^6$  interactions. Most of the long-range solvers perform their computation in K-space, hence the name of this command.

When such a solver is used in conjunction with an appropriate pair style, the cutoff for Coulombic or  $1/r^N$  interactions is effectively infinite. If the Coulombic case, this means each charge in the system interacts with charges in an infinite array of periodic images of the simulation domain.

Note that using a long-range solver requires use of a matching *pair style* to perform consistent short-range pairwise calculations. This means that the name of the pair style contains a matching keyword to the name of the KSpace style, as in this table:

Pair style	KSpace style
coul/long	ewald or ppm
coul/msm	msm
lj/long or buck/long	disp (for dispersion)
tip4p/long	tip4p
dipole/long	dipole

The *ewald* style performs a standard Ewald summation as described in any solid-state physics text.

The *ewald/disp* style adds a long-range dispersion sum option for  $1/r^6$  potentials and is useful for simulation of interfaces (*Veld*). It also performs standard Coulombic Ewald summations, but in a more efficient manner than the *ewald* style. The  $1/r^6$  capability means that Lennard-Jones or Buckingham potentials can be used without a cutoff, i.e. they become full long-range potentials.

The *ewald/disp/dipole* style can also be used with point-dipoles, see (*Toukmaji*).

The *ewald/dipole* style adds long-range standard Ewald summations for dipole-dipole interactions, see (*Toukmaji*).

The *ewald/dipole/spin* style adds long-range standard Ewald summations for magnetic dipole-dipole interactions between magnetic spins.

The *pppm* style invokes a particle-particle particle-mesh solver (*Hockney*) which maps atom charge to a 3d mesh, uses 3d FFTs to solve Poisson's equation on the mesh, then interpolates electric fields on the mesh points back to the atoms. It is closely related to the particle-mesh Ewald technique (PME) (*Darden*) used in AMBER and CHARMM. The cost of traditional Ewald summation scales as  $N^{3/2}$  where  $N$  is the number of atoms in the system. The PPPM solver scales as  $N \log N$  due to the FFTs, so it is almost always a faster choice (*Pollock*).

The *pppm/cg* style is identical to the *pppm* style except that it has an optimization for systems where most particles are uncharged. Similarly the *msm/cg* style implements the same optimization for *msm*. The optional *smallq* argument defines the cutoff for the absolute charge value which determines whether a particle is considered charged or not. Its default value is 1.0e-5.

The *pppm/dipole* style invokes a particle-particle particle-mesh solver for dipole-dipole interactions, following the method of (*Cerda*).

The *pppm/dipole/spin* style invokes a particle-particle particle-mesh solver for magnetic dipole-dipole interactions between magnetic spins.

The *pppm/tip4p* style is identical to the *pppm* style except that it adds a charge at the massless fourth site in each TIP4P water molecule. It should be used with *pair styles* with a *tip4p/long* in their style name.

The *pppm/stagger* style performs calculations using two different meshes, one shifted slightly with respect to the other. This can reduce force aliasing errors and increase the accuracy of the method for a given mesh size. Or a coarser mesh can be used for the same target accuracy, which saves CPU time. However, there is a trade-off since FFTs on two meshes are now performed which increases the computation required. See (*Cerutti*), (*Neelov*), and (*Hockney*) for details of the method.

For high relative accuracy, using staggered PPPM allows the mesh size to be reduced by a factor of 2 in each dimension as compared to regular PPPM (for the same target accuracy). This can give up to a 4x speedup in the KSpace time (8x less mesh points, 2x more expensive). However, for low relative accuracy, the staggered PPPM mesh size may be essentially the same as for regular PPPM, which means the method will be up to 2x slower in the KSpace time (simply 2x more expensive). For more details and timings, see the *Speed tips* doc page.

---

**Note:** Using *pppm/stagger* may not give the same increase in the accuracy of energy and pressure as it does in forces, so some caution must be used if energy and/or pressure are quantities of interest, such as when using a barostat.

---

The *pppm/disp* and *pppm/disp/tip4p* styles add a mesh-based long-range dispersion sum option for  $1/r^6$  potentials (*Isele-Holder*), similar to the *ewald/disp* style. The  $1/r^6$  capability means that Lennard-Jones or Buckingham potentials can be used without a cutoff, i.e. they become full long-range potentials.

For these styles, you will possibly want to adjust the default choice of parameters by using the *kpace\_modify* command. This can be done by either choosing the Ewald and grid parameters, or by specifying separate accuracies for the real and kspace calculations. When not making any settings, the simulation will stop with an error message. Further information on the influence of the parameters and how to choose them is described in (*Isele-Holder*), (*Isele-Holder2*) and the *Howto dispersion* doc page.

---

**Note:** All of the PPPM styles can be used with single-precision FFTs by using the compiler switch `-DFFT_SINGLE` for the `FFT_INC` setting in your low-level Makefile. This setting also changes some of the PPPM operations (e.g. mapping charge to mesh and interpolating electric fields to particles) to be performed in single precision. This option can speed-up long-range calculations, particularly in parallel or on GPUs. The use of the `-DFFT_SINGLE` flag is discussed on the *Build settings* doc page. MSM does not currently support the `-DFFT_SINGLE` compiler switch.

---

The *electrode* styles add methods that are required for the constant potential method implemented in *fix electrode/\**. The styles *ewald/electrode*, *pppm/electrode* and *pppm/electrode/intel* are available. These styles do not support the *kpace\_modify slab nozforce* command.

---

The *msm* style invokes a multi-level summation method MSM solver, (*Hardy*) or (*Hardy2*), which maps atom charge to a 3d mesh, and uses a multi-level hierarchy of coarser and coarser meshes on which direct Coulomb solvers are done. This method does not use FFTs and scales as  $N$ . It may therefore be faster than the other K-space solvers for relatively large problems when running on large core counts. MSM can also be used for non-periodic boundary conditions and for mixed periodic and non-periodic boundaries.

MSM is most competitive versus Ewald and PPPM when only relatively low accuracy forces, about  $1e-4$  relative error or less accurate, are needed. Note that use of a larger Coulombic cutoff (i.e. 15 angstroms instead of 10 angstroms) provides better MSM accuracy for both the real space and grid computed forces.

Currently calculation of the full pressure tensor in MSM is expensive. Using the *kpace\_modify pressure/scalar yes* command provides a less expensive way to compute the scalar pressure  $(P_{xx} + P_{yy} + P_{zz})/3.0$ . The scalar pressure can be used, for example, to run an isotropic barostat. If the full pressure tensor is needed, then calculating the pressure at every timestep or using a fixed pressure simulation with MSM will cause the code to run slower.

The *scafacos* style is a wrapper on the [ScaFaCoS Coulomb solver library](#) which provides a variety of solver methods which can be used with LAMMPS. The paper by ([Sutman](#)) gives an overview of ScaFaCoS.

ScaFaCoS was developed by a consortium of German research facilities with a BMBF (German Ministry of Science and Education) funded project in 2009-2012. Participants of the consortium were the Universities of Bonn, Chemnitz, Stuttgart, and Wuppertal as well as the Forschungszentrum Juelich.

The library is available for download at “<http://scafacos.de>” or can be cloned from the git-repository “<https://github.com/scafacos/scafacos.git>”.

In order to use this KSpace style, you must download and build the ScaFaCoS library, then build LAMMPS with the SCAFACOS package installed package which links LAMMPS to the ScaFaCoS library. See details on [this page](#).

---

**Note:** Unlike other KSpace solvers in LAMMPS, ScaFaCoS computes all Coulombic interactions, both short- and long-range. Thus you should NOT use a Coulombic pair style when using `kspace_style scafacos`. This also means the total Coulombic energy (short- and long-range) will be tallied for *thermodynamic output* command as part of the *elong* keyword; the *ecoul* keyword will be zero.

---

---

**Note:** See the current restriction below about use of ScaFaCoS in LAMMPS with molecular charged systems or the TIP4P water model.

---

The specified *method* determines which ScaFaCoS algorithm is used. These are the ScaFaCoS methods currently available from LAMMPS:

- *fmm* = Fast Multi-Pole method
- *p2nfft* = FFT-based Coulomb solver
- *ewald* = Ewald summation
- *direct* = direct  $O(N^2)$  summation
- *p3m* = PPPM

We plan to support additional ScaFaCoS solvers from LAMMPS in the future. For an overview of the included solvers, refer to ([Sutmann](#))

The specified *accuracy* is similar to the accuracy setting for other LAMMPS KSpace styles, but is passed to ScaFaCoS, which can interpret it in different ways for different methods it supports. Within the ScaFaCoS library the *accuracy* is treated as a tolerance level (either absolute or relative) for the chosen quantity, where the quantity can be either the Columic field values, the per-atom Columic energy or the total Columic energy. To select from these options, see the [kpace\\_modify scafacos accuracy](#) doc page.

The *kpace\_modify scafacos* command also explains other ScaFaCoS options currently exposed to LAMMPS.

---

The specified *accuracy* determines the relative RMS error in per-atom forces calculated by the long-range solver. It is set as a dimensionless number, relative to the force that two unit point charges (e.g. 2 monovalent ions) exert on each other at a distance of 1 Angstrom. This reference value was chosen as representative of the magnitude of electrostatic forces in atomic systems. Thus an accuracy value of 1.0e-4 means that the RMS error will be a factor of 10000 smaller than the reference force.

The accuracy setting is used in conjunction with the pairwise cutoff to determine the number of K-space vectors for style *ewald* or the grid size for style *pppm* or *msm*.

Note that style *pppm* only computes the grid size at the beginning of a simulation, so if the length or triclinic tilt of the simulation cell increases dramatically during the course of the simulation, the accuracy of the simulation may degrade. Likewise, if the *kpace\_modify slab* option is used with shrink-wrap boundaries in the z-dimension, and the box size changes dramatically in z. For example, for a triclinic system with all three tilt factors set to the maximum limit, the PPPM grid should be increased roughly by a factor of 1.5 in the y direction and 2.0 in the z direction as compared to the same system using a cubic orthogonal simulation cell. One way to handle this issue if you have a long simulation where the box size changes dramatically, is to break it into shorter simulations (multiple *run* commands). This works because the grid size is re-computed at the beginning of each run. Another way to ensure the described accuracy requirement is met is to run a short simulation at the maximum expected tilt or length, note the required grid size, and then use the *kpace\_modify mesh* command to manually set the PPPM grid size to this value for the long run. The simulation then will be “too accurate” for some portion of the run.

RMS force errors in real space for *ewald* and *pppm* are estimated using equation 18 of (Kolafa), which is also referenced as equation 9 of (Petersen). RMS force errors in K-space for *ewald* are estimated using equation 11 of (Petersen), which is similar to equation 32 of (Kolafa). RMS force errors in K-space for *pppm* are estimated using equation 38 of (Deserno). RMS force errors for *msm* are estimated using ideas from chapter 3 of (Hardy), with equation 3.197 of particular note. When using *msm* with non-periodic boundary conditions, it is expected that the error estimation will be too pessimistic. RMS force errors for dipoles when using *ewald/disp* or *ewald/dipole* are estimated using equations 33 and 46 of (Wang). The RMS force errors for *pppm/dipole* are estimated using the equations in (Cerda).

See the *kpace\_modify* command for additional options of the K-space solvers that can be set, including a *force* option for setting an absolute RMS error in forces, as opposed to a relative RMS error.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

**Note:** For the GPU package, the *pppm/gpu* style performs charge assignment and force interpolation calculations on the GPU. These processes are performed either in single or double precision, depending on whether the `-DFFT_SINGLE` setting was specified in your low-level Makefile, as discussed above. The FFTs themselves are still calculated on the CPU. If *pppm/gpu* is used with a GPU-enabled pair style, part of the PPPM calculation can be performed concurrently on the GPU while other calculations for non-bonded and bonded force calculation are performed on the CPU.

---

**Note:** For the KOKKOS package, the *pppm/kk* style performs charge assignment and force interpolation calculations, along with the FFTs themselves, on the GPU or (optionally) threaded on the CPU when using OpenMP and FFTW3.

---

## 16.66.4 Restrictions

Note that the long-range electrostatic solvers in LAMMPS assume conducting metal (tin foil) boundary conditions for both charge and dipole interactions. Vacuum boundary conditions are not currently supported.

The *ewald/disp*, *ewald*, *pppm*, and *msm* styles support non-orthogonal (triclinic symmetry) simulation boxes. However, triclinic simulation cells may not yet be supported by all suffix versions of these styles.

Most of the base *kspace* styles are part of the KSPACE package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The *msm/dielectric* and *pppm/dielectric* *kspace* styles are part of the DIELECTRIC package. They are only enabled if LAMMPS was built with that package **and** the KSPACE package. See the *Build package* page for more info.

For MSM, a simulation must be 3d and one can use any combination of periodic, non-periodic, but not shrink-wrapped boundaries (specified using the *boundary* command).

For Ewald and PPPM, a simulation must be 3d and periodic in all dimensions. The only exception is if the slab option is set with *kpace\_modify*, in which case the xy dimensions must be periodic and the z dimension must be non-periodic.

The *scafacos* KSpace style will only be enabled if LAMMPS is built with the SCAFACOS package. See the *Build package* doc page for more info.

The use of ScaFaCos in LAMMPS does not yet support molecular charged systems where the short-range Coulombic interactions between atoms in the same bond/angle/dihedral are weighted by the *special\_bonds* command. Likewise it does not support the “TIP4P water style” where a fictitious charge site is introduced in each water molecule. Finally, the methods *p3m* and *ewald* do not support computing the virial, so this contribution is not included.

## 16.66.5 Related commands

*kpace\_modify*, *pair\_style lj/cut/coul/long*, *pair\_style lj/charmm/coul/long*, *pair\_style lj/long/coul/long*, *pair\_style buck/coul/long*

## 16.66.6 Default

```
kpace_style none
```

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**(Deserno)** Deserno and Holm, J Chem Phys, 109, 7694 (1998).

**(Hockney)** Hockney and Eastwood, Computer Simulation Using Particles, Adam Hilger, NY (1989).

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**(Petersen)** Petersen, J Chem Phys, 103, 3668 (1995).

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**(Pollock)** Pollock and Glosli, Comp Phys Comm, 95, 93 (1996).

**(Cerutti)** Cerutti, Duke, Darden, Lybrand, Journal of Chemical Theory and Computation 5, 2322 (2009)

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**(Isele-Holder)** Isele-Holder, Mitchell, Ismail, J Chem Phys, 137, 174107 (2012).

**(Isele-Holder2)** Isele-Holder, Mitchell, Hammond, Kohlmeyer, Ismail, J Chem Theory Comput 9, 5412 (2013).

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**(Hardy2)** Hardy, Stone, Schulten, Parallel Computing, 35, 164-177 (2009).

**(Sutmann)** Sutmann, Arnold, Fahrenberger, et. al., Physical review / E 88(6), 063308 (2013)

**(Cerda)** Cerda, Ballenegger, Lenz, Holm, J Chem Phys 129, 234104 (2008)

**(Sutmann) G. Sutmann. ScaFaCoS - a Scalable library of Fast Coulomb Solvers for particle Systems.**

In Bajaj, Zavattieri, Koslowski, Siegmund, Proceedings of the Society of Engineering Science 51st Annual Technical Meeting. 2014.

## 16.67 label command

### 16.67.1 Syntax

```
label ID
```

- ID = string used as label name

### 16.67.2 Examples

```
label xyz  
label loop
```

### 16.67.3 Description

Label this line of the input script with the chosen ID. Unless a jump command was used previously, this does nothing. But if a *jump* command was used with a label argument to begin invoking this script file, then all command lines in the script prior to this line will be ignored. I.e. execution of the script will begin at this line. This is useful for looping over a section of the input script as discussed in the *jump* command.

### 16.67.4 Restrictions

none

### 16.67.5 Related commands

*jump, next*

## 16.67.6 Default

none

## 16.68 labelmap command

### 16.68.1 Syntax

```
labelmap option args
```

- *option* = *atom* or *bond* or *angle* or *dihedral* or *improper* or *clear* or *write*  
*clear* = no args  
*write* arg = filename  
*atom* or *bond* or *angle* or *dihedral* or *improper*  
args = list of one or more numeric-type/type-label pairs

### 16.68.2 Examples

```
labelmap atom 3 carbon 4 'c3"' 5 "c1'" 6 "c#"
labelmap bond 1 carbonyl 2 nitrile 3 "" c1'-c2" ""
labelmap atom $(label2type(atom,carbon)) C # change type label from 'carbon' to 'C'
labelmap clear
labelmap write mymap.include
```

### 16.68.3 Description

New in version 15Sep2022.

Define alphanumeric type labels to associate with one or more numeric atom, bond, angle, dihedral or improper types. A collection of type labels for all atom types, bond types, etc. is stored as a label map.

The label map can also be defined by the *read\_data* command when it reads these sections in a data file: Atom Type Labels, Bond Type Labels, etc. See the *Howto type labels* doc page for a general discussion of how type labels can be used.

Valid type labels can contain any alphanumeric character, but must not start with a number, a '#', or a '\*' character. They can contain other standard ASCII characters such as angular or square brackets '<' and '>' or '[' and ']', parenthesis '(' and ')', dash '-', underscore '\_', plus '+' and equals '=' signs and more. They must not contain blanks or any other whitespace. Note that type labels must be put in single or double quotation marks if they contain the '#' character or if they contain a double (") or single quotation mark ('). If the label contains both a single and a double quotation mark, then triple quotation (""") must be used. When enclosing a type label with quotation marks, the LAMMPS input parser may require adding leading or trailing blanks around the type label so it can identify the enclosing quotation marks. Those blanks will be removed when defining the label.

A *labelmap* command can only modify the label map for one type-kind (atom types, bond types, etc). Any number of numeric-type/type-label pairs may follow. If a type label already exists for the same numeric type, it will be overwritten. Type labels must be unique; assigning the same type label to multiple numeric types within the same type-kind is not allowed. When reading and writing data files, it is required that there is a label defined for *every* numeric type within a given type-kind in order to write out the type label section for that type-kind.

The *clear* option resets the label map and thus discards all previous settings.

The *write* option takes a filename as argument and writes the current label mappings to a file as a sequence of *labelmap* commands, so the file can be copied into a new LAMMPS input file or read in using the *include* command.

---

## 16.68.4 Restrictions

This command must come after the simulation box is defined by a *read\_data*, *read\_restart*, or *create\_box* command.

Label maps are currently not supported when using the KOKKOS package.

## 16.68.5 Related commands

*read\_data*, *write\_data*, *molecule*, *fix bond/react*

## 16.68.6 Default

none

## 16.69 lattice command

### 16.69.1 Syntax

```
lattice style scale keyword values ...
```

- style = *none* or *sc* or *bcc* or *fcc* or *hcp* or *diamond* or *sq* or *sq2* or *hex* or *custom*
- scale = scale factor between lattice and simulation box  
scale = reduced density rho\* (for LJ units)  
scale = lattice constant in distance units (for all other units)
- zero or more keyword/value pairs may be appended
- keyword = *origin* or *orient* or *spacing* or *a1* or *a2* or *a3* or *basis*  
*origin* values = x y z  
x,y,z = fractions of a unit cell ( $0 \leq x,y,z < 1$ )  
*orient* values = dim i j k  
dim = x or y or z  
i,j,k = integer lattice directions  
*spacing* values = dx dy dz  
dx,dy,dz = lattice spacings in the x,y,z box directions  
*a1,a2,a3* values = x y z  
x,y,z = primitive vector components that define unit cell  
*basis* values = x y z  
x,y,z = fractional coords of a basis atom ( $0 \leq x,y,z < 1$ )



## 16.69.2 Examples

```
lattice fcc 3.52
lattice hex 0.85
lattice sq 0.8 origin 0.0 0.5 0.0 orient x 1 1 0 orient y -1 1 0
lattice custom 3.52 a1 1.0 0.0 0.0 a2 0.5 1.0 0.0 a3 0.0 0.0 0.5 &
                basis 0.0 0.0 0.0 basis 0.5 0.5 0.5
lattice none 2.0
```

## 16.69.3 Description

Define a lattice for use by other commands. In LAMMPS, a lattice is simply a set of points in space, determined by a unit cell with basis atoms, that is replicated infinitely in all dimensions. The arguments of the lattice command can be used to define a wide variety of crystallographic lattices.

A lattice is used by LAMMPS in two ways. First, the *create\_atoms* command creates atoms on the lattice points inside the simulation box. Note that the *create\_atoms* command allows different atom types to be assigned to different basis atoms of the lattice. Second, the lattice spacing in the x,y,z dimensions implied by the lattice, can be used by other commands as distance units (e.g. *create\_box*, *region* and *velocity*), which are often convenient to use when the underlying problem geometry is atoms on a lattice.

The lattice style must be consistent with the dimension of the simulation - see the *dimension* command. Styles *sc* or *bcc* or *fcc* or *hcp* or *diamond* are for 3d problems. Styles *sq* or *sq2* or *hex* are for 2d problems. Style *custom* can be used for either 2d or 3d problems.

A lattice consists of a unit cell, a set of basis atoms within that cell, and a set of transformation parameters (scale, origin, orient) that map the unit cell into the simulation box. The vectors *a1*, *a2*, *a3* are the edge vectors of the unit cell. This is the nomenclature for “primitive” vectors in solid-state crystallography, but in LAMMPS the unit cell they determine does not have to be a “primitive cell” of minimum volume.

Note that the lattice command can be used multiple times in an input script. Each time it is invoked, the lattice attributes are re-defined and are used for all subsequent commands (that use lattice attributes). For example, a sequence of *lattice*, *region*, and *create\_atoms* commands can be repeated multiple times to build a poly-crystalline model with different geometric regions populated with atoms in different lattice orientations.

A lattice of style *none* does not define a unit cell and basis set, so it cannot be used with the *create\_atoms* command. However it does define a lattice spacing via the specified scale parameter. As explained above the lattice spacings in x,y,z can be used by other commands as distance units. No additional keyword/value pairs can be specified for the *none* style. By default, a “lattice none 1.0” is defined, which means the lattice spacing is the same as one distance unit, as defined by the *units* command.

Lattices of style *sc*, *fcc*, *bcc*, and *diamond* are 3d lattices that define a cubic unit cell with edge length = 1.0. This means *a1* = 1 0 0, *a2* = 0 1 0, and *a3* = 0 0 1. Style *hcp* has *a1* = 1 0 0, *a2* = 0 sqrt(3) 0, and *a3* = 0 0 sqrt(8/3). The placement of the basis atoms within the unit cell are described in any solid-state physics text. A *sc* lattice has 1 basis atom at the lower-left-bottom corner of the cube. A *bcc* lattice has 2 basis atoms, one at the corner and one at the center of the cube. A *fcc* lattice has 4 basis atoms, one at the corner and 3 at the cube face centers. A *hcp* lattice has 4 basis atoms, two in the *z* = 0 plane and 2 in the *z* = 0.5 plane. A *diamond* lattice has 8 basis atoms.

Lattices of style *sq* and *sq2* are 2d lattices that define a square unit cell with edge length = 1.0. This means *a1* = 1 0 0 and *a2* = 0 1 0. A *sq* lattice has 1 basis atom at the lower-left corner of the square. A *sq2* lattice has 2 basis atoms, one at the corner and one at the center of the square. A *hex* style is also a 2d lattice, but the unit cell is rectangular, with *a1* = 1 0 0 and *a2* = 0 sqrt(3) 0. It has 2 basis atoms, one at the corner and one at the center of the rectangle.

A lattice of style *custom* allows you to specify *a1*, *a2*, *a3*, and a list of basis atoms to put in the unit cell. By default, *a1* and *a2* and *a3* are 3 orthogonal unit vectors (edges of a unit cube). But you can specify them to be of any length and

non-orthogonal to each other, so that they describe a tilted parallelepiped. Via the *basis* keyword you add atoms, one at a time, to the unit cell. Its arguments are fractional coordinates ( $0.0 \leq x,y,z < 1.0$ ). The position vector  $x$  of a basis atom within the unit cell is thus a linear combination of the unit cell's 3 edge vectors, i.e.  $x = b_x a_1 + b_y a_2 + b_z a_3$ , where  $b_x, b_y, b_z$  are the 3 values specified for the *basis* keyword.

---

This sub-section discusses the arguments that determine how the idealized unit cell is transformed into a lattice of points within the simulation box.

The *scale* argument determines how the size of the unit cell will be scaled when mapping it into the simulation box. I.e. it determines a multiplicative factor to apply to the unit cell, to convert it to a lattice of the desired size and distance units in the simulation box. The meaning of the *scale* argument depends on the *units* being used in your simulation.

For all unit styles except *lj*, the scale argument is specified in the distance units defined by the unit style. For example, in *real* or *metal* units, if the unit cell is a unit cube with edge length 1.0, specifying *scale* = 3.52 would create a cubic lattice with a spacing of 3.52 Angstroms. In *cgs* units, the spacing would be 3.52 cm.

For unit style *lj*, the scale argument is the Lennard-Jones reduced density, typically written as  $\rho^*$ . LAMMPS converts this value into the multiplicative factor via the formula “ $\text{factor}^{\text{dim}} = \rho/\rho^*$ ”, where  $\rho = N/V$  with  $V$  = the volume of the lattice unit cell and  $N$  = the number of basis atoms in the unit cell (described below), and  $\text{dim} = 2$  or  $3$  for the dimensionality of the simulation. Effectively, this means that if LJ particles of size  $\sigma = 1.0$  are used in the simulation, the lattice of particles will be at the desired reduced density.

The *origin* option specifies how the unit cell will be shifted or translated when mapping it into the simulation box. The  $x,y,z$  values are fractional values ( $0.0 \leq x,y,z < 1.0$ ) meaning shift the lattice by a fraction of the lattice spacing in each dimension. The meaning of “lattice spacing” is discussed below.

The *orient* option specifies how the unit cell will be rotated when mapping it into the simulation box. The *dim* argument is one of the 3 coordinate axes in the simulation box. The other 3 arguments are the crystallographic direction in the lattice that you want to orient along that axis, specified as integers. E.g. “orient x 2 1 0” means the x-axis in the simulation box will be the [210] lattice direction, and similarly for y and z. The 3 lattice directions you specify do not have to be unit vectors, but they must be mutually orthogonal and obey the right-hand rule, i.e. (X cross Y) points in the Z direction.

---

**Note:** The preceding paragraph describing lattice directions is only valid for orthogonal cubic unit cells (or square in 2d). If you are using a *hcp* or *hex* lattice or the more general lattice style *custom* with non-orthogonal  $a_1, a_2, a_3$  vectors, then you should think of the 3 *orient* vectors as creating a 3x3 rotation matrix which is applied to  $a_1, a_2, a_3$  to rotate the original unit cell to a new orientation in the simulation box.

---

Several LAMMPS commands have the option to use distance units that are inferred from “lattice spacings” in the  $x,y,z$  box directions. E.g. the *region* command can create a block of size 10x20x20, where 10 means 10 lattice spacings in the x direction.

---

**Note:** Though they are called lattice spacings, all the commands that have a “units lattice” option, simply use the 3 values as scale factors on the distance units defined by the *units* command. Thus if you do not like the lattice spacings computed by LAMMPS (e.g. for a non-orthogonal or rotated unit cell), you can define the 3 values to be whatever you wish, via the *spacing* option.

---

If the *spacing* option is not specified, the lattice spacings are computed by LAMMPS in the following way. A unit cell of the lattice is mapped into the simulation box (scaled and rotated), so that it now has (perhaps) a modified size and orientation. The lattice spacing in X is defined as the difference between the min/max extent of the x coordinates of the 8 corner points of the modified unit cell (4 in 2d). Similarly, the Y and Z lattice spacings are defined as the difference in the min/max of the y and z coordinates.

Note that if the unit cell is orthogonal with axis-aligned edges (no rotation via the *orient* keyword), then the lattice spacings in each dimension are simply the scale factor (described above) multiplied by the length of  $a_1, a_2, a_3$ . Thus a *hex* style lattice with a scale factor of 3.0 Angstroms, would have a lattice spacing of 3.0 in x and  $3*\sqrt{3}$  in y.

**Note:** For non-orthogonal unit cells and/or when a rotation is applied via the *orient* keyword, then the lattice spacings computed by LAMMPS are typically less intuitive. In particular, in these cases, there is no guarantee that a particular lattice spacing is an integer multiple of the periodicity of the lattice in that direction. Thus, if you create an orthogonal periodic simulation box whose size in a dimension is a multiple of the lattice spacing, and then fill it with atoms via the *create\_atoms* command, you will NOT necessarily create a periodic system. I.e. atoms may overlap incorrectly at the faces of the simulation box.

The *spacing* option sets the 3 lattice spacings directly. All must be non-zero (use 1.0 for dz in a 2d simulation). The specified values are multiplied by the multiplicative factor described above that is associated with the scale factor. Thus a spacing of 1.0 means one unit cell edge length independent of the scale factor. As mentioned above, this option can be useful if the spacings LAMMPS computes are inconvenient to use in subsequent commands, which can be the case for non-orthogonal or rotated lattices.

Note that whenever the lattice command is used, the values of the lattice spacings LAMMPS calculates are printed out. Thus their effect in commands that use the spacings should be decipherable.

Example commands for generating a Wurtzite crystal. The lattice constants approximate those of CdSe. The  $\sqrt{3} \times 1$  orthorhombic supercell is used with the x, y, and z directions oriented along  $[\bar{1}230]$ ,  $[10\bar{1}0]$ , and  $[0001]$ , respectively.

```
variable a equal 4.34
variable b equal $a*sqrt(3.0)
variable c equal $a*sqrt(8.0/3.0)

variable third equal 1.0/3.0
variable five6 equal 5.0/6.0

lattice custom 1.0 &
  a1 $b 0.0 0.0 &
  a2 0.0 $a 0.0 &
  a3 0.0 0.0 $c &
  basis 0.0 0.0 0.0 &
  basis 0.5 0.5 0.0 &
  basis ${third} 0.0 0.5 &
  basis ${five6} 0.5 0.5 &
  basis 0.0 0.0 0.625 &
  basis 0.5 0.5 0.625 &
  basis ${third} 0.0 0.125 &
  basis ${five6} 0.5 0.125

region myreg block 0 1 0 1 0 1
create_box 2 myreg
create_atoms 1 box &
  basis 5 2 &
  basis 6 2 &
  basis 7 2 &
  basis 8 2
```

## 16.69.4 Restrictions

The *a1,a2,a3,basis* keywords can only be used with style *custom*.

## 16.69.5 Related commands

*dimension, create\_atoms, region*

## 16.69.6 Default

```
lattice none 1.0
```

For other lattice styles, the option defaults are origin = 0.0 0.0 0.0, orient = x 1 0 0, orient = y 0 1 0, orient = z 0 0 1, a1 = 1 0 0, a2 = 0 1 0, and a3 = 0 0 1.

## 16.70 log command

### 16.70.1 Syntax

```
log file keyword
```

- file = name of new logfile
- keyword = *append* if output should be appended to logfile (optional)

### 16.70.2 Examples

```
log log.equil  
log log.equil append
```

### 16.70.3 Description

This command closes the current LAMMPS log file, opens a new file with the specified name, and begins logging information to it. If the specified file name is *none*, then no new log file is opened. If the optional keyword *append* is specified, then output will be appended to an existing log file, instead of overwriting it.

If multiple processor partitions are being used, the file name should be a variable, so that different processors do not attempt to write to the same log file.

The file “log.lammps” is the default log file for a LAMMPS run. The name of the initial log file can also be set by the *-log command-line switch*.

## 16.70.4 Restrictions

none

## 16.70.5 Related commands

none

## 16.70.6 Default

The default LAMMPS log file is named log.lammps

# 16.71 mass command

## 16.71.1 Syntax

```
mass I value
```

- I = atom type (see asterisk form below), or type label
- value = mass

## 16.71.2 Examples

```
mass 1 1.0
mass * 62.5
mass 2* 62.5

labelmap atom 1 C
mass C 12.01
```

## 16.71.3 Description

Set the mass for all atoms of one or more atom types. Per-type mass values can also be set in the *read\_data* data file using the “Masses” keyword. See the *units* command for what mass units to use.

The I index can be specified in one of several ways. An explicit numeric value can be used, as in the first example above. Or I can be a type label, which is an alphanumeric string defined by the *labelmap* command or in a section of a data file read by the *read\_data* command, and which converts internally to a numeric type. Or a wild-card asterisk can be used to set the mass for multiple atom types. This takes the form “\*” or “\*n” or “n\*” or “m\*n”, where m and n are numbers. If N = the number of atom types, then an asterisk with no numeric values means all types from 1 to N. A leading asterisk means all types from 1 to n (inclusive). A trailing asterisk means all types from n to N (inclusive). A middle asterisk means all types from m to n (inclusive).

A line in a *data file* that follows the “Masses” keyword specifies mass using the same format as the arguments of the mass command in an input script, except that no wild-card asterisk can be used. For example, under the “Masses” section of a data file, the line that corresponds to the first example above would be listed as

1 1.0

Note that the mass command can only be used if the *atom style* requires per-type atom mass to be set. Currently, all but the *sphere* and *ellipsoid* and *peri* styles do. They require mass to be set for individual particles, not types. Per-atom masses are defined in the data file read by the *read\_data* command, or set to default values by the *create\_atoms* command. Per-atom masses can also be set to new values by the *set mass* or *set density* commands.

Also note that *pair\_style eam* and *pair\_style bop* commands define the masses of atom types in their respective potential files, in which case the mass command is normally not used.

If you define a *hybrid atom style* which includes one (or more) sub-styles which require per-type mass and one (or more) sub-styles which require per-atom mass, then you must define both. However, in this case the per-type mass will be ignored; only the per-atom mass will be used by LAMMPS.

### 16.71.4 Restrictions

This command must come after the simulation box is defined by a *read\_data*, *read\_restart*, or *create\_box* command.

All masses must be defined before a simulation is run. They must also all be defined before a *velocity* or *fix shake* command is used.

The mass assigned to any type or atom must be > 0.0.

### 16.71.5 Related commands

none

### 16.71.6 Default

none

## 16.72 mdi command

### 16.72.1 Syntax

mdi option args

- option = *engine* or *plugin* or *connect* or *exit*
  - engine* args = zero or more keyword/args pairs
    - keywords = *elements*
    - elements* args = N\_1 N\_2 ... N\_ntypes
    - N\_1,N\_2,...N\_ntypes = atomic number for each of ntypes LAMMPS atom types
  - plugin* args = name keyword value keyword value ...
    - name = name of plugin library, e.g. lammps means a liblammps.so library will be loaded
    - keyword/value pairs in any order, some are required, some are optional
    - keywords = *mdi* or *infile* or *extra* or *command*
    - mdi* value = args passed to MDI for driver to operate with plugins (required)
    - infile* value = filename the engine will read at start-up (optional)

`extra` value = additional command-line args to pass to engine library when loaded.  
 ↔(optional)  
`command` value = a LAMMPS input script command to execute (required)  
`connect` args = none  
`exit` args = none

## 16.72.2 Examples

```

mdi engine
mdi engine elements 13 29
mdi plugin lammmps mdi "-role ENGINE -name lammmps -method LINK" &
    infile in.aimd.engine extra "-log log.aimd.engine.plugin" &
    command "run 5"
mdi connect
mdi exit
  
```

## 16.72.3 Description

This command implements operations within LAMMPS to use the *MDI Library* <[https://molssi-mdi.github.io/MDI\\_Library/html/index.html](https://molssi-mdi.github.io/MDI_Library/html/index.html)> for coupling to other codes in a client/server protocol.

See the Howto MDI doc page for a discussion of all the different ways 2 or more codes can interact via MDI.

The `examples/mdi` directory has examples which use LAMMPS in 4 different modes: as a driver using an engine as either a stand-alone code or as a plugin, and as an engine operating as either a stand-alone code or as a plugin. The README file in that directory shows how to launch and couple codes for all the 4 usage modes, and so they communicate via the MDI library using either MPI or sockets.

The scripts in that directory illustrate the use of all the options for this command.

The *engine* option enables LAMMPS to act as an MDI engine (server), responding to requests from an MDI driver (client) code.

The *plugin* option enables LAMMPS to act as an MDI driver (client), and load the MDI engine (server) code as a library plugin. In this case the MDI engine is a library plugin. An MDI engine can also be a stand-alone code, launched separately from LAMMPS, in which case the `mdi plugin` command is not used.

The *connect* and *exit* options are only used when LAMMPS is acting as an MDI driver. As explained below, these options are normally not needed, except for a specific kind of use case.

The *mdi engine* command is used to make LAMMPS operate as an MDI engine. It is typically used in an input script after LAMMPS has setup the system it is going to model consistent with what the driver code expects. Depending on when the driver code tells the LAMMPS engine to exit, other commands can be executed after this command, but typically it is used at the end of a LAMMPS input script.

To act as an MDI engine operating as an MD code (or surrogate QM code), this is the list of standard MDI commands issued by a driver code which LAMMPS currently recognizes. Using standard commands defined by the MDI library means that a driver code can work interchangeably with LAMMPS or other MD codes or with QM codes which support the MDI standard. See more details about these commands in the [MDI library documentation](#)

These commands are valid at the @DEFAULT node defined by MDI. Commands that start with “>” mean the driver is sending information to LAMMPS. Commands that start with “<” are requests by the driver for LAMMPS to send it information. Commands that start with an alphabetic letter perform actions. Commands that start with “@” are MDI “node” commands, which are described further below.

Command name	Action
>CELL or <CELL	Send/request 3 simulation box edge vectors (9 values)
>CELL_DISPL or <CELL_DISPL	Send/request displacement of the simulation box from the origin (3 values)
>CHARGES or <CHARGES	Send/request charge on each atom (N values)
>COORDS or <COORDS	Send/request coordinates of each atom (3N values)
>ELEMENTS	Send elements (atomic numbers) for each atom (N values)
<ENERGY	Request total energy (potential + kinetic) of the system (1 value)
>FORCES or <FORCES	Send/request forces on each atom (3N values)
>+FORCES	Send forces to add to each atom (3N values)
<LABELS	Request string label of each atom (N values)
<MASSES	Request mass of each atom (N values)
MD	Perform an MD simulation for N timesteps (most recent >NSTEPS value)
OPTG	Perform an energy minimization to convergence (most recent >TOLERANCE values)
>NATOMS or <NATOMS	Sends/request number of atoms in the system (1 value)
>NSTEPS	Send number of timesteps for next MD dynamics run via MD command
<PE	Request potential energy of the system (1 value)
<STRESS	Request symmetric stress tensor (virial) of the system (9 values)
>TOLERANCE	Send 4 tolerance parameters for next MD minimization via OPTG command
>TYPES or <TYPES	Send/request the LAMMPS atom type for each atom (N values)
>VELOCITIES or <VELOCITIES	Send/request the velocity of each atom (3N values)
@INIT_MD or @INIT_OPTG	Driver tells LAMMPS to start single-step dynamics or minimization (see below)
EXIT	Driver tells LAMMPS to exit engine mode

**Note:** The <ENERGY, <FORCES, <PE, and <STRESS commands trigger LAMMPS to compute atomic interactions for the current configuration of atoms and size/shape of the simulation box. I.e. LAMMPS invokes its pair, bond, angle, ..., kspace styles. If the driver is updating the atom coordinates and/or box incrementally (as in an MD simulation which the driver is managing), then the LAMMPS engine will do the same, and only occasionally trigger neighbor list builds. If the change in atom positions is large (since the previous >COORDS command), then LAMMPS will do a more expensive operation to migrate atoms to new processors as needed and re-neighbor. If the >NATOMS or >TYPES or >ELEMENTS commands have been sent (since the previous >COORDS command), then LAMMPS assumes the system is new and re-initializes an entirely new simulation.

**Note:** The >TYPES or >ELEMENTS commands are how the MDI driver tells the LAMMPS engine which LAMMPS atom type to assign to each atom. If both the MDI driver and the LAMMPS engine are initialized so that atom type values are consistent in both codes, then the >TYPES command can be used. If not, the optional *elements* keyword can be used to specify what element each LAMMPS atom type corresponds to. This is specified by the atomic number of the element, e.g. 13 for Al. An atomic number must be specified for each of the ntypes LAMMPS atom types. Ntypes is typically specified via the `create_box` command or in the data file read by the `read_data` command. In this has been done, the MDI driver can send an >ELEMENTS command to the LAMMPS driver with the atomic number of each atom.



The MD and OPTG commands perform an entire MD simulation or energy minimization (to convergence) with no communication from the driver until the simulation is complete. By contrast, the @INIT\_MD and @INIT\_OPTG commands allow the driver to communicate with the engine at each timestep of a dynamics run or iteration of a minimization; see more info below.

The MD command performs a simulation using the most recent >NSTEPS value. The OPTG command performs a minimization using the 4 convergence parameters from the most recent >TOLERANCE command. The 4 parameters sent are those used by the *minimize* command in LAMMPS: etol, ftol, maxiter, and maxeval.

The mdi engine command also implements the following custom MDI commands which are LAMMPS-specific. These commands are also valid at the @DEFAULT node defined by MDI:

- – Command name
  - Action
- – >NBYTES
  - Send # of datums in a subsequent command (1 value)
- – >COMMAND
  - Send a LAMMPS input script command as a string (Nbytes in length)
- – >COMMANDS
  - Send multiple LAMMPS input script commands as a newline-separated string (Nbytes in length)
- – >INFILE
  - Send filename of an input script to execute (filename Nbytes in length)
- – <KE
  - Request kinetic energy of the system (1 value)

Note that other custom commands can easily be added if these are not sufficient to support what a user-written driver code needs. Code to support new commands can be added to the MDI package within LAMMPS, specifically to the src/MDI/mdi\_engine.cpp file.

MDI also defines a standard mechanism for the driver to request that an MD engine (LAMMPS) perform a dynamics simulation one step at a time or an energy minimization one iteration at a time. This is so that the driver can (optionally) communicate with LAMMPS at intermediate points of the timestep or iteration by issuing MDI node commands which start with “@”.

To tell LAMMPS to run dynamics in single-step mode, the driver sends as @INIT\_MD command followed by the these commands. The driver can interact with LAMMPS at 3 node locations within each timestep: @COORDS, @FORCES, @ENDSTEP:

- – Command name
  - Action
- – @COORDS
  - Proceed to next @COORDS node = post-integrate location in LAMMPS timestep
- – @FORCES
  - Proceed to next @FORCES node = post-force location in LAMMPS timestep
- – @ENDSTEP
  - Proceed to next @ENDSTEP node = end-of-step location in LAMMPS timestep
- – @DEFAULT

- Exit MD simulation, return to @DEFAULT node
- – EXIT
- Driver tells LAMMPS to exit the MD simulation and engine mode

To tell LAMMPS to run an energy minimization in single-iteration mode. The driver can interact with LAMMPS at 2 node locations within each iteration of the minimizer: @COORDS, @FORCES:

- – Command name
  - Action
- – @COORDS
  - Proceed to next @COORDS node = min-pre-force location in LAMMPS min iteration
- – @FORCES
  - Proceed to next @FORCES node = min-post-force location in LAMMPS min iteration
- – @DEFAULT
  - Exit minimization, return to @DEFAULT node
- – EXIT
  - Driver tells LAMMPS to exit the minimization and engine mode

While LAMMPS is at its @COORDS node, the following standard MDI commands are supported, as documented above: >COORDS or <COORDS, @COORDS, @FORCES, @ENDSTEP, @DEFAULT, EXIT.

While LAMMPS is at its @FORCES node, the following standard MDI commands are supported, as documented above: <COORDS, <ENERGY, >FORCES or >+FORCES or <FORCES, <KE, <PE, <STRESS, @COORDS, @FORCES, @ENDSTEP, @DEFAULT, EXIT.

While LAMMPS is at its @ENDSTEP node, the following standard MDI commands are supported, as documented above: <ENERGY, <FORCES, <KE, <PE, <STRESS, @COORDS, @FORCES, @ENDSTEP, @DEFAULT, EXIT.

---

The *mdi plugin* command is used to make LAMMPS operate as an MDI driver which loads an MDI engine as a plugin library. It is typically used in an input script after LAMMPS has setup the system it is going to model consistent with the engine code.

The *name* argument specifies which plugin library to load. A name like “lammps” is converted to a filename liblammps.so. The path for where this file is located is specified by the *-plugin\_path* switch within the *-mdi* command-line switch, which is specified when LAMMPS is launched. See the examples/mdi/README files for examples of how this is done.

The *mdi* keyword is required and is used as the *-mdi* argument passed to the library when it is launched. The *-role* and *-method* settings are required. The *-name* setting can be anything you choose. MDI drivers and engines can query their names to verify they are values they expect.

The *infile* keyword is optional. It sets the name of an input script which the engine will open and process. MDI will pass it as a command-line argument to the library when it is launched. The file typically contains settings that an MD or QM code will use for its calculations.

The *extra* keyword is optional. It contains additional command-line arguments which MDI will pass to the library when it is launched.

The *command* keyword is required. It specifies a LAMMPS input script command (as a single argument in quotes if it is multiple words). Once the plugin library is launched, LAMMPS will execute this command. Other previously-defined commands in the input script, such as the *fix mdi/qm* command, should perform MDI communication with the engine,

while the specified *command* executes. Note that if *command* is an *include* command, then it could specify a filename with multiple LAMMPS commands.

---

**Note:** When the *command* is complete, LAMMPS will send an MDI EXIT command to the plugin engine and the plugin will be removed. The “mdi plugin” command will then exit and the next command (if any) in the LAMMPS input script will be processed. A subsequent “mdi plugin” command could then load the same or a different MDI plugin if desired.

---

The *mdi connect* and *mdi exit* commands are only used when LAMMPS is operating as an MDI driver. And when other LAMMPS command(s) which send MDI commands and associated data to/from the MDI engine are not able to initiate and terminate the connection to the engine code.

The only current MDI driver command in LAMMPS is the *fix mdi/qm* command. If it is only used once in an input script then it can initiate and terminate the connection. But if it is being issued multiple times, e.g. in a loop that issues a *clear* command, then it cannot initiate or terminate the connection multiple times. Instead, the *mdi connect* and *mdi exit* commands should be used outside the loop to initiate or terminate the connection.

See the examples/mdi/in.series.driver script for an example of how this is done. The LOOP in that script is reading a series of data file configurations and passing them to an MDI engine (e.g. quantum code) for energy and force evaluation. A *clear* command inside the loop wipes out the current system so a new one can be defined. This operation also destroys all fixes. So the *fix mdi/qm* command is issued once per loop iteration. Note that it includes a “connect no” option which disables the initiate/terminate logic within that fix.

## 16.72.4 Restrictions

This command is part of the MDI package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

To use LAMMPS in conjunction with other MDI-enabled atomistic codes, the *units* command should be used to specify *real* or *metal* units. This will ensure the correct unit conversions between LAMMPS and MDI units, which the other codes will also perform in their preferred units.

LAMMPS can also be used as an MDI engine in other unit choices it supports, e.g. *lj*, but then no unit conversion is performed.

## 16.72.5 Related commands

*fix mdi/qm*

## 16.72.6 Default

None

## 16.73 min\_modify command

### 16.73.1 Syntax

```
min_modify keyword values ...
```

- one or more keyword/value pairs may be listed

keyword = *dmax* or *line* or *norm* or *alpha\_damp* or *discrete\_factor* or *integrator* or *tmax*

→ *tmax*

*dmax* value = max

max = maximum distance for line search to move (distance units)

*line* value = *backtrack* or *quadratic* or *forcezero* or *spin\_cubic* or *spin\_none*

*backtrack*, *quadratic*, *forcezero*, *spin\_cubic*, *spin\_none* = style of linesearch to use

*norm* value = *two* or *inf* or *max*

*two* = Euclidean two-norm (length of 3N vector)

*inf* = max force component across all 3-vectors

*max* = max force norm across all 3-vectors

*alpha\_damp* value = damping

damping = fictitious Gilbert damping for spin minimization (adim)

*discrete\_factor* value = factor

factor = discretization factor for adaptive spin timestep (adim)

*integrator* value = *eulerimplicit* or *verlet*

time integration scheme for fire minimization

*tmax* value = factor

factor = maximum adaptive timestep for fire minimization (adim)

### 16.73.2 Examples

```
min_modify dmax 0.2
min_modify integrator verlet tmax 4
```

### 16.73.3 Description

This command sets parameters that affect the energy minimization algorithms selected by the *min\_style* command. The various settings may affect the convergence rate and overall number of force evaluations required by a minimization, so users can experiment with these parameters to tune their minimizations.

The *cg* and *sd* minimization styles have an outer iteration and an inner iteration which is steps along a one-dimensional line search in a particular search direction. The *dmax* parameter is how far any atom can move in a single line search in any dimension (x, y, or z). For the *quickmin* and *fire* minimization styles, the *dmax* setting is how far any atom can move in a single iteration (timestep). Thus a value of 0.1 in real *units* means no atom will move further than 0.1 Angstroms in a single outer iteration. This prevents highly overlapped atoms from being moved long distances (e.g. through another atom) due to large forces.

The choice of line search algorithm for the *cg* and *sd* minimization styles can be selected via the *line* keyword. The default *quadratic* line search algorithm starts out using the robust backtracking method described below. However, once the system gets close to a local minimum and the linesearch steps get small, so that the energy is approximately quadratic in the step length, it uses the estimated location of zero gradient as the linesearch step, provided the energy change is downhill. This becomes more efficient than backtracking for highly-converged relaxations. The *forcezero* line search algorithm is similar to *quadratic*. It may be more efficient than *quadratic* on some systems.

The backtracking search is robust and should always find a local energy minimum. However, it will “converge” when it can no longer reduce the energy of the system. Individual atom forces may still be larger than desired at this point, because the energy change is measured as the difference of two large values (energy before and energy after) and that difference may be smaller than machine epsilon even if atoms could move in the gradient direction to reduce forces further.

The choice of a norm can be modified for the min styles *cg*, *sd*, *quickmin*, *fire*, *fire/old*, *spin*, *spin/cg* and *spin/lbfgs* using the *norm* keyword. The default *two* norm computes the 2-norm (Euclidean length) of the global force vector:

$$\|\vec{F}\|_2 = \sqrt{\vec{F}_1^2 + \dots + \vec{F}_N^2}$$

The *max* norm computes the length of the 3-vector force for each atom (2-norm), and takes the maximum value of those across all atoms

$$\|\vec{F}\|_{max} = \max\left(\|\vec{F}_1\|, \dots, \|\vec{F}_N\|\right)$$

The *inf* norm takes the maximum component across the forces of all atoms in the system:

$$\|\vec{F}\|_{inf} = \max(|F_1^1|, |F_1^2|, |F_1^3| \dots, |F_N^1|, |F_N^2|, |F_N^3|)$$

For the min styles *spin*, *spin/cg* and *spin/lbfgs*, the force norm is replaced by the spin-torque norm.

Keywords *alpha\_damp* and *discrete\_factor* only make sense when a *min\_spin* command is declared. Keyword *alpha\_damp* defines an analog of a magnetic Gilbert damping. It defines a relaxation rate toward an equilibrium for a given magnetic system. Keyword *discrete\_factor* defines a discretization factor for the adaptive timestep used in the *spin* minimization. See *min\_spin* for more information about those quantities.

The choice of a line search algorithm for the *spin/cg* and *spin/lbfgs* styles can be specified via the *line* keyword. The *spin\_cubic* and *spin\_none* keywords only make sense when one of those two minimization styles is declared. The *spin\_cubic* performs the line search based on a cubic interpolation of the energy along the search direction. The *spin\_none* keyword deactivates the line search procedure. The *spin\_none* is a default value for *line* keyword for both *spin/lbfgs* and *spin/cg*. Convergence of *spin/lbfgs* can be more robust if *spin\_cubic* line search is used.

The Newton *integrator* used for *fire* minimization can be selected to be either the symplectic Euler (*eulerimplicit*) or velocity Verlet (*verlet*). *tmax* defines the maximum value for the adaptive timestep during a *fire* minimization. It is a multiplication factor applied to the current *timestep* (not in time unit). For example, *tmax* = 4.0 with a *timestep* of 2fs, means that the maximum value the timestep can reach during a *fire* minimization is 4fs. Note that parameter defaults has been chosen to be reliable in most cases, but one should consider adjusting *timestep* and *tmax* to optimize the minimization for large or complex systems. Other parameters of the *fire* minimization can be tuned (*tmin*, *delaystep*, *dtgrow*, *dtshrink*, *alpha0*, and *alphashrink*). Please refer to the references describing the *min\_style fire*. An additional stopping criteria *vdffmax* is used by *fire* in order to avoid unnecessary looping when it is reasonable to think the system will not be relaxed further. Note that in this case the system will NOT have reached your minimization criteria. This could happen when the system comes to be stuck in a local basin of the phase space. *vdffmax* is the maximum number of consecutive iterations with  $P(t) < 0$ .

The *min\_style fire* is an optimized implementation of *min\_style fire/old*. It can however behave similarly to the *fire/old* style by using the following set of parameters:

```
min_modify integrator eulerexplicit tmax 10.0 tmin 0.0 delaystep 5 &
           dtgrow 1.1 dtshrink 0.5 alpha0 0.1 alphashrink 0.99 &
           vdffmax 100000 halfstepback no initialdelay no
```

### 16.73.4 Restrictions

For magnetic GNEB calculations, only *spin\_none* value for *line* keyword can be used when minimization styles *spin/cg* and *spin/lbfgs* are employed. See *neb/spin* for more explanation.

### 16.73.5 Related commands

*min\_style*, *minimize*

### 16.73.6 Default

The option defaults are *dmax* = 0.1, *line* = quadratic and *norm* = two.

For the *spin*, *spin/cg* and *spin/lbfgs* styles, the option defaults are *alpha\_damp* = 1.0, *discrete\_factor* = 10.0, *line* = *spin\_none*, and *norm* = euclidean.

For the *fire* style, the option defaults are *integrator* = eulerimplicit, *tmax* = 10.0, *tmin* = 0.02, *delaystep* = 20, *dtgrow* = 1.1, *dtshrink* = 0.5, *alpha0* = 0.25, *alphashrink* = 0.99, *vdffmax* = 2000, *halfstepback* = yes and *initialdelay* = yes.

## 16.74 min\_style spin command

## 16.75 min\_style spin/cg command

## 16.76 min\_style spin/lbfgs command

### 16.76.1 Syntax

```
min_style spin
min_style spin/cg
min_style spin/lbfgs
```

### 16.76.2 Examples

```
min_style spin/lbfgs
min_modify line spin_cubic discrete_factor 10.0
```

### 16.76.3 Description

Apply a minimization algorithm to use when a *minimize* command is performed.

Style *spin* defines a damped spin dynamics with an adaptive timestep, according to:

$$\frac{d\vec{s}_i}{dt} = \lambda \vec{s}_i \times (\vec{\omega}_i \times \vec{s}_i)$$

with  $\lambda$  a damping coefficient (similar to a Gilbert damping).  $\lambda$  can be defined by setting the *alpha\_damp* keyword with the *min\_modify* command.

The minimization procedure solves this equation using an adaptive timestep. The value of this timestep is defined by the largest precession frequency that has to be solved in the system:

$$\Delta t_{\max} = \frac{2\pi}{\kappa |\vec{\omega}_{\max}|}$$

with  $|\vec{\omega}_{\max}|$  the norm of the largest precession frequency in the system (across all processes, and across all replicas if a spin/neb calculation is performed).

$\kappa$  defines a discretization factor *discrete\_factor* for the definition of this timestep. *discrete\_factor* can be defined with the *min\_modify* command.

Style *spin/cg* defines an orthogonal spin optimization (OSO) combined to a conjugate gradient (CG) algorithm. The *min\_modify* command can be used to couple the *spin/cg* to a line search procedure, and to modify the discretization factor *discrete\_factor*. By default, style *spin/cg* does not employ the line search procedure and uses the adaptive timestep technique in the same way as style *spin*.

Style *spin/lbfgs* defines an orthogonal spin optimization (OSO) combined to a limited-memory Broyden-Fletcher-Goldfarb-Shanno (L-BFGS) algorithm. By default, style *spin/lbfgs* does not employ line search procedure. If the line search procedure is not used then the discrete factor defines the maximum root mean squared rotation angle of spins by equation  $\pi/(5*Kappa)$ . The default value for Kappa is 10. The *spin\_cubic* line search option can improve the convergence of the *spin/lbfgs* algorithm.

The *min\_modify* command can be used to activate the line search procedure, and to modify the discretization factor *discrete\_factor*.

For more information about styles *spin/cg* and *spin/lbfgs*, see their implementation reported in (*Ivanov*).

---

**Note:** All the *spin* styles replace the force tolerance by a torque tolerance. See *minimize* for more explanation.

---

---

**Note:** The *spin/cg* and *spin/lbfgs* styles can be used for magnetic NEB calculations only if the line search procedure is deactivated. See *neb/spin* for more explanation.

---

## 16.76.4 Restrictions

The *spin*, *spin/cg*, and *spin/lbfgs* styles are part of the SPIN package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This minimization procedure is only applied to spin degrees of freedom for a frozen lattice configuration.

## 16.76.5 Related commands

*min\_style*, *minimize*, *min\_modify*

### 16.76.6 Default

The option defaults are *alpha\_damp* = 1.0, *discrete\_factor* = 10.0, *line* = *spin\_none* and *norm* = *euclidean*.

---

(Ivanov) Ivanov, Uzdin, Jonsson. arXiv preprint arXiv:1904.02669, (2019).

### 16.77 `min_style cg` command

### 16.78 `min_style hftn` command

### 16.79 `min_style sd` command

### 16.80 `min_style quickmin` command

### 16.81 `min_style fire` command

### 16.82 `min_style fire/old` command

### 16.83 `min_style spin` command

### 16.84 `min_style spin/cg` command

### 16.85 `min_style spin/lbfgs` command

#### 16.85.1 Syntax

```
min_style style
```

- style = *cg* or *hftn* or *sd* or *quickmin* or *fire* or *fire/old* or *spin* or *spin/cg* or *spin/lbfgs*  
*spin* is discussed briefly here and fully on [min\\_style spin](#) doc page  
*spin/cg* is discussed briefly here and fully on [min\\_style spin](#) doc page  
*spin/lbfgs* is discussed briefly here and fully on [min\\_style spin](#) doc page

#### 16.85.2 Examples

```
min_style cg  
min_style fire  
min_style spin
```



### 16.85.3 Description

Choose a minimization algorithm to use when a *minimize* command is performed.

Style *cg* is the Polak-Ribiere version of the conjugate gradient (CG) algorithm. At each iteration the force gradient is combined with the previous iteration information to compute a new search direction perpendicular (conjugate) to the previous search direction. The PR variant affects how the direction is chosen and how the CG method is restarted when it ceases to make progress. The PR variant is thought to be the most effective CG choice for most problems.

Style *hfn* is a Hessian-free truncated Newton algorithm. At each iteration a quadratic model of the energy potential is solved by a conjugate gradient inner iteration. The Hessian (second derivatives) of the energy is not formed directly, but approximated in each conjugate search direction by a finite difference directional derivative. When close to an energy minimum, the algorithm behaves like a Newton method and exhibits a quadratic convergence rate to high accuracy. In most cases the behavior of *hfn* is similar to *cg*, but it offers an alternative if *cg* seems to perform poorly. This style is not affected by the *min\_modify* command.

Style *sd* is a steepest descent algorithm. At each iteration, the search direction is set to the downhill direction corresponding to the force vector (negative gradient of energy). Typically, steepest descent will not converge as quickly as CG, but may be more robust in some situations.

Style *quickmin* is a damped dynamics method described in (*Sheppard*), where the damping parameter is related to the projection of the velocity vector along the current force vector for each atom. The velocity of each atom is initialized to 0.0 by this style, at the beginning of a minimization.

Style *fire* is a damped dynamics method described in (*Bitzek*), which is similar to *quickmin* but adds a variable timestep and alters the projection operation to maintain components of the velocity non-parallel to the current force vector. The velocity of each atom is initialized to 0.0 by this style, at the beginning of a minimization. This style correspond to an optimized version described in (*Guenole*) that include different time integration schemes and defaults parameters. The default parameters can be modified with the command *min\_modify*.

Style *fire/old* is the original implementation of *fire* in Lammmps, conserved for backward compatibility. The main differences regarding the current version *fire* are: time integration by Explicit Euler only, different sequence in maintaining velocity components non-parallel to the current force vector and hard-coded minimization parameters. A complete description of the differences between *fire/old* and *fire* can be found in (*Guenole*) (where the current *fire* in LAMMPS is called *fire2.0*). By using an appropriate set of parameters, *fire* can behave similar to *fire/old*, as described in the *min\_modify* command.

Style *spin* is a damped spin dynamics with an adaptive timestep.

Style *spin/cg* uses an orthogonal spin optimization (OSO) combined to a conjugate gradient (CG) approach to minimize spin configurations.

Style *spin/lbfgs* uses an orthogonal spin optimization (OSO) combined to a limited-memory Broyden-Fletcher-Goldfarb-Shanno (LBFGS) approach to minimize spin configurations.

See the *min/spin* page for more information about the *spin*, *spin/cg* and *spin/lbfgs* styles.

Either the *quickmin*, *fire* and *fire/old* styles are useful in the context of nudged elastic band (NEB) calculations via the *neb* command.

Either the *spin*, *spin/cg* and *spin/lbfgs* styles are useful in the context of magnetic geodesic nudged elastic band (GNEB) calculations via the *neb/spin* command.

---

**Note:** The damped dynamic minimizers use whatever timestep you have defined via the *timestep* command. Often they will converge more quickly if you use a timestep about 10x larger than you would normally use for dynamics simulations. For *fire*, the default timestep is recommended to be equal to the one you would normally use for dynamics simulations.

---

---

**Note:** The *quickmin*, *fire*, *fire/old*, *hftn*, and *cg/kk* styles do not yet support the use of the *fix box/relax* command or minimizations involving the electron radius in *eFF* models.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 16.85.4 Restrictions

The *spin*, *spin/cg*, and *spin/lbfgs* styles are part of the SPIN package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 16.85.5 Related commands

*min\_modify*, *minimize*, *neb*

## 16.85.6 Default

```
min_style cg
```

---

**(Sheppard)** Sheppard, Terrell, Henkelman, J Chem Phys, 128, 134106 (2008). See ref 1 in this paper for original reference to Qmin in Jonsson, Mills, Jacobsen.

**(Bitzek)** Bitzek, Koskinen, Gahler, Moseler, Gumbusch, Phys Rev Lett, 97, 170201 (2006).

**(Guenole)** Guenole, Noehring, Vaid, Houille, Xie, Prakash, Bitzek, Comput Mater Sci, 175, 109584 (2020).

## 16.86 minimize command

## 16.87 minimize/kk command

### 16.87.1 Syntax

```
minimize etol ftol maxiter maxeval
```

---

- etol = stopping tolerance for energy (unitless)
- ftol = stopping tolerance for force (force units)
- maxiter = max iterations of minimizer
- maxeval = max number of force/energy evaluations

## 16.87.2 Examples

```
minimize 1.0e-4 1.0e-6 100 1000
minimize 0.0 1.0e-8 1000 100000
```

## 16.87.3 Description

Perform an energy minimization of the system, by iteratively adjusting atom coordinates. Iterations are terminated when one of the stopping criteria is satisfied. At that point the configuration will hopefully be in a local potential energy minimum. More precisely, the configuration should approximate a critical point for the objective function (see below), which may or may not be a local minimum.

The minimization algorithm used is set by the *min\_style* command. Other options are set by the *min\_modify* command. Minimize commands can be interspersed with *run* commands to alternate between relaxation and dynamics. The minimizers bound the distance atoms may move in one iteration, so that you can relax systems with highly overlapped atoms (large energies and forces) by pushing the atoms off of each other.

---

### Neighbor list update settings

The distance that atoms can move during individual minimization steps can be quite large, especially at the beginning of a minimization. Thus *neighbor list settings* *<neigh\_modify>* of *every = 1* and *delay = 0* are **required**. This may be combined with either *check = no* (always update the neighbor list) or *check = yes* (only update the neighbor list if at least one atom has moved more than half the *neighbor list skin* *<neighbor>* distance since the last reneighboring). Using *check = yes* is recommended since it avoids unneeded reneighboring steps when the system is closer to the minimum and thus atoms move only small distances. Using *check = no* may be required for debugging or when coupling LAMMPS with external codes that require a predictable sequence of neighbor list updates.

If the settings are **not** *every = 1* and *delay = 0*, LAMMPS will temporarily apply a *neigh\_modify every 1 delay 0 check yes <neigh\_modify>* setting during the minimization and restore the original setting at the end of the minimization. A corresponding message will be printed to the screen and log file, if this happens.

---

Alternate means of relaxing a system are to run dynamics with a small or *limited timestep*. Or dynamics can be run using *fix viscous* to impose a damping force that slowly drains all kinetic energy from the system. The *pair\_style soft* potential can be used to un-overlap atoms while running dynamics.

Note that you can minimize some atoms in the system while holding the coordinates of other atoms fixed by applying *fix setforce 0.0 0.0 0.0* to the other atoms. See a more detailed discussion of *using fixes while minimizing below*.

The *minimization styles* *cg*, *sd*, and *hftn* involves an outer iteration loop which sets the search direction along which atom coordinates are changed. An inner iteration is then performed using a line search algorithm. The line search typically evaluates forces and energies several times to set new coordinates. Currently, a backtracking algorithm is used which may not be optimal in terms of the number of force evaluations performed, but appears to be more robust than previous line searches we have tried. The backtracking method is described in Nocedal and Wright's Numerical Optimization (Procedure 3.1 on p 41).

The *minimization styles* *quickmin*, *fire* and *fire/old* perform damped dynamics using an Euler integration step. Thus they require a *timestep* be defined.

**Note:** The damped dynamic minimizer algorithms will use the timestep you have defined via the *timestep* command or its default value. Often they will converge more quickly if you use a timestep about 10x larger than you would normally use for regular molecular dynamics simulations.

---

In all cases, the objective function being minimized is the total potential energy of the system as a function of the N atom coordinates:

$$E(r_1, r_2, \dots, r_N) = \sum_{i,j} E_{pair}(r_i, r_j) + \sum_{ij} E_{bond}(r_i, r_j) + \sum_{ijk} E_{angle}(r_i, r_j, r_k) + \sum_{ijkl} E_{dihedral}(r_i, r_j, r_k, r_l) + \sum_{ijkl} E_{improper}(r_i, r_j, r_k, r_l) + \sum_i E_{fix}(r_i)$$

where the first term is the sum of all non-bonded *pairwise interactions* including *long-range Coulombic interactions*, the second through fifth terms are *bond*, *angle*, *dihedral*, and *improper* interactions respectively, and the last term is energy due to *fixes* which can act as constraints or apply force to atoms, such as through interaction with a wall. See the discussion below about how fix commands affect minimization.

The starting point for the minimization is the current configuration of the atoms.

---

The minimization procedure stops if any of several criteria are met:

- the change in energy between outer iterations is less than *etol*
- the 2-norm (length) of the global force vector is less than the *ftol*
- the line search fails because the step distance backtracks to 0.0
- the number of outer iterations or timesteps exceeds *maxiter*
- the number of total force evaluations exceeds *maxeval*

---

**Note:** the *minimization style spin*, *spin/cg*, and *spin/lbfgs* replace the force tolerance *ftol* by a torque tolerance. The minimization procedure stops if the 2-norm (length) of the torque vector on atom (defined as the cross product between the atomic spin and its precession vectors omega) is less than *ftol*, or if any of the other criteria are met. Torque have the same units as the energy.

---

**Note:** You can also use the *fix halt* command to specify a general criterion for exiting a minimization, that is a calculation performed on the state of the current system, as defined by an *equal-style variable*.

---

For the first criterion, the specified energy tolerance *etol* is unitless; it is met when the energy change between successive iterations divided by the energy magnitude is less than or equal to the tolerance. For example, a setting of 1.0e-4 for *etol* means an energy tolerance of one part in 10<sup>4</sup>. For the damped dynamics minimizers this check is not performed for a few steps after velocities are reset to 0, otherwise the minimizer would prematurely converge.

For the second criterion, the specified force tolerance *ftol* is in force units, since it is the length of the global force vector for all atoms, e.g. a vector of size 3N for N atoms. Since many of the components will be near zero after minimization, you can think of *ftol* as an upper bound on the final force on any component of any atom. For example, a setting of 1.0e-4 for *ftol* means no x, y, or z component of force on any atom will be larger than 1.0e-4 (in force units) after minimization.

Either or both of the *etol* and *ftol* values can be set to 0.0, in which case some other criterion will terminate the minimization.

During a minimization, the outer iteration count is treated as a timestep. Output is triggered by this timestep, e.g. thermodynamic output or dump and restart files.

Using the *thermo\_style custom* command with the *fmax* or *fnorm* keywords can be useful for monitoring the progress of the minimization. Note that these outputs will be calculated only from forces on the atoms, and will not include any extra degrees of freedom, such as from the *fix box/relax* command.

Following minimization, a statistical summary is printed that lists which convergence criterion caused the minimizer to stop, as well as information about the energy, force, final line search, and iteration counts. An example is as follows:

```
Minimization stats:
  Stopping criterion = max iterations
  Energy initial, next-to-last, final =
    -0.626828169302    -2.82642039062    -2.82643549739
  Force two-norm initial, final = 2052.1 91.9642
  Force max component initial, final = 346.048 9.78056
  Final line search alpha, max atom move = 2.23899e-06 2.18986e-05
  Iterations, force evaluations = 2000 12724
```

The 3 energy values are for before and after the minimization and on the next-to-last iteration. This is what the *etol* parameter checks.

The two-norm force values are the length of the global force vector before and after minimization. This is what the *ftol* parameter checks.

The max-component force values are the absolute value of the largest component (x,y,z) in the global force vector, i.e. the infinity-norm of the force vector.

The alpha parameter for the line-search, when multiplied by the max force component (on the last iteration), gives the max distance any atom moved during the last iteration. Alpha will be 0.0 if the line search could not reduce the energy. Even if alpha is non-zero, if the “max atom move” distance is tiny compared to typical atom coordinates, then it is possible the last iteration effectively caused no atom movement and thus the evaluated energy did not change and the minimizer terminated. Said another way, even with non-zero forces, it’s possible the effect of those forces is to move atoms a distance less than machine precision, so that the energy cannot be further reduced.

The iterations and force evaluation values are what is checked by the *maxiter* and *maxeval* parameters.

---

**Note:** There are several force fields in LAMMPS which have discontinuities or other approximations which may prevent you from performing an energy minimization to tight tolerances. For example, you should use a *pair\_style* that goes to 0.0 at the cutoff distance when performing minimization (even if you later change it when running dynamics). If you do not do this, the total energy of the system will have discontinuities when the relative distance between any pair of atoms changes from cutoff *plus* epsilon to cutoff *minus* epsilon and the minimizer may thus behave poorly. Some of the many-body potentials use splines and other internal cutoffs that inherently have this problem. The *long-range Coulombic styles* (PPPM, Ewald) are approximate to within the user-specified tolerance, which means their energy and forces may not agree to a higher precision than the Kspace-specified tolerance. This agreement is further reduced when using tabulation to speed up the computation of the real-space part of the Coulomb interactions, which is enabled by default. In all these cases, the minimizer may give up and stop before finding a minimum to the specified energy or force tolerance.

---

Note that a cutoff Lennard-Jones potential (and others) can be shifted so that its energy is 0.0 at the cutoff via the *pair\_modify* command. See the doc pages for individual *pair\_styles* for details. Note that most Coulombic potentials have a cutoff, unless versions with a long-range component are used (e.g. *pair\_style lj/cut/coul/long*) or some other damping/smoothing schemes are used. The CHARMM potentials go to 0.0 at the cutoff (e.g. *pair\_style lj/charmm/coul/charmm*), as do the GROMACS potentials (e.g. *pair\_style lj/gromacs*).

If a soft potential (*pair\_style soft*) is used the *Astop* value is used for the prefactor (no time dependence).

The *fix box/relax* command can be used to apply an external pressure to the simulation box and allow it to shrink/expand during the minimization.

Only a few other fixes (typically those that add forces) are invoked during minimization. See the doc pages for individual *fix* commands to see which ones are relevant. Current examples of fixes that can be used include:

- *fix addforce*
- *fix addtorque*
- *fix efield*
- *fix enforce2d*
- *fix indent*
- *fix lineforce*
- *fix planeforce*
- *fix setforce*
- *fix spring*
- *fix spring/self*
- *fix viscous*
- *fix wall*
- *fix wall/region*

---

**Note:** Some fixes which are invoked during minimization have an associated potential energy. For that energy to be included in the total potential energy of the system (the quantity being minimized), you **MUST** enable the *fix\_modify energy* option for that fix. The doc pages for individual *fix* commands specify if this should be done.

---

---

**Note:** The minimizers in LAMMPS do not allow for bonds (or angles, etc) to be held fixed while atom coordinates are being relaxed, e.g. via *fix shake* or *fix rigid*. See more info in the Restrictions section below.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 16.87.4 Restrictions

Features that are not yet implemented are listed here, in case someone knows how they could be coded:

It is an error to use *fix shake* with minimization because it turns off bonds that should be included in the potential energy of the system. The effect of a fix shake can be approximated during a minimization by using stiff spring constants for the bonds and/or angles that would normally be constrained by the SHAKE algorithm.

*Fix rigid* is also not supported by minimization. It is not an error to have it defined, but the energy minimization will not keep the defined body(s) rigid during the minimization. Note that if bonds, angles, etc internal to a rigid body have been turned off (e.g. via *neigh\_modify exclude*), they will not contribute to the potential energy which is probably not what is desired.

Pair potentials that produce torque on a particle (e.g. *granular potentials* or the *GayBerne potential* for ellipsoidal particles) are not relaxed by a minimization. More specifically, radial relaxations are induced, but no rotations are induced by a minimization, so such a system will not fully relax.

## 16.87.5 Related commands

*min\_modify*, *min\_style*, *run\_style*

## 16.87.6 Default

none

## 16.88 molecule command

### 16.88.1 Syntax

```
molecule ID file1 keyword values ... file2 keyword values ... fileN ...
```

- ID = user-assigned name for the molecule template
- file1,file2,... = names of files containing molecule descriptions
- zero or more keyword/value pairs may be appended after each file
- keyword = *offset* or *toff* or *boff* or *aoff* or *doff* or *ioff* or *scale*

```
offset values = Toff Boff Aoff Doft Ioff
```

```
  Toff = offset to add to atom types
```

```
  Boff = offset to add to bond types
```

```
  Aoff = offset to add to angle types
```

```
  Doft = offset to add to dihedral types
```

```
  Ioff = offset to add to improper types
```

```
toff value = Toff
```

```
  Toff = offset to add to atom types
```

```
boff value = Boff
```

```
  Boff = offset to add to bond types
```

```
aoff value = Aoff
```

```
  Aoff = offset to add to angle types
```

```
doff value = Doft
```

```
  Doft = offset to add to dihedral types
```

```
ioff value = Ioff
  Ioff = offset to add to improper types
scale value = sfactor
  sfactor = scale factor to apply to the size and mass of the molecule
```

## 16.88.2 Examples

```
molecule 1 mymol.txt
molecule 1 co2.txt h2o.txt
molecule C02 co2.txt boff 3 aoff 2
molecule 1 mymol.txt offset 6 9 18 23 14
molecule objects file.1 scale 1.5 file.1 scale 2.0 file.2 scale 1.3
```

## 16.88.3 Description

Define a molecule template that can be used as part of other LAMMPS commands, typically to define a collection of particles as a bonded molecule or a rigid body. Commands that currently use molecule templates include:

- *fix deposit*
- *fix pour*
- *fix rigid/small*
- *fix shake*
- *fix gcmc*
- *fix bond/react*
- *create\_atoms*
- *atom\_style template*

The ID of a molecule template can only contain alphanumeric characters and underscores.

A single template can contain multiple molecules, listed one per file. Some of the commands listed above currently use only the first molecule in the template, and will issue a warning if the template contains multiple molecules. The *atom\_style template* command allows multiple-molecule templates to define a system with more than one templated molecule.

Each filename can be followed by optional keywords which are applied only to the molecule in the file as used in this template. This is to make it easy to use the same molecule file in different molecule templates or in different simulations. You can specify the same file multiple times with different optional keywords.

The *offset*, *toff*, *boff*, *aoff*, *doff*, *ioff* keywords add the specified offset values to the atom types, bond types, angle types, dihedral types, and/or improper types as they are read from the molecule file. E.g. if *toff* = 2, and the file uses atom types 1,2,3, then each created molecule will have atom types 3,4,5. For the *offset* keyword, all five offset values must be specified, but individual values will be ignored if the molecule template does not use that attribute (e.g. no bonds).

---

**Note:** Offsets are **ignored** on lines using type labels, as the type labels will determine the actual types directly depending on the current *labelmap* settings.

---

The *scale* keyword scales the size of the molecule. This can be useful for modeling polydisperse granular rigid bodies. The scale factor is applied to each of these properties in the molecule file, if they are defined: the individual particle coordinates (Coords section), the individual mass of each particle (Masses section), the individual diameters of each



particle (Diameters section), the total mass of the molecule (header keyword = mass), the center-of-mass of the molecule (header keyword = com), and the moments of inertia of the molecule (header keyword = inertia).

---

**Note:** The molecule command can be used to define molecules with bonds, angles, dihedrals, impropers, or special bond lists of neighbors within a molecular topology, so that you can later add the molecules to your simulation, via one or more of the commands listed above. Since this topology-related information requires that suitable storage is reserved when LAMMPS creates the simulation box (e.g. when using the *create\_box* command or the *read\_data* command) suitable space has to be reserved so you do not overflow those pre-allocated data structures when adding molecules later. Both the *create\_box* command and the *read\_data* command have “extra” options which insure space is allocated for storing topology info for molecules that are added later.

---

The format of an individual molecule file is similar but (not identical) to the data file read by the *read\_data* commands, and is as follows.

A molecule file has a header and a body. The header appears first. The first line of the header and thus of the molecule file is *always* skipped; it typically contains a description of the file or a comment from the software that created the file.

Then lines are read one line at a time. Lines can have a trailing comment starting with ‘#’ that is ignored. There *must* be at least one blank between any valid content and the comment. If the line is blank (i.e. contains only white-space after comments are deleted), it is skipped. If the line contains a header keyword, the corresponding value(s) is/are read from the line. A line that is *not* blank and does *not* contain a header keyword begins the body of the file.

The body of the file contains zero or more sections. The first line of a section has only a keyword. The next line is skipped. The remaining lines of the section contain values. The number of lines depends on the section keyword as described below. Zero or more blank lines can be used between sections. Sections can appear in any order, with a few exceptions as noted below.

These are the recognized header keywords. Header lines can come in any order. The numeric value(s) are read from the beginning of the line. The keyword should appear at the end of the line. All these settings have default values, as explained below. A line need only appear if the value(s) are different than the default.

- N *atoms* = # of atoms N in molecule, default = 0
- Nb *bonds* = # of bonds Nb in molecule, default = 0
- Na *angles* = # of angles Na in molecule, default = 0
- Nd *dihedrals* = # of dihedrals Nd in molecule, default = 0
- Ni *impropers* = # of impropers Ni in molecule, default = 0
- Nf *fragments* = # of fragments in molecule, default = 0
- Mtotal *mass* = total mass of molecule
- Xc Yc Zc *com* = coordinates of center-of-mass of molecule
- Ixx Iyy Izz Ixy Ixz Iyz *inertia* = 6 components of inertia tensor of molecule

For *mass*, *com*, and *inertia*, the default is for LAMMPS to calculate this quantity itself if needed, assuming the molecule consists of a set of point particles or finite-size particles (with a non-zero diameter) that do not overlap. If finite-size particles in the molecule do overlap, LAMMPS will not account for the overlap effects when calculating any of these 3 quantities, so you should pre-compute them yourself and list the values in the file.

The mass and center-of-mass coordinates (Xc,Yc,Zc) are self-explanatory. The 6 moments of inertia (ixx,iyy,izz,ixy,ixz,iyz) should be the values consistent with the current orientation of the rigid body around its center of mass. The values are with respect to the simulation box XYZ axes, not with respect to the principal axes of the rigid body itself. LAMMPS performs the latter calculation internally.

These are the allowed section keywords for the body of the file.

- *Coords, Types, Molecules, Fragments, Charges, Diameters, Masses* = atom-property sections
- *Bonds, Angles, Dihedrals, Improvers* = molecular topology sections
- *Special Bond Counts, Special Bonds* = special neighbor info
- *Shake Flags, Shake Atoms, Shake Bond Types* = SHAKE info

For the Types, Bonds, Angles, Dihedrals, and Improvers sections, each atom/bond/angle/etc type can be specified either as a number (numeric type) or as an alphanumeric type label. The latter is only allowed if type labels have been defined, either by the *labelmap* command or in data files read by the *read\_data* command which have sections for Atom Type Labels, Bond Type Labels, Angle Type Labels, etc. See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used. When using type labels, any values specified as *offset* are ignored.

If a Bonds section is specified then the Special Bond Counts and Special Bonds sections can also be used, if desired, to explicitly list the 1-2, 1-3, 1-4 neighbors within the molecule topology (see details below). This is optional since if these sections are not included, LAMMPS will auto-generate this information. Note that LAMMPS uses this info to properly exclude or weight bonded pairwise interactions between bonded atoms. See the *special\_bonds* command for more details. One reason to list the special bond info explicitly is for the *thermalized Drude oscillator model* which treats the bonds between nuclear cores and Drude electrons in a different manner.

---

**Note:** Whether a section is required depends on how the molecule template is used by other LAMMPS commands. For example, to add a molecule via the *fix deposit* command, the Coords and Types sections are required. To add a rigid body via the *fix pour* command, the Bonds (Angles, etc) sections are not required, since the molecule will be treated as a rigid body. Some sections are optional. For example, the *fix pour* command can be used to add “molecules” which are clusters of finite-size granular particles. If the Diameters section is not specified, each particle in the molecule will have a default diameter of 1.0. See the doc pages for LAMMPS commands that use molecule templates for more details.

---

Each section is listed below in alphabetic order. The format of each section is described including the number of lines it must contain and rules (if any) for whether it can appear in the data file. For per-atom sections, entries should be numbered from 1 to Natoms (where Natoms is the number of atoms in the template), indicating which atom (or bond, etc) the entry applies to. Per-atom sections need to include a setting for every atom, but the atoms can be listed in any order.

---

*Coords* section:

- one line per atom
  - line syntax: ID x y z
  - x,y,z = coordinate of atom
- 

*Types* section:

- one line per atom
  - line syntax: ID type
  - type = atom type of atom (1-Natomtype, or type label)
- 

*Molecules* section:

- one line per atom
-

- line syntax: ID molecule-ID
  - molecule-ID = molecule ID of atom
- 

*Fragments* section:

- one line per fragment
- line syntax: ID a b c d ...
- a,b,c,d,... = IDs of atoms in fragment

The ID of a fragment can only contain alphanumeric characters and underscores. The atom IDs should be values from 1 to Natoms, where Natoms = # of atoms in the molecule.

---

*Charges* section:

- one line per atom
- line syntax: ID q
- q = charge on atom

This section is only allowed for *atom styles* that support charge. If this section is not included, the default charge on each atom in the molecule is 0.0.

---

*Diameters* section:

- one line per atom
- line syntax: ID diam
- diam = diameter of atom

This section is only allowed for *atom styles* that support finite-size spherical particles, e.g. atom\_style sphere. If not listed, the default diameter of each atom in the molecule is 1.0.

---

*Masses* section:

- one line per atom
- line syntax: ID mass
- mass = mass of atom

This section is only allowed for *atom styles* that support per-atom mass, as opposed to per-type mass. See the *mass* command for details. If this section is not included, the default mass for each atom is derived from its volume (see *Diameters* section) and a default density of 1.0, in *units* of mass/volume.

---

*Bonds* section:

- one line per bond
- line syntax: ID type atom1 atom2
- type = bond type (1-Nbondtype, or type label)
- atom1,atom2 = IDs of atoms in bond

The IDs for the two atoms in each bond should be values from 1 to Natoms, where Natoms = # of atoms in the molecule.

---

*Angles* section:

- one line per angle
- line syntax: ID type atom1 atom2 atom3
- type = angle type (1-Nangletype, or type label)
- atom1,atom2,atom3 = IDs of atoms in angle

The IDs for the three atoms in each angle should be values from 1 to Natoms, where Natoms = # of atoms in the molecule. The 3 atoms are ordered linearly within the angle. Thus the central atom (around which the angle is computed) is the atom2 in the list.

---

*Dihedrals* section:

- one line per dihedral
- line syntax: ID type atom1 atom2 atom3 atom4
- type = dihedral type (1-Ndihedralttype, or type label)
- atom1,atom2,atom3,atom4 = IDs of atoms in dihedral

The IDs for the four atoms in each dihedral should be values from 1 to Natoms, where Natoms = # of atoms in the molecule. The 4 atoms are ordered linearly within the dihedral.

---

*Impropers* section:

- one line per improper
- line syntax: ID type atom1 atom2 atom3 atom4
- type = improper type (1-Nimproptype, or type label)
- atom1,atom2,atom3,atom4 = IDs of atoms in improper

The IDs for the four atoms in each improper should be values from 1 to Natoms, where Natoms = # of atoms in the molecule. The ordering of the 4 atoms determines the definition of the improper angle used in the formula for the defined *improper style*. See the doc pages for individual styles for details.

---

*Special Bond Counts* section:

- one line per atom
- line syntax: ID N1 N2 N3
- N1 = # of 1-2 bonds
- N2 = # of 1-3 bonds
- N3 = # of 1-4 bonds

N1, N2, N3 are the number of 1-2, 1-3, 1-4 neighbors respectively of this atom within the topology of the molecule. See the [special\\_bonds](#) page for more discussion of 1-2, 1-3, 1-4 neighbors. If this section appears, the Special Bonds section must also appear.

As explained above, LAMMPS will auto-generate this information if this section is not specified. If specified, this section will override what would be auto-generated.

---

*Special Bonds* section:

- one line per atom
- line syntax: ID a b c d ...
- a,b,c,d,... = IDs of atoms in N1+N2+N3 special bonds

A, b, c, d, etc are the IDs of the n1+n2+n3 atoms that are 1-2, 1-3, 1-4 neighbors of this atom. The IDs should be values from 1 to Natoms, where Natoms = # of atoms in the molecule. The first N1 values should be the 1-2 neighbors, the next N2 should be the 1-3 neighbors, the last N3 should be the 1-4 neighbors. No atom ID should appear more than once. See the *special\_bonds* doc page for more discussion of 1-2, 1-3, 1-4 neighbors. If this section appears, the Special Bond Counts section must also appear.

As explained above, LAMMPS will auto-generate this information if this section is not specified. If specified, this section will override what would be auto-generated.

---

*Shake Flags* section:

- one line per atom
- line syntax: ID flag
- flag = 0,1,2,3,4

This section is only needed when molecules created using the template will be constrained by SHAKE via the “fix shake” command. The other two Shake sections must also appear in the file, following this one.

The meaning of the flag for each atom is as follows. See the *fix shake* page for a further description of SHAKE clusters.

- 0 = not part of a SHAKE cluster
  - 1 = part of a SHAKE angle cluster (two bonds and the angle they form)
  - 2 = part of a 2-atom SHAKE cluster with a single bond
  - 3 = part of a 3-atom SHAKE cluster with two bonds
  - 4 = part of a 4-atom SHAKE cluster with three bonds
- 

*Shake Atoms* section:

- one line per atom
- line syntax: ID a b c d
- a,b,c,d = IDs of atoms in cluster

This section is only needed when molecules created using the template will be constrained by SHAKE via the “fix shake” command. The other two Shake sections must also appear in the file.

The a,b,c,d values are atom IDs (from 1 to Natoms) for all the atoms in the SHAKE cluster that this atom belongs to. The number of values that must appear is determined by the shake flag for the atom (see the Shake Flags section above). All atoms in a particular cluster should list their a,b,c,d values identically.

If flag = 0, no a,b,c,d values are listed on the line, just the (ignored) ID.

If flag = 1, a,b,c are listed, where a = ID of central atom in the angle, and b,c the other two atoms in the angle.

If flag = 2, a,b are listed, where a = ID of atom in bond with the lowest ID, and b = ID of atom in bond with the highest ID.

If flag = 3, a,b,c are listed, where a = ID of central atom, and b,c = IDs of other two atoms bonded to the central atom.

If flag = 4, a,b,c,d are listed, where a = ID of central atom, and b,c,d = IDs of other three atoms bonded to the central atom.

See the *fix shake* page for a further description of SHAKE clusters.

---

*Shake Bond Types* section:

- one line per atom
- line syntax: ID a b c
- a,b,c = bond types (or angle type) of bonds (or angle) in cluster

This section is only needed when molecules created using the template will be constrained by SHAKE via the “fix shake” command. The other two Shake sections must also appear in the file.

The a,b,c values are bond types for all bonds in the SHAKE cluster that this atom belongs to. Bond types may be either numbers (from 1 to Nbondtypes) or bond type labels as defined by the *labelmap* command or a “Bond Type Labels” section of a data file.

The number of values that must appear is determined by the shake flag for the atom (see the Shake Flags section above). All atoms in a particular cluster should list their a,b,c values identically.

If flag = 0, no a,b,c values are listed on the line, just the (ignored) ID.

If flag = 1, a,b,c are listed, where a = bondtype of the bond between the central atom and the first non-central atom (value b in the Shake Atoms section), b = bondtype of the bond between the central atom and the second non-central atom (value c in the Shake Atoms section), and c = the angle type (1 to Nangletypes, or angle type label) of the angle between the 3 atoms.

If flag = 2, only a is listed, where a = bondtype of the bond between the 2 atoms in the cluster.

If flag = 3, a,b are listed, where a = bondtype of the bond between the central atom and the first non-central atom (value b in the Shake Atoms section), and b = bondtype of the bond between the central atom and the second non-central atom (value c in the Shake Atoms section).

If flag = 4, a,b,c are listed, where a = bondtype of the bond between the central atom and the first non-central atom (value b in the Shake Atoms section), b = bondtype of the bond between the central atom and the second non-central atom (value c in the Shake Atoms section), and c = bondtype of the bond between the central atom and the third non-central atom (value d in the Shake Atoms section).

See the *fix shake* page for a further description of SHAKE clusters.

---

## 16.88.4 Restrictions

None

## 16.88.5 Related commands

*fix deposit, fix pour, fix gcmc*

## 16.88.6 Default

The default keywords values are offset 0 0 0 0 and scale = 1.0.

## 16.89 neb command

### 16.89.1 Syntax

```
neb etol ftol N1 N2 Nevery file-style arg keyword
```

- etol = stopping tolerance for energy (energy units)
- ftol = stopping tolerance for force (force units)
- N1 = max # of iterations (timesteps) to run initial NEB
- N2 = max # of iterations (timesteps) to run barrier-climbing NEB
- Nevery = print replica energies and reaction coordinates every this many timesteps
- file-style = *final* or *each* or *none*

*final* arg = filename

filename = file with initial coords for final replica  
 coords for intermediate replicas are linearly interpolated  
 between first and last replica

*each* arg = filename

filename = unique filename for each replica (except first)  
 with its initial coords

*none* arg = no argument all replicas assumed to already have  
 their initial coords

keyword = *verbose*

### 16.89.2 Examples

```
neb 0.1 0.0 1000 500 50 final coords.final
neb 0.0 0.001 1000 500 50 each coords.initial.$i
neb 0.0 0.001 1000 500 50 none verbose
```

### 16.89.3 Description

Perform a nudged elastic band (NEB) calculation using multiple replicas of a system. Two or more replicas must be used; the first and last are the end points of the transition path.

NEB is a method for finding both the atomic configurations and height of the energy barrier associated with a transition state, e.g. for an atom to perform a diffusive hop from one energy basin to another in a coordinated fashion with its neighbors. The implementation in LAMMPS follows the discussion in these 4 papers: (*HenkelmanA*), (*HenkelmanB*), (*Nakano*) and (*Maras*).

Each replica runs on a partition of one or more processors. Processor partitions are defined at run-time using the *-partition command-line switch*. Note that if you have MPI installed, you can run a multi-replica simulation with more replicas (partitions) than you have physical processors, e.g you can run a 10-replica simulation on just one or two processors. You will simply not get the performance speed-up you would see with one or more physical processors per replica. See the *Howto replica* doc page for further discussion.

---

**Note:** As explained below, a NEB calculation performs a damped dynamics minimization across all the replicas. The minimizer uses whatever timestep you have defined in your input script, via the *timestep* command. Often NEB will converge more quickly if you use a timestep about 10x larger than you would normally use for dynamics simulations.

---

When a NEB calculation is performed, it is assumed that each replica is running the same system, though LAMMPS does not check for this. I.e. the simulation domain, the number of atoms, the interaction potentials, and the starting configuration when the *neb* command is issued should be the same for every replica.

In a NEB calculation each replica is connected to other replicas by inter-replica nudging forces. These forces are imposed by the *fix neb* command, which must be used in conjunction with the *neb* command. The group used to define the *fix neb* command defines the NEB atoms which are the only ones that inter-replica springs are applied to. If the group does not include all atoms, then non-NEB atoms have no inter-replica springs and the forces they feel and their motion is computed in the usual way due only to other atoms within their replica. Conceptually, the non-NEB atoms provide a background force field for the NEB atoms. They can be allowed to move during the NEB minimization procedure (which will typically induce different coordinates for non-NEB atoms in different replicas), or held fixed using other LAMMPS commands such as *fix setforce*. Note that the *partition* command can be used to invoke a command on a subset of the replicas, e.g. if you wish to hold NEB or non-NEB atoms fixed in only the end-point replicas.

The initial atomic configuration for each of the replicas can be specified in different manners via the *file-style* setting, as discussed below. Only atoms whose initial coordinates should differ from the current configuration need be specified.

Conceptually, the initial and final configurations for the first replica should be states on either side of an energy barrier.

As explained below, the initial configurations of intermediate replicas can be atomic coordinates interpolated in a linear fashion between the first and last replicas. This is often adequate for simple transitions. For more complex transitions, it may lead to slow convergence or even bad results if the minimum energy path (MEP, see below) of states over the barrier cannot be correctly converged to from such an initial path. In this case, you will want to generate initial states for the intermediate replicas that are geometrically closer to the MEP and read them in.

---

For a *file-style* setting of *final*, a filename is specified which contains atomic coordinates for zero or more atoms, in the format described below. For each atom that appears in the file, the new coordinates are assigned to that atom in the final replica. Each intermediate replica also assigns a new position to that atom in an interpolated manner. This is done by using the current position of the atom as the starting point and the read-in position as the final point. The distance between them is calculated, and the new position is assigned to be a fraction of the distance. E.g. if there are 10 replicas, the second replica will assign a position that is 10% of the distance along a line between the starting and final point, and the 9th replica will assign a position that is 90% of the distance along the line. Note that for this procedure to produce consistent coordinates across all the replicas, the current coordinates need to be the same in all replicas. LAMMPS does not check for this, but invalid initial configurations will likely result if it is not the case.



---

**Note:** The “distance” between the starting and final point is calculated in a minimum-image sense for a periodic simulation box. This means that if the two positions are on opposite sides of a box (periodic in that dimension), the distance between them will be small, because the periodic image of one of the atoms is close to the other. Similarly, even if the assigned position resulting from the interpolation is outside the periodic box, the atom will be wrapped back into the box when the NEB calculation begins.

---

For a *file-style* setting of *each*, a filename is specified which is assumed to be unique to each replica. This can be done by using a variable in the filename, e.g.

```
variable i equal part
neb 0.0 0.001 1000 500 50 each coords.initial.$i
```

which in this case will substitute the partition ID (0 to N-1) for the variable I, which is also effectively the replica ID. See the *variable* command for other options, such as using *world-*, *universe-*, or *uloop-style* variables.

Each replica (except the first replica) will read its file, formatted as described below, and for any atom that appears in the file, assign the specified coordinates to its atom. The various files do not need to contain the same set of atoms.

For a *file-style* setting of *none*, no filename is specified. Each replica is assumed to already be in its initial configuration at the time the *neb* command is issued. This allows each replica to define its own configuration by reading a replica-specific data or restart or dump file, via the *read\_data*, *read\_restart*, or *read\_dump* commands. The replica-specific names of these files can be specified as in the discussion above for the *each* *file-style*. Also see the section below for how a NEB calculation can produce restart files, so that a long calculation can be restarted if needed.

---

**Note:** None of the *file-style* settings change the initial configuration of any atom in the first replica. The first replica must thus be in the correct initial configuration at the time the *neb* command is issued.

---

A NEB calculation proceeds in two stages, each of which is a minimization procedure, performed via damped dynamics. To enable this, you must first define a damped dynamics *min\_style*, such as *quickmin* or *fire*. The *cg*, *sd*, and *hftn* styles cannot be used, since they perform iterative line searches in their inner loop, which cannot be easily synchronized across multiple replicas.

The minimizer tolerances for energy and force are set by *etol* and *ftol*, the same as for the *minimize* command.

A non-zero *etol* means that the NEB calculation will terminate if the energy criterion is met by every replica. The energies being compared to *etol* do not include any contribution from the inter-replica nudging forces, since these are non-conservative. A non-zero *ftol* means that the NEB calculation will terminate if the force criterion is met by every replica. The forces being compared to *ftol* include the inter-replica nudging forces.

The maximum number of iterations in each stage is set by *N1* and *N2*. These are effectively timestep counts since each iteration of damped dynamics is like a single timestep in a dynamics *run*. During both stages, the potential energy of each replica and its normalized distance along the reaction path (reaction coordinate RD) will be printed to the screen and log file every *Nevery* timesteps. The RD is 0 and 1 for the first and last replica. For intermediate replicas, it is the cumulative distance (normalized by the total cumulative distance) between adjacent replicas, where “distance” is defined as the length of the 3N-vector of differences in atomic coordinates, where N is the number of NEB atoms involved in the transition. These outputs allow you to monitor NEB’s progress in finding a good energy barrier. *N1* and *N2* must both be multiples of *Nevery*.

In the first stage of NEB, the set of replicas should converge toward a minimum energy path (MEP) of conformational states that transition over a barrier. The MEP for a transition is defined as a sequence of 3N-dimensional states, each of which has a potential energy gradient parallel to the MEP itself. The configuration of highest energy along a MEP corresponds to a saddle point. The replica states will also be roughly equally spaced along the MEP due to the inter-replica nudging force added by the *fix\_neb* command.

In the second stage of NEB, the replica with the highest energy is selected and the inter-replica forces on it are converted to a force that drives its atom coordinates to the top or saddle point of the barrier, via the barrier-climbing calculation described in (*HenkelmanB*). As before, the other replicas rearrange themselves along the MEP so as to be roughly equally spaced.

When both stages are complete, if the NEB calculation was successful, the configurations of the replicas should be along (close to) the MEP and the replica with the highest energy should be an atomic configuration at (close to) the saddle point of the transition. The potential energies for the set of replicas represents the energy profile of the transition along the MEP.

---

A few other settings in your input script are required or advised to perform a NEB calculation. See the NOTE about the choice of timestep at the beginning of this doc page.

An atom map must be defined which it is not by default for *atom\_style atomic* problems. The *atom\_modify map* command can be used to do this.

The minimizers in LAMMPS operate on all atoms in your system, even non-NEB atoms, as defined above. To prevent non-NEB atoms from moving during the minimization, you should use the *fix setforce* command to set the force on each of those atoms to 0.0. This is not required, and may not even be desired in some cases, but if those atoms move too far (e.g. because the initial state of your system was not well-minimized), it can cause problems for the NEB procedure.

The damped dynamics *minimizers*, such as *quickmin* and *fire*), adjust the position and velocity of the atoms via an Euler integration step. Thus you must define an appropriate *timestep* to use with NEB. As mentioned above, NEB will often converge more quickly if you use a timestep about 10x larger than you would normally use for dynamics simulations.

---

Each file read by the *neb* command containing atomic coordinates used to initialize one or more replicas must be formatted as follows.

The file can be ASCII text or a gzipped text file (detected by a *.gz* suffix). The file can contain initial blank lines or comment lines starting with “#” which are ignored. The first non-blank, non-comment line should list *N* = the number of lines to follow. The *N* successive lines contain the following information:

```
ID1 x1 y1 z1
ID2 x2 y2 z2
...
IDN xN yN zN
```

The fields are the atom ID, followed by the x,y,z coordinates. The lines can be listed in any order. Additional trailing information on the line is OK, such as a comment.

Note that for a typical NEB calculation you do not need to specify initial coordinates for very many atoms to produce differing starting and final replicas whose intermediate replicas will converge to the energy barrier. Typically only new coordinates for atoms geometrically near the barrier need be specified.

Also note there is no requirement that the atoms in the file correspond to the NEB atoms in the group defined by the *fix neb* command. Not every NEB atom need be in the file, and non-NEB atoms can be listed in the file.

---

Four kinds of output can be generated during a NEB calculation: energy barrier statistics, thermodynamic output by each replica, dump files, and restart files.

When running with multiple partitions (each of which is a replica in this case), the print-out to the screen and master log.lammps file contains a line of output, printed once every *Nevery* timesteps. It contains the timestep, the maximum force per replica, the maximum force per atom (in any replica), potential gradients in the initial, final, and climbing replicas, the forward and backward energy barriers, the total reaction coordinate (RDT), and the normalized reaction coordinate and potential energy of each replica.

The “maximum force per replica” is the two-norm of the  $3N$ -length force vector for the atoms in each replica, maximized across replicas, which is what the *ftol* setting is checking against. In this case,  $N$  is all the atoms in each replica. The “maximum force per atom” is the maximum force component of any atom in any replica. The potential gradients are the two-norm of the  $3N$ -length force vector solely due to the interaction potential i.e. without adding in inter-replica forces.

The “reaction coordinate” (RD) for each replica is the two-norm of the  $3N$ -length vector of distances between its atoms and the preceding replica’s atoms, added to the RD of the preceding replica. The RD of the first replica  $RD_1 = 0.0$ ; the RD of the final replica  $RD_N = RDT$ , the total reaction coordinate. The normalized RDs are divided by  $RDT$ , so that they form a monotonically increasing sequence from zero to one. When computing RD,  $N$  only includes the atoms being operated on by the *fix neb* command.

The forward (reverse) energy barrier is the potential energy of the highest replica minus the energy of the first (last) replica.

Supplementary information for all replicas can be printed out to the screen and master *log.lammps* file by adding the verbose keyword. This information include the following. The “path angle” (*pathangle*) for the replica  $i$  which is the angle between the  $3N$ -length vectors  $(R_{i-1} - R_i)$  and  $(R_{i+1} - R_i)$  (where  $R_i$  is the atomic coordinates of replica  $i$ ). A “path angle” of 180 indicates that replicas  $i-1$ ,  $i$  and  $i+1$  are aligned. “*angletangrad*” is the angle between the  $3N$ -length tangent vector and the  $3N$ -length force vector at image  $i$ . The tangent vector is calculated as in (*HenkelmanA*) for all intermediate replicas and at  $R_2 - R_1$  and  $R_M - R_{M-1}$  for the first and last replica, respectively. “*anglegrad*” is the angle between the  $3N$ -length energy gradient vector of replica  $i$  and that of replica  $i+1$ . It is not defined for the final replica and reads *nan*. *gradV* is the norm of the energy gradient of image  $i$ . *ReplicaForce* is the two-norm of the  $3N$ -length force vector (including nudging forces) for replica  $i$ . *MaxAtomForce* is the maximum force component of any atom in replica  $i$ .

When a NEB calculation does not converge properly, the supplementary information can help understanding what is going wrong. For instance when the path angle becomes acute, the definition of tangent used in the NEB calculation is questionable and the NEB cannot may diverge (*Maras*).

When running on multiple partitions, LAMMPS produces additional log files for each partition, e.g. *log.lammps.0*, *log.lammps.1*, etc. For a NEB calculation, these contain the thermodynamic output for each replica.

If *dump* commands in the input script define a filename that includes a *universe* or *uloop* style *variable*, then one dump file (per dump command) will be created for each replica. At the end of the NEB calculation, the final snapshot in each file will contain the sequence of snapshots that transition the system over the energy barrier. Earlier snapshots will show the convergence of the replicas to the MEP.

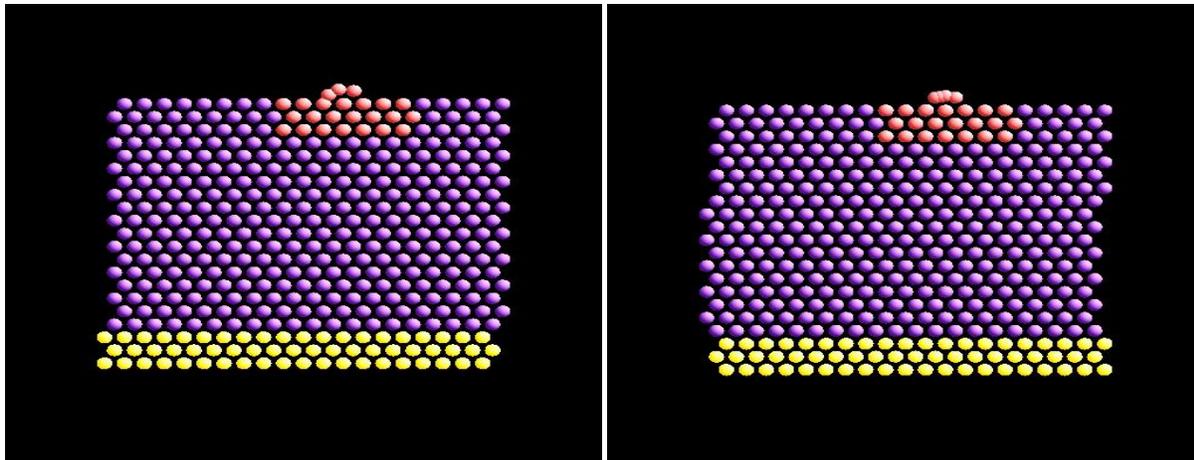
Likewise, *restart* filenames can be specified with a *universe* or *uloop* style *variable*, to generate restart files for each replica. These may be useful if the NEB calculation fails to converge properly to the MEP, and you wish to restart the calculation from an intermediate point with altered parameters.

There are 2 Python scripts provided in the *tools/python* directory, *neb\_combine.py* and *neb\_final.py*, which are useful in analyzing output from a NEB calculation. Assume a NEB simulation with  $M$  replicas, and the NEB atoms labeled with a specific atom type.

The *neb\_combine.py* script extracts atom coords for the NEB atoms from all  $M$  dump files and creates a single dump file where each snapshot contains the NEB atoms from all the replicas and one copy of non-NEB atoms from the first replica (presumed to be identical in other replicas). This can be visualized/animated to see how the NEB atoms relax as the NEB calculation proceeds.

The *neb\_final.py* script extracts the final snapshot from each of the  $M$  dump files to create a single dump file with  $M$  snapshots. This can be visualized to watch the system make its transition over the energy barrier.

To illustrate, here are images from the final snapshot produced by the *neb\_combine.py* script run on the dump files produced by the two example input scripts in *examples/neb*.



---

### 16.89.4 Restrictions

This command can only be used if LAMMPS was built with the REPLICIA package. See the *Build package* doc page for more info.

---

### 16.89.5 Related commands

*prd, temper, fix langevin, fix viscous*

### 16.89.6 Default

none

---

**(HenkelmanA)** Henkelman and Jonsson, J Chem Phys, 113, 9978-9985 (2000).

**(HenkelmanB)** Henkelman, Uberuaga, Jonsson, J Chem Phys, 113, 9901-9904 (2000).

**(Nakano)** Nakano, Comp Phys Comm, 178, 280-289 (2008).

**(Maras)** Maras, Trushin, Stukowski, Ala-Nissila, Jonsson, Comp Phys Comm, 205, 13-21 (2016)

## 16.90 neb/spin command

### 16.90.1 Syntax

```
neb/spin etol ttol N1 N2 Nevery file-style arg keyword
```

- etol = stopping tolerance for energy (energy units)
- ttol = stopping tolerance for torque ( units)
- N1 = max # of iterations (timesteps) to run initial NEB

- N2 = max # of iterations (timesteps) to run barrier-climbing NEB
- Nevery = print replica energies and reaction coordinates every this many timesteps
- file-style = *final* or *each* or *none*

```

final arg = filename
  filename = file with initial coords for final replica
  coords for intermediate replicas are linearly interpolated
  between first and last replica
each arg = filename
  filename = unique filename for each replica (except first)
  with its initial coords
none arg = no argument all replicas assumed to already have
  their initial coords

```

- keyword = *verbose*  
*verbose* = print supplemental information

## 16.90.2 Examples

```

neb/spin 0.1 0.0 1000 500 50 final coords.final
neb/spin 0.0 0.001 1000 500 50 each coords.initial.$i
neb/spin 0.0 0.001 1000 500 50 none verbose

```

## 16.90.3 Description

Perform a geodesic nudged elastic band (GNEB) calculation using multiple replicas of a system. Two or more replicas must be used; the first and last are the end points of the transition path.

GNEB is a method for finding both the spin configurations and height of the energy barrier associated with a transition state, e.g. spins to perform a collective rotation from one energy basin to another. The implementation in LAMMPS follows the discussion in the following paper: ([Bessaraba](#)).

Each replica runs on a partition of one or more processors. Processor partitions are defined at run-time using the *partition command-line switch*. Note that if you have MPI installed, you can run a multi-replica simulation with more replicas (partitions) than you have physical processors, e.g you can run a 10-replica simulation on just one or two processors. You will simply not get the performance speed-up you would see with one or more physical processors per replica. See the *Howto replica* doc page for further discussion.

---

**Note:** As explained below, a GNEB calculation performs a minimization across all the replicas. One of the *spin* style minimizers has to be defined in your input script.

---

When a GNEB calculation is performed, it is assumed that each replica is running the same system, though LAMMPS does not check for this. I.e. the simulation domain, the number of magnetic atoms, the interaction potentials, and the starting configuration when the neb command is issued should be the same for every replica.

In a GNEB calculation each replica is connected to other replicas by inter-replica nudging forces. These forces are imposed by the *fix neb/spin* command, which must be used in conjunction with the neb command. The group used to define the fix neb/spin command defines the GNEB magnetic atoms which are the only ones that inter-replica springs are applied to. If the group does not include all magnetic atoms, then non-GNEB magnetic atoms have no inter-replica springs and the torques they feel and their precession motion is computed in the usual way due only to other magnetic

atoms within their replica. Conceptually, the non-GNEB atoms provide a background force field for the GNEB atoms. Their magnetic spins can be allowed to evolve during the GNEB minimization procedure.

The initial spin configuration for each of the replicas can be specified in different manners via the *file-style* setting, as discussed below. Only atomic spins whose initial coordinates should differ from the current configuration need to be specified.

Conceptually, the initial and final configurations for the first replica should be states on either side of an energy barrier.

As explained below, the initial configurations of intermediate replicas can be spin coordinates interpolated in a linear fashion between the first and last replicas. This is often adequate for simple transitions. For more complex transitions, it may lead to slow convergence or even bad results if the minimum energy path (MEP, see below) of states over the barrier cannot be correctly converged to from such an initial path. In this case, you will want to generate initial states for the intermediate replicas that are geometrically closer to the MEP and read them in.

---

For a *file-style* setting of *final*, a filename is specified which contains atomic and spin coordinates for zero or more atoms, in the format described below. For each atom that appears in the file, the new coordinates are assigned to that atom in the final replica. Each intermediate replica also assigns a new spin to that atom in an interpolated manner. This is done by using the current direction of the spin at the starting point and the read-in direction as the final point. The “angular distance” between them is calculated, and the new direction is assigned to be a fraction of the angular distance.

---

**Note:** The “angular distance” between the starting and final point is evaluated in the geodesic sense, as described in (*BessarabA*).

---

**Note:** The angular interpolation between the starting and final point is achieved using Rodrigues formula:

---

$$\vec{m}_i^\nu = \vec{m}_i^I \cos(\omega_i^\nu) + (\vec{k}_i \times \vec{m}_i^I) \sin(\omega_i^\nu) + (1.0 - \cos(\omega_i^\nu)) \vec{k}_i (\vec{k}_i \cdot \vec{m}_i^I)$$

where  $\vec{m}_i^I$  is the initial spin configuration for spin  $i$ ,  $\omega_i^\nu$  is a rotation angle defined as:

$$\omega_i^\nu = (\nu - 1) \Delta\omega_i \quad \text{and} \quad \Delta\omega_i = \frac{\omega_i}{Q - 1}$$

with  $\nu$  the image number,  $Q$  the total number of images, and  $\omega_i$  the total rotation between the initial and final spins.  $\vec{k}_i$  defines a rotation axis such as:

$$\vec{k}_i = \frac{\vec{m}_i^I \times \vec{m}_i^F}{|\vec{m}_i^I \times \vec{m}_i^F|}$$

if the initial and final spins are not aligned. If the initial and final spins are aligned, then their cross product is null, and the expression above does not apply. If they point toward the same direction, the intermediate images conserve the same orientation. If the initial and final spins are aligned, but point toward opposite directions, an arbitrary rotation vector belonging to the plane perpendicular to initial and final spins is chosen. In this case, a warning message is displayed.

For a *file-style* setting of *each*, a filename is specified which is assumed to be unique to each replica. See the *neb* documentation page for more information about this option.

For a *file-style* setting of *none*, no filename is specified. Each replica is assumed to already be in its initial configuration at the time the neb command is issued. This allows each replica to define its own configuration by reading a replica-specific data or restart or dump file, via the *read\_data*, *read\_restart*, or *read\_dump* commands. The replica-specific names of these files can be specified as in the discussion above for the *each* file-style. Also see the section below for how a NEB calculation can produce restart files, so that a long calculation can be restarted if needed.

---

---

**Note:** None of the *file-style* settings change the initial configuration of any atom in the first replica. The first replica must thus be in the correct initial configuration at the time the *neb* command is issued.

---

A NEB calculation proceeds in two stages, each of which is a minimization procedure. To enable this, you must first define a *min\_style*, using either the *spin*, *spin/cg*, or *spin/lbfgs* style (see *min\_spin* for more information). The other styles cannot be used, since they relax the lattice degrees of freedom instead of the spins.

The minimizer tolerances for energy and force are set by *etol* and *ttol*, the same as for the *minimize* command.

A non-zero *etol* means that the GNEB calculation will terminate if the energy criterion is met by every replica. The energies being compared to *etol* do not include any contribution from the inter-replica nudging forces, since these are non-conservative. A non-zero *ttol* means that the GNEB calculation will terminate if the torque criterion is met by every replica. The torques being compared to *ttol* include the inter-replica nudging forces.

The maximum number of iterations in each stage is set by *N1* and *N2*. These are effectively timestep counts since each iteration of damped dynamics is like a single timestep in a dynamics *run*. During both stages, the potential energy of each replica and its normalized distance along the reaction path (reaction coordinate RD) will be printed to the screen and log file every *Nevery* timesteps. The RD is 0 and 1 for the first and last replica. For intermediate replicas, it is the cumulative angular distance (normalized by the total cumulative angular distance) between adjacent replicas, where “distance” is defined as the length of the 3N-vector of the geodesic distances in spin coordinates, with N the number of GNEB spins involved (see equation (13) in (*Bessaraba*)). These outputs allow you to monitor NEB’s progress in finding a good energy barrier. *N1* and *N2* must both be multiples of *Nevery*.

In the first stage of GNEB, the set of replicas should converge toward a minimum energy path (MEP) of conformational states that transition over a barrier. The MEP for a transition is defined as a sequence of 3N-dimensional spin states, each of which has a potential energy gradient parallel to the MEP itself. The configuration of highest energy along a MEP corresponds to a saddle point. The replica states will also be roughly equally spaced along the MEP due to the inter-replica nudging force added by the *fix neb* command.

In the second stage of GNEB, the replica with the highest energy is selected and the inter-replica forces on it are converted to a force that drives its spin coordinates to the top or saddle point of the barrier, via the barrier-climbing calculation described in (*Bessaraba*). As before, the other replicas rearrange themselves along the MEP so as to be roughly equally spaced.

When both stages are complete, if the GNEB calculation was successful, the configurations of the replicas should be along (close to) the MEP and the replica with the highest energy should be a spin configuration at (close to) the saddle point of the transition. The potential energies for the set of replicas represents the energy profile of the transition along the MEP.

---

An atom map must be defined which it is not by default for *atom\_style atomic* problems. The *atom\_modify map* command can be used to do this.

An initial value can be defined for the timestep. Although, the *spin* minimization algorithm is an adaptive timestep methodology, so that this timestep is likely to evolve during the calculation.

The minimizers in LAMMPS operate on all spins in your system, even non-GNEB atoms, as defined above.

---

Each file read by the *neb/spin* command containing spin coordinates used to initialize one or more replicas must be formatted as follows.

The file can be ASCII text or a gzipped text file (detected by a *.gz* suffix). The file can contain initial blank lines or comment lines starting with “#” which are ignored. The first non-blank, non-comment line should list N = the number of lines to follow. The N successive lines contain the following information:

```
ID1 g1 x1 y1 z1 sx1 sy1 sz1
ID2 g2 x2 y2 z2 sx2 sy2 sz2
...
IDN gN yN zN sxN syN szN
```

The fields are the atom ID, the norm of the associated magnetic spin, followed by the  $x,y,z$  coordinates and the  $s_x,s_y,s_z$  spin coordinates. The lines can be listed in any order. Additional trailing information on the line is OK, such as a comment.

Note that for a typical GNEB calculation you do not need to specify initial spin coordinates for very many atoms to produce differing starting and final replicas whose intermediate replicas will converge to the energy barrier. Typically only new spin coordinates for atoms geometrically near the barrier need be specified.

Also note there is no requirement that the atoms in the file correspond to the GNEB atoms in the group defined by the *fix neb* command. Not every GNEB atom need be in the file, and non-GNEB atoms can be listed in the file.

---

Four kinds of output can be generated during a GNEB calculation: energy barrier statistics, thermodynamic output by each replica, dump files, and restart files.

When running with multiple partitions (each of which is a replica in this case), the print-out to the screen and master log.lammps file contains a line of output, printed once every *Nevery* timesteps. It contains the timestep, the maximum torque per replica, the maximum torque per atom (in any replica), potential gradients in the initial, final, and climbing replicas, the forward and backward energy barriers, the total reaction coordinate (RDT), and the normalized reaction coordinate and potential energy of each replica.

The “maximum torque per replica” is the two-norm of the  $3N$ -length vector given by the cross product of a spin by its precession vector  $\omega$ , in each replica, maximized across replicas, which is what the *ttol* setting is checking against. In this case,  $N$  is all the atoms in each replica. The “maximum torque per atom” is the maximum torque component of any atom in any replica. The potential gradients are the two-norm of the  $3N$ -length magnetic precession vector solely due to the interaction potential i.e. without adding in inter-replica forces, and projected along the path tangent (as detailed in Appendix D of (*Bessaraba*)).

The “reaction coordinate” (RD) for each replica is the two-norm of the  $3N$ -length vector of geodesic distances between its spins and the preceding replica’s spins (see equation (13) of (*Bessaraba*)), added to the RD of the preceding replica. The RD of the first replica  $RD1 = 0.0$ ; the RD of the final replica  $RDN = RDT$ , the total reaction coordinate. The normalized RDs are divided by RDT, so that they form a monotonically increasing sequence from zero to one. When computing RD,  $N$  only includes the spins being operated on by the *fix neb/spin* command.

The forward (reverse) energy barrier is the potential energy of the highest replica minus the energy of the first (last) replica.

Supplementary information for all replicas can be printed out to the screen and master log.lammps file by adding the *verbose* keyword. This information include the following. The “GradVidottan” are the projections of the potential gradient for the replica  $i$  on its tangent vector (as detailed in Appendix D of (*Bessaraba*)). The “DNi” are the non normalized geodesic distances (see equation (13) of (*Bessaraba*)), between a replica  $i$  and the next replica  $i+1$ . For the last replica, this distance is not defined and a “NAN” value is the corresponding output.

When a NEB calculation does not converge properly, the supplementary information can help understanding what is going wrong.

When running on multiple partitions, LAMMPS produces additional log files for each partition, e.g. log.lammps.0, log.lammps.1, etc. For a GNEB calculation, these contain the thermodynamic output for each replica.

If *dump* commands in the input script define a filename that includes a *universe* or *uloop* style *variable*, then one dump file (per dump command) will be created for each replica. At the end of the GNEB calculation, the final snapshot in each file will contain the sequence of snapshots that transition the system over the energy barrier. Earlier snapshots will show the convergence of the replicas to the MEP.



Likewise, *restart* filenames can be specified with a *universe* or *uloop* style *variable*, to generate restart files for each replica. These may be useful if the GNEB calculation fails to converge properly to the MEP, and you wish to restart the calculation from an intermediate point with altered parameters.

A c file script is provided in the `tool/spin/interpolate_gneb` directory, that interpolates the MEP given the information provided by the *verbose* output option (as detailed in Appendix D of (*BessarabA*)).

---

## 16.90.4 Restrictions

This command can only be used if LAMMPS was built with the SPIN package. See the *Build package* doc page for more info.

For magnetic GNEB calculations, only the *spin\_none* value for the *line* keyword can be used when minimization styles *spin/cg* and *spin/lbfgs* are employed.

---

## 16.90.5 Related commands

*min/spin*, *fix neb/spin*

## 16.90.6 Default

none

---

(*BessarabA*) Bessarab, Uzdin, Jonsson, Comp Phys Comm, 196, 335-347 (2015).

## 16.91 neigh\_modify command

### 16.91.1 Syntax

**neigh\_modify** keyword values ...

- one or more keyword/value pairs may be listed

keyword = *delay* or *every* or *check* or *once* or *cluster* or *include* or *exclude* or *page\_*  
↔ or *one* or *binsize* or *collection/type* or *collection/interval*

*delay* value = N

N = delay building until this many steps since last build

*every* value = M

M = build neighbor list every this many steps

*check* value = *yes* or *no*

*yes* = only build if some atom has moved half the skin distance or more

*no* = always build on 1st step that *every* and *delay* are satisfied

*once* value = *yes* or *no*

*yes* = only build neighbor list once at start of run and never rebuild

*no* = rebuild neighbor list according to other settings

*cluster* value = *yes* or *no*

```

yes = check bond,angle,etc neighbor list for nearby clusters
no = do not check bond,angle,etc neighbor list for nearby clusters
include value = group-ID
group-ID = only build pair neighbor lists for atoms in this group
exclude values:
type M N
M,N = exclude if one atom in pair is type M, other is type N
group group1-ID group2-ID
group1-ID,group2-ID = exclude if one atom is in 1st group, other in 2nd
molecule/intra group-ID
group-ID = exclude if both atoms are in the same molecule and in group
molecule/inter group-ID
group-ID = exclude if both atoms are in different molecules and in group
none
delete all exclude settings
page value = N
N = number of pairs stored in a single neighbor page
one value = N
N = max number of neighbors of one atom
binsize value = size
size = bin size for neighbor list construction (distance units)
collection/type values = N arg1 ... argN
N = number of custom collections
arg = N separate lists of types (see below)
collection/interval values = N arg1 ... argN
N = number of custom collections
arg = N separate cutoffs for intervals (see below)

```

## 16.91.2 Examples

```

neigh_modify every 2 delay 10 check yes page 100000
neigh_modify exclude type 2 3
neigh_modify exclude group frozen frozen check no
neigh_modify exclude group residuel chain3
neigh_modify exclude molecule/intra rigid
neigh_modify collection/type 2 1*2,5 3*4
neigh_modify collection/interval 2 1.0 10.0

```

## 16.91.3 Description

This command sets parameters that affect the building and use of pairwise neighbor lists. Depending on what pair interactions and other commands are defined, a simulation may require one or more neighbor lists.

The *every*, *delay*, *check*, and *once* options affect how often lists are built as a simulation runs. The *delay* setting means never build new lists until at least N steps after the previous build. The *every* setting means build lists every M steps (after the delay has passed). If the *check* setting is *no*, the lists are built on the first step that satisfies the *delay* and *every* settings. If the *check* setting is *yes*, then the *every* and *delay* settings determine when a build may possibly be performed, but an actual build only occurs if some atom has moved more than half the skin distance (specified in the *neighbor* command) since the last build.

If the *once* setting is *yes*, then the neighbor list is only built once at the beginning of each run, and never rebuilt, except on steps when a restart file is written, or steps when a fix forces a rebuild to occur (e.g. fixes that create or delete atoms,

such as *fix deposit* or *fix evaporate*). This setting should only be made if you are certain atoms will not move far enough that the neighbor list should be rebuilt, e.g. running a simulation of a cold crystal. Note that it is not that expensive to check if neighbor lists should be rebuilt.

When the rRESPA integrator is used (see the *run\_style* command), the *every* and *delay* parameters refer to the longest (outermost) timestep.

The *cluster* option does a sanity test every time neighbor lists are built for bond, angle, dihedral, and improper interactions, to check that each set of 2, 3, or 4 atoms is a cluster of nearby atoms. It does this by computing the distance between pairs of atoms in the interaction and insuring they are not further apart than half the periodic box length. If they are, an error is generated, since the interaction would be computed between far-away atoms instead of their nearby periodic images. The only way this should happen is if the pairwise cutoff is so short that atoms that are part of the same interaction are not communicated as ghost atoms. This is an unusual model (e.g. no pair interactions at all) and the problem can be fixed by use of the *comm\_modify cutoff* command. Note that to save time, the default *cluster* setting is *no*, so that this check is not performed.

The *include* option limits the building of pairwise neighbor lists to atoms in the specified group. This can be useful for models where a large portion of the simulation is particles that do not interact with other particles or with each other via pairwise interactions. The group specified with this option must also be specified via the *atom\_modify first* command. Note that specifying “all” as the group-ID effectively turns off the *include* option.

The *exclude* option turns off pairwise interactions between certain pairs of atoms, by not including them in the neighbor list. These are sample scenarios where this is useful:

- In crack simulations, pairwise interactions can be shut off between 2 slabs of atoms to effectively create a crack.
- When a large collection of atoms is treated as frozen, interactions between those atoms can be turned off to save needless computation. E.g. Using the *fix setforce* command to freeze a wall or portion of a bio-molecule.
- When one or more rigid bodies are specified, interactions within each body can be turned off to save needless computation. See the *fix rigid* command for more details.

The *exclude type* option turns off the pairwise interaction if one atom is of type M and the other of type N. M can equal N. The *exclude group* option turns off the interaction if one atom is in the first group and the other is the second. Group1-ID can equal group2-ID. The *exclude molecule/intra* option turns off the interaction if both atoms are in the specified group and in the same molecule, as determined by their molecule ID. The *exclude molecule/inter* turns off the interaction between pairs of atoms that have different molecule IDs and are both in the specified group.

Each of the exclude options can be specified multiple times. The *exclude type* option is the most efficient option to use; it requires only a single check, no matter how many times it has been specified. The other exclude options are more expensive if specified multiple times; they require one check for each time they have been specified.

Note that the exclude options only affect pairwise interactions; see the *delete\_bonds* command for information on turning off bond interactions.

---

**Note:** Excluding pairwise interactions will not work correctly when also using a long-range solver via the *kpace\_style* command. LAMMPS will give a warning to this effect. This is because the short-range pairwise interaction needs to subtract off a term from the total energy for pairs whose short-range interaction is excluded, to compensate for how the long-range solver treats the interaction. This is done correctly for pairwise interactions that are excluded (or weighted) via the *special\_bonds* command. But it is not done for interactions that are excluded via these *neigh\_modify exclude* options.

---

The *page* and *one* options affect how memory is allocated for the neighbor lists. For most simulations the default settings for these options are fine, but if a very large problem is being run or a very long cutoff is being used, these parameters can be tuned. The indices of neighboring atoms are stored in “pages”, which are allocated one after another as they fill up. The size of each page is set by the *page* value. A new page is allocated when the next atom’s neighbors could potentially overflow the list. This threshold is set by the *one* value which tells LAMMPS the maximum number of neighbor’s one atom can have.

---

**Note:** LAMMPS can crash without an error message if the number of neighbors for a single particle is larger than the *page* setting, which means it is much, much larger than the *one* setting. This is because LAMMPS does not error check these limits for every pairwise interaction (too costly), but only after all the particle's neighbors have been found. This problem usually means something is very wrong with the way you have setup your problem (particle spacing, cutoff length, neighbor skin distance, etc). If you really expect that many neighbors per particle, then boost the *one* and *page* settings accordingly.

---

The *binsize* option allows you to specify what size of bins will be used in neighbor list construction to sort and find neighboring atoms. By default, for *neighbor style bin*, LAMMPS uses bins that are 1/2 the size of the maximum pair cutoff. For *neighbor style multi*, the bins are 1/2 the size of the collection interaction cutoff. Typically these are good values for minimizing the time for neighbor list construction. This setting overrides the default. If you make it too big, there is little overhead due to looping over bins, but more atoms are checked. If you make it too small, the optimal number of atoms is checked, but bin overhead goes up. If you set the *binsize* to 0.0, LAMMPS will use the default *binsize* of 1/2 the cutoff.

The *collection/type* option allows you to define collections of atom types, used by the *multi* neighbor mode. By grouping atom types with similar physical size or interaction cutoff lengths, one may be able to improve performance by reducing overhead. You must first specify the number of collections N to be defined followed by N lists of types. Each list consists of a series of type ranges separated by commas. The range can be specified as a single numeric value, or a wildcard asterisk can be used to specify a range of values. This takes the form "\*" or "\*n" or "n\*" or "m\*n". For example, if M = the number of atom types, then an asterisk with no numeric values means all types from 1 to M. A leading asterisk means all types from 1 to n (inclusive). A trailing asterisk means all types from n to M (inclusive). A middle asterisk means all types from m to n (inclusive). Note that all atom types must be included in exactly one of the N collections.

The *collection/interval* option provides a similar capability. This command allows a user to define collections by specifying a series of cutoff intervals. LAMMPS will automatically sort atoms into these intervals based on their type-dependent cutoffs or their finite size. You must first specify the number of collections N to be defined followed by N values representing the upper cutoff of each interval. This command is particularly useful for granular pair styles where the interaction distance of particles depends on their radius and may not depend on their atom type.

#### 16.91.4 Restrictions

If the "delay" setting is non-zero, then it must be a multiple of the "every" setting.

The *molecule/intra* and *molecule/inter* exclude options can only be used with atom styles that define molecule IDs.

The value of the *page* setting must be at least 10x larger than the *one* setting. This insures neighbor pages are not mostly empty space.

## 16.91.5 Related commands

*neighbor*, *delete\_bonds*

## 16.91.6 Default

The option defaults are delay = 10, every = 1, check = yes, once = no, cluster = no, include = all (same as no include option defined), exclude = none, page = 100000, one = 2000, and binsize = 0.0.

## 16.92 neighbor command

### 16.92.1 Syntax

```
neighbor skin style
```

- skin = extra distance beyond force cutoff (distance units)
- style = *bin* or *nsq* or *multi* or *multi/old*

### 16.92.2 Examples

```
neighbor 0.3 bin
neighbor 2.0 nsq
```

### 16.92.3 Description

This command sets parameters that affect the building of pairwise neighbor lists. All atom pairs within a neighbor cutoff distance equal to the their force cutoff plus the *skin* distance are stored in the list. Typically, the larger the skin distance, the less often neighbor lists need to be built, but more pairs must be checked for possible force interactions every timestep. The default value for *skin* depends on the choice of units for the simulation; see the default values below.

The *skin* distance is also used to determine how often atoms migrate to new processors if the *check* option of the *neigh\_modify* command is set to *yes*. Atoms are migrated (communicated) to new processors on the same timestep that neighbor lists are re-built.

The *style* value selects what algorithm is used to build the list. The *bin* style creates the list by binning which is an operation that scales linearly with N/P, the number of atoms per processor where N = total number of atoms and P = number of processors. It is almost always faster than the *nsq* style which scales as (N/P)<sup>2</sup>. For unsolvated small molecules in a non-periodic box, the *nsq* choice can sometimes be faster. Either style should give the same answers.

The *multi* style is a modified binning algorithm that is useful for systems with a wide range of cutoff distances, e.g. due to different size particles. For granular pair styles, cutoffs are set to the sum of the maximum atomic radii for each atom type. For the *bin* style, the bin size is set to 1/2 of the largest cutoff distance between any pair of atom types and a single set of bins is defined to search over for all atom types. This can be inefficient if one pair of types has a very long cutoff, but other type pairs have a much shorter cutoff. The *multi* style uses different sized bins for collections of different sized particles, where “size” may mean the physical size of the particle or its cutoff distance for interacting with other particles. Different sets of bins are then used to construct the neighbor lists as as further described by Shire, Hanley, and Stratford (*Shire*). This imposes some extra setup overhead, but the searches themselves may be much faster. By default, each atom type defines a separate collection of particles. For systems where two or more atom types have the same

size (either physical size or cutoff distance), the definition of collections can be customized, which can result in less overhead and faster performance. See the *neigh\_modify* command for how to define custom collections. Whether the collection definition is customized or not, also see the *comm\_modify mode multi* command for communication options that further improve performance in a manner consistent with neighbor style multi.

An alternate style, *multi/old*, sets the bin size to 1/2 of the shortest cutoff distance and multiple sets of bins are defined to search over for different atom types. This algorithm used to be the default *multi* algorithm in LAMMPS but was found to be significantly slower than the new approach. For now we are keeping the old option in case there are use cases where multi/old outperforms the new multi style.

---

**Note:** If there are multiple sub-styles in a *hybrid/overlay pair style* that cover the same atom types, but have significantly different cutoffs, the *multi* style does not apply. Instead, the *pair\_modify neigh/trim* setting applies (which is *yes* by default). Please check the neighbor list summary printed at the beginning of a calculation to verify that the desired set of neighbor list builds is performed.

---

The *neigh\_modify* command has additional options that control how often neighbor lists are built and which pairs are stored in the list.

When a run is finished, counts of the number of neighbors stored in the pairwise list and the number of times neighbor lists were built are printed to the screen and log file. See the *Run output* page for details.

## 16.92.4 Restrictions

none

## 16.92.5 Related commands

*neigh\_modify*, *units*, *comm\_modify*

## 16.92.6 Default

0.3 bin for units = lj, skin = 0.3 sigma  
2.0 bin for units = real or metal, skin = 2.0 Angstroms  
0.001 bin for units = si, skin = 0.001 meters = 1.0 mm  
0.1 bin for units = cgs, skin = 0.1 cm = 1.0 mm

---

(Shire) Shire, Hanley and Stratford, Comp Part Mech, (2020).

## 16.93 newton command

### 16.93.1 Syntax

```
newton flag  
newton flag1 flag2
```

- flag = *on* or *off* for both pairwise and bonded interactions
- flag1 = *on* or *off* for pairwise interactions
- flag2 = *on* or *off* for bonded interactions

### 16.93.2 Examples

```
newton off  
newton on off
```

### 16.93.3 Description

This command turns Newton's third law *on* or *off* for pairwise and bonded interactions. For most problems, setting Newton's third law to *on* means a modest savings in computation at the cost of two times more communication. Whether this is faster depends on problem size, force cutoff lengths, a machine's compute/communication ratio, and how many processors are being used.

Setting the pairwise newton flag to *off* means that if two interacting atoms are on different processors, both processors compute their interaction and the resulting force information is not communicated. Similarly, for bonded interactions, *newton off* means that if a bond, angle, dihedral, or improper interaction contains atoms on 2 or more processors, the interaction is computed by each processor.

LAMMPS should produce the same answers for any newton flag settings, except for round-off issues.

With *run\_style respa* and only bonded interactions (bond, angle, etc) computed in the innermost timestep, it may be faster to turn *newton off* for bonded interactions, to avoid extra communication in the innermost loop.

### 16.93.4 Restrictions

The newton bond setting cannot be changed after the simulation box is defined by a *read\_data* or *create\_box* command.

### 16.93.5 Related commands

*run\_style respa*

## 16.93.6 Default

```
newton on
```

## 16.94 next command

### 16.94.1 Syntax

```
next variables
```

- variables = one or more variable names

### 16.94.2 Examples

```
next x  
next a t x myTemp
```

### 16.94.3 Description

This command is used with variables defined by the *variable* command. It assigns the next value to the variable from the list of values defined for that variable by the *variable* command. Thus when that variable is subsequently substituted for in an input script command, the new value is used.

See the *variable* command for info on how to define and use different kinds of variables in LAMMPS input scripts. If a variable name is a single lower-case character from “a” to “z”, it can be used in an input script command as \$a or \$z. If it is multiple letters, it can be used as \${myTemp}.

If multiple variables are used as arguments to the *next* command, then all must be of the same variable style: *index*, *loop*, *file*, *universe*, or *uloop*. An exception is that *universe*- and *uloop*-style variables can be mixed in the same *next* command.

All the variables specified with the *next* command are incremented by one value from their respective list of values. A *file*-style variable reads the next line from its associated file. An *atomfile*-style variable reads the next set of lines (one per atom) from its associated file. *String*- or *atom*- or *equal*- or *world*-style variables cannot be used with the *next* command, since they only store a single value.

When any of the variables in the *next* command has no more values, a flag is set that causes the input script to skip the next *jump* command encountered. This enables a loop containing a *next* command to exit. As explained in the *variable* command, the variable that has exhausted its values is also deleted. This allows it to be used and re-defined later in the input script. *File*-style and *atomfile*-style variables are exhausted when the end-of-file is reached.

When the *next* command is used with *index*- or *loop*-style variables, the next value is assigned to the variable for all processors. When the *next* command is used with *file*-style variables, the next line is read from its file and the string assigned to the variable. When the *next* command is used with *atomfile*-style variables, the next set of per-atom values is read from its file and assigned to the variable.

When the *next* command is used with *universe*- or *uloop*-style variables, all *universe*- or *uloop*-style variables must be listed in the *next* command. This is because of the manner in which the incrementing is done, using a single lock file for all variables. The next value (for each variable) is assigned to whichever processor partition executes the command first. All processors in the partition are assigned the same value(s). Running LAMMPS on multiple partitions of processors via the *-partition command-line switch*. *Universe*- and *uloop*-style variables are incremented using the files



“tmp.lammps.variable” and “tmp.lammps.variable.lock” which you will see in your directory during and after such a LAMMPS run.

Here is an example of running a series of simulations using the next command with an *index*-style variable. If this input script is named in.polymer, 8 simulations would be run using data files from directories run1 through run8.

```
variable d index run1 run2 run3 run4 run5 run6 run7 run8
shell cd $d
read_data data.polymer
run 10000
shell cd ..
clear
next d
jump in.polymer
```

If the variable “d” were of style *universe*, and the same in.polymer input script were run on 3 partitions of processors, then the first 3 simulations would begin, one on each set of processors. Whichever partition finished first, it would assign variable “d” the fourth value and run another simulation, and so forth until all 8 simulations were finished.

Jump and next commands can also be nested to enable multi-level loops. For example, this script will run 15 simulations in a double loop.

```
variable i loop 3
  variable j loop 5
  clear
  ...
  read_data data.polymer.$i$j
  print Running simulation $i.$j
  run 10000
  next j
  jump in.script
next i
jump in.script
```

Here is an example of a double loop which uses the *if* and *jump* commands to break out of the inner loop when a condition is met, then continues iterating through the outer loop.

```
label      loopa
variable  a loop 5
  label   loopb
  variable b loop 5
  print   "A,B = $a,$b"
  run     10000
  if      $b > 2 then "jump in.script break"
  next    b
  jump    in.script loopb
label     break
variable  b delete

next      a
jump     in.script loopa
```

## 16.94.4 Restrictions

As described above.

## 16.94.5 Related commands

*jump, include, shell, variable,*

## 16.94.6 Default

none

# 16.95 package command

## 16.95.1 Syntax

```
package style args
```

- *style* = *gpu* or *intel* or *kokkos* or *omp*
  - *args* = arguments specific to the style
- gpu* args = *Ngpu* keyword value ...  
*Ngpu* = # of GPUs per node  
zero or more keyword/value pairs may be appended  
keywords = *neigh* or *newton* or *pair/only* or *binsize* or *split* or *gpuID* or *tpa* or *↪**blocksize* or *omp* or *platform* or *device\_type* or *ocl\_args*  
*neigh* value = *yes* or *no*  
  *yes* = neighbor list build on GPU (default)  
  *no* = neighbor list build on CPU  
*newton* = *off* or *on*  
  *off* = set Newton pairwise flag off (default and required)  
  *on* = set Newton pairwise flag on (currently not allowed)  
*pair/only* = *off* or *on*  
  *off* = apply "gpu" suffix to all available styles in the GPU package (default)  
  *on* = apply "gpu" suffix only pair styles  
*binsize* value = *size*  
  *size* = bin size for neighbor list construction (distance units)  
*split* = *fraction*  
  *fraction* = fraction of atoms assigned to GPU (default = 1.0)  
*tpa* value = *Nlanes*  
  *Nlanes* = # of GPU vector lanes (CUDA threads) used per atom  
*blocksize* value = *size*  
  *size* = thread block size for pair force computation  
*omp* value = *Nthreads*  
  *Nthreads* = number of OpenMP threads to use on CPU (default = 0)  
*platform* value = *id*  
  *id* = For OpenCL, platform ID for the GPU or accelerator  
*gpuID* values = *id*  
  *id* = ID of first GPU to be used on each node

```

    device_type value = intelgpu or nvidiagpu or amdgpu or applegpu or generic or
↳ custom, val1, val2, ...
    val1, val2, ... = custom OpenCL accelerator configuration parameters (see below
↳ for details)
    ocl_args value = args
    args = List of additional OpenCL compiler arguments delimited by colons
intel args = NPhi keyword value ...
Nphi = # of co-processors per node
zero or more keyword/value pairs may be appended
keywords = mode or omp or lrt or balance or ghost or tpc or tptask or no_affinity
mode value = single or mixed or double
    single = perform force calculations in single precision
    mixed = perform force calculations in mixed precision
    double = perform force calculations in double precision
omp value = Nthreads
    Nthreads = number of OpenMP threads to use on CPU (default = 0)
lrt value = yes or no
    yes = use additional thread dedicated for some PPPM calculations
    no = do not dedicate an extra thread for some PPPM calculations
balance value = split
    split = fraction of work to offload to co-processor, -1 for dynamic
ghost value = yes or no
    yes = include ghost atoms for offload
    no = do not include ghost atoms for offload
tpc value = Ntpc
    Ntpc = max number of co-processor threads per co-processor core (default = 4)
tptask value = Ntptask
    Ntptask = max number of co-processor threads per MPI task (default = 240)
no_affinity values = none
kokkos args = keyword value ...
zero or more keyword/value pairs may be appended
keywords = neigh or neigh/req or neigh/thread or neigh/transpose or newton or
↳ binsize or comm or comm/exchange or comm/forward or comm/pair/forward or comm/fix/
↳ forward or comm/reverse or comm/pair/reverse or gpu/aware or pair/only
neigh value = full or half
    full = full neighbor list
    half = half neighbor list built in thread-safe manner
neigh/req value = full or half
    full = full neighbor list
    half = half neighbor list built in thread-safe manner
neigh/thread value = off or on
    off = thread only over atoms
    on = thread over both atoms and neighbors
neigh/transpose value = off or on
    off = use same memory layout for GPU neigh list build as pair style
    on = use transposed memory layout for GPU neigh list build
newton = off or on
    off = set Newton pairwise and bonded flags off
    on = set Newton pairwise and bonded flags on
binsize value = size
    size = bin size for neighbor list construction (distance units)
comm value = no or host or device
    use value for comm/exchange and comm/forward and comm/pair/forward and comm/
↳ fix/forward and comm/reverse

```

```

comm/exchange value = no or host or device
comm/forward value = no or host or device
comm/pair/forward value = no or device
comm/fix/forward value = no or device
comm/reverse value = no or host or device
  no = perform communication pack/unpack in non-KOKKOS mode
  host = perform pack/unpack on host (e.g. with OpenMP threading)
  device = perform pack/unpack on device (e.g. on GPU)
comm/pair/reverse value = no or device
  no = perform communication pack/unpack in non-KOKKOS mode
  device = perform pack/unpack on device (e.g. on GPU)
gpu/aware = off or on
  off = do not use GPU-aware MPI
  on = use GPU-aware MPI (default)
pair/only = off or on
  off = use device acceleration (e.g. GPU) for all available styles in the
↳KOKKOS package (default)
  on = use device acceleration only for pair styles (and host acceleration for
↳others)
omp args = Nthreads keyword value ...
Nthreads = # of OpenMP threads to associate with each MPI process
zero or more keyword/value pairs may be appended
keywords = neigh
  neigh value = yes or no
  yes = threaded neighbor list build (default)
  no = non-threaded neighbor list build

```

## 16.95.2 Examples

```

package gpu 0
package gpu 1 split 0.75
package gpu 2 split -1.0
package gpu 0 omp 2 device_type intelgpu
package kokkos neigh half comm device
package omp 0 neigh no
package omp 4
package intel 1
package intel 2 omp 4 mode mixed balance 0.5

```

## 16.95.3 Description

This command invokes package-specific settings for the various accelerator packages available in LAMMPS. Currently the following packages use settings from this command: GPU, INTEL, KOKKOS, and OPENMP.

If this command is specified in an input script, it must be near the top of the script, before the simulation box has been defined. This is because it specifies settings that the accelerator packages use in their initialization, before a simulation is defined.

This command can also be specified from the command-line when launching LAMMPS, using the “-pk” *command-line switch*. The syntax is exactly the same as when used in an input script.

Note that all of the accelerator packages require the package command to be specified (except the OPT package), if the package is to be used in a simulation (LAMMPS can be built with an accelerator package without using it in a particular

simulation). However, in all cases, a default version of the command is typically invoked by other accelerator settings.

The KOKKOS package requires a “-k on” *command-line switch* respectively, which invokes a “package kokkos” command with default settings.

For the GPU, INTEL, and OPENMP packages, if a “-sf gpu” or “-sf intel” or “-sf omp” *command-line switch* is used to auto-append accelerator suffixes to various styles in the input script, then those switches also invoke a “package gpu”, “package intel”, or “package omp” command with default settings.

---

**Note:** A package command for a particular style can be invoked multiple times when a simulation is setup, e.g. by the *-c on*, *-k on*, *-sf*, and *-pk* *command-line switches*, and by using this command in an input script. Each time it is used all of the style options are set, either to default values or to specified settings. I.e. settings from previous invocations do not persist across multiple invocations.

---

See the *Accelerator packages* page for more details about using the various accelerator packages for speeding up LAMMPS simulations.

---

The *gpu* style invokes settings associated with the use of the GPU package.

The *Ngpu* argument sets the number of GPUs per node. If *Ngpu* is 0 and no other keywords are specified, GPU or accelerator devices are auto-selected. In this process, all platforms are searched for accelerator devices and GPUs are chosen if available. The device with the highest number of compute cores is selected. The number of devices is increased to be the number of matching accelerators with the same number of compute cores. If there are more devices than MPI tasks, the additional devices will be unused. The auto-selection of GPUs/ accelerator devices and platforms can be restricted by specifying a non-zero value for *Ngpu* and / or using the *gpuID*, *platform*, and *device\_type* keywords as described below. If there are more MPI tasks (per node) than GPUs, multiple MPI tasks will share each GPU.

Optional keyword/value pairs can also be specified. Each has a default value as listed below.

The *neigh* keyword specifies where neighbor lists for pair style computation will be built. If *neigh* is *yes*, which is the default, neighbor list building is performed on the GPU. If *neigh* is *no*, neighbor list building is performed on the CPU. GPU neighbor list building currently cannot be used with a triclinic box. GPU neighbor lists are not compatible with commands that are not GPU-enabled. When a non-GPU enabled command requires a neighbor list, it will also be built on the CPU. In these cases, it will typically be more efficient to only use CPU neighbor list builds.

The *newton* keyword sets the Newton flags for pairwise (not bonded) interactions to *off* or *on*, the same as the *newton* command allows. Currently, only an *off* value is allowed, since all the GPU package pair styles require this setting. This means more computation is done, but less communication. In the future a value of *on* may be allowed, so the *newton* keyword is included as an option for compatibility with the package command for other accelerator styles. Note that the *newton* setting for bonded interactions is not affected by this keyword.

The *pair/only* keyword can change how any “gpu” suffix is applied. By default a suffix is applied to all styles for which an accelerated variant is available. However, that is not always the most effective way to use an accelerator. With *pair/only* set to *on* the suffix will only be applied to supported pair styles, which tend to be the most effective in using an accelerator and their operation can be overlapped with all other computations on the CPU.

The *binsize* keyword sets the size of bins used to bin atoms in neighbor list builds performed on the GPU, if *neigh* = *yes* is set. If *binsize* is set to 0.0 (the default), then the binsize is set automatically using heuristics in the GPU package.

The *split* keyword can be used for load balancing force calculations between CPU and GPU cores in GPU-enabled pair styles. If  $0 < split < 1.0$ , a fixed fraction of particles is offloaded to the GPU while force calculation for the other particles occurs simultaneously on the CPU. If  $split < 0.0$ , the optimal fraction (based on CPU and GPU timings) is calculated every 25 timesteps, i.e. dynamic load-balancing across the CPU and GPU is performed. If  $split = 1.0$ , all force calculations for GPU accelerated pair styles are performed on the GPU. In this case, other *hybrid* pair interactions, *bond*, *angle*, *dihedral*, *improper*, and *long-range* calculations can be performed on the CPU while the GPU is performing force calculations for the GPU-enabled pair style. If all CPU force computations complete before the GPU completes, LAMMPS will block until the GPU has finished before continuing the timestep.

As an example, if you have two GPUs per node and 8 CPU cores per node, and would like to run on 4 nodes (32 cores) with dynamic balancing of force calculation across CPU and GPU cores, you could specify

```
mpirun -np 32 -sf gpu -in in.script      # launch command
package gpu 2 split -1                  # input script command
```

In this case, all CPU cores and GPU devices on the nodes would be utilized. Each GPU device would be shared by 4 CPU cores. The CPU cores would perform force calculations for some fraction of the particles at the same time the GPUs performed force calculation for the other particles.

The *gpuID* keyword is used to specify the first ID for the GPU or other accelerator that LAMMPS will use. For example, if the ID is 1 and *Ngpu* is 3, GPUs 1-3 will be used. Device IDs should be determined from the output of `nvc_get_devices`, `ocl_get_devices`, or `hip_get_devices` as provided in the `lib/gpu` directory. When using OpenCL with accelerators that have main memory NUMA, the accelerators can be split into smaller virtual accelerators for more efficient use with MPI.

The *tpa* keyword sets the number of GPU vector lanes per atom used to perform force calculations. With a default value of 1, the number of lanes will be chosen based on the pair style, however, the value can be set explicitly with this keyword to fine-tune performance. For large cutoffs or with a small number of particles per GPU, increasing the value can improve performance. The number of lanes per atom must be a power of 2 and currently cannot be greater than the SIMD width for the GPU / accelerator. In the case it exceeds the SIMD width, it will automatically be decreased to meet the restriction.

The *blocksize* keyword allows you to tweak the number of threads used per thread block. This number should be a multiple of 32 (for GPUs) and its maximum depends on the specific GPU hardware. Typical choices are 64, 128, or 256. A larger block size increases occupancy of individual GPU cores, but reduces the total number of thread blocks, thus may lead to load imbalance. On modern hardware, the sensitivity to the blocksize is typically low.

The *Nthreads* value for the *omp* keyword sets the number of OpenMP threads allocated for each MPI task. This setting controls OpenMP parallelism only for routines run on the CPUs. For more details on setting the number of OpenMP threads, see the discussion of the *Nthreads* setting on this page for the “package omp” command. The meaning of *Nthreads* is exactly the same for the GPU, INTEL, and GPU packages.

The *platform* keyword is only used with OpenCL to specify the ID for an OpenCL platform. See the output from `ocl_get_devices` in the `lib/gpu` directory. In LAMMPS only one platform can be active at a time and by default (`id=-1`) the platform is auto-selected to find the GPU with the most compute cores. When *Ngpu* or other keywords are specified, the auto-selection is appropriately restricted. For example, if *Ngpu* is 3, only platforms with at least 3 accelerators are considered. Similar restrictions can be enforced by the *gpuID* and *device\_type* keywords.

The *device\_type* keyword can be used for OpenCL to specify the type of GPU to use or specify a custom configuration for an accelerator. In most cases this selection will be automatic and there is no need to use the keyword. The *applegpu* type is not specific to a particular GPU vendor, but is separate due to the more restrictive Apple OpenCL implementation. For expert users, to specify a custom configuration, the *custom* keyword followed by the next parameters can be specified:

CONFIG\_ID, SIMD\_SIZE, MEM\_THREADS, SHUFFLE\_AVAIL, FAST\_MATH, THREADS\_PER\_ATOM, THREADS\_PER\_CHARGE, THREADS\_PER\_THREE, BLOCK\_PAIR, BLOCK\_BIO\_PAIR, BLOCK\_ELLIPSE, PPPM\_BLOCK\_1D, BLOCK\_NBOR\_BUILD, BLOCK\_CELL\_2D, BLOCK\_CELL\_ID, MAX\_SHARED\_TYPES, MAX\_BIO\_SHARED\_TYPES, PPPM\_MAX\_SPLINE.

CONFIG\_ID can be 0. SHUFFLE\_AVAIL in {0,1} indicates that inline-PTX (NVIDIA) or OpenCL extensions (Intel) should be used for horizontal vector operations. FAST\_MATH in {0,1} indicates that OpenCL fast math optimizations are used during the build and hardware-accelerated transcendental functions are used when available. THREADS\_PER\_\* give the default *tpa* values for ellipsoidal models, styles using charge, and any other styles. The BLOCK\_\* parameters specify the block sizes for various kernel calls and the MAX\_\*SHARED\*\_ parameters are used to determine the amount of local shared memory to use for storing model parameters.

For OpenCL, the routines are compiled at runtime for the specified GPU or accelerator architecture. The *ocl\_args* keyword can be used to specify additional flags for the runtime build.

The *intel* style invokes settings associated with the use of the INTEL package. All of its settings, except the *omp* and *mode* keywords, are ignored if LAMMPS was not built with Xeon Phi co-processor support. All of its settings, including the *omp* and *mode* keyword are applicable if LAMMPS was built with co-processor support.

The *Nphi* argument sets the number of co-processors per node. This can be set to any value, including 0, if LAMMPS was not built with co-processor support.

Optional keyword/value pairs can also be specified. Each has a default value as listed below.

The *Nthreads* value for the *omp* keyword sets the number of OpenMP threads allocated for each MPI task. This setting controls OpenMP parallelism only for routines run on the CPUs. For more details on setting the number of OpenMP threads, see the discussion of the *Nthreads* setting on this page for the “package omp” command. The meaning of *Nthreads* is exactly the same for the GPU, INTEL, and GPU packages.

The *mode* keyword determines the precision mode to use for computing pair style forces, either on the CPU or on the co-processor, when using a INTEL supported *pair style*. It can take a value of *single*, *mixed* which is the default, or *double*. *Single* means single precision is used for the entire force calculation. *Mixed* means forces between a pair of atoms are computed in single precision, but accumulated and stored in double precision, including storage of forces, torques, energies, and virial quantities. *Double* means double precision is used for the entire force calculation.

The *lrt* keyword can be used to enable “Long Range Thread (LRT)” mode. It can take a value of *yes* to enable and *no* to disable. LRT mode generates an extra thread (in addition to any OpenMP threads specified with the OMP\_NUM\_THREADS environment variable or the *omp* keyword). The extra thread is dedicated for performing part of the *PPPM solver* computations and communications. This can improve parallel performance on processors supporting Simultaneous Multithreading (SMT) such as Hyper-Threading (HT) on Intel processors. In this mode, one additional thread is generated per MPI process. LAMMPS will generate a warning in the case that more threads are used than available in SMT hardware on a node. If the *PPPM solver* from the INTEL package is not used, then the LRT setting is ignored and no extra threads are generated. Enabling LRT will replace the *run\_style* with the *verlet/lrt/intel* style that is identical to the default *verlet* style aside from supporting the LRT feature. This feature requires setting the pre-processor flag `-DLMP_INTEL_USELRT` in the makefile when compiling LAMMPS.

The *balance* keyword sets the fraction of *pair style* work offloaded to the co-processor for split values between 0.0 and 1.0 inclusive. While this fraction of work is running on the co-processor, other calculations will run on the host, including neighbor and pair calculations that are not offloaded, as well as angle, bond, dihedral, kspace, and some MPI communications. If *split* is set to -1, the fraction of work is dynamically adjusted automatically throughout the run. This typically give performance within 5 to 10 percent of the optimal fixed fraction.

The *ghost* keyword determines whether or not ghost atoms, i.e. atoms at the boundaries of processor sub-domains, are offloaded for neighbor and force calculations. When the value = “no”, ghost atoms are not offloaded. This option can reduce the amount of data transfer with the co-processor and can also overlap MPI communication of forces with computation on the co-processor when the *newton pair* setting is “on”. When the value = “yes”, ghost atoms are offloaded. In some cases this can provide better performance, especially if the *balance* fraction is high.

The *tpc* keyword sets the max # of co-processor threads *Ntpc* that will run on each core of the co-processor. The default value = 4, which is the number of hardware threads per core supported by the current generation Xeon Phi chips.

The *tptask* keyword sets the max # of co-processor threads ( $N_{tptask} \times$  assigned to each MPI task. The default value = 240, which is the total # of threads an entire current generation Xeon Phi chip can run ( $240 = 60 \text{ cores} \times 4 \text{ threads/core}$ ). This means each MPI task assigned to the Phi will have enough threads for the chip to run the max allowed, even if only 1 MPI task is assigned. If 8 MPI tasks are assigned to the Phi, each will run with 30 threads. If you wish to limit the number of threads per MPI task, set *tptask* to a smaller value. E.g. for *tptask* = 16, if 8 MPI tasks are assigned, each will run with 16 threads, for a total of 128.

Note that the default settings for *tpc* and *tptask* are fine for most problems, regardless of how many MPI tasks you assign to a Phi.

The *no\_affinity* keyword will turn off automatic setting of core affinity for MPI tasks and OpenMP threads on the host when using offload to a co-processor. Affinity settings are used when possible to prevent MPI tasks and OpenMP threads

from being on separate NUMA domains and to prevent offload threads from interfering with other processes/threads used for LAMMPS.

---

The *kokkos* style invokes settings associated with the use of the KOKKOS package.

All of the settings are optional keyword/value pairs. Each has a default value as listed below.

The *neigh* keyword determines how neighbor lists are built. A value of *half* uses a thread-safe variant of half-neighbor lists, the same as used by most pair styles in LAMMPS, which is the default when running on CPUs (i.e. the Kokkos CUDA back end is not enabled).

A value of *full* uses a full neighbor lists and is the default when running on GPUs. This performs twice as much computation as the *half* option, however that is often a win because it is thread-safe and does not require atomic operations in the calculation of pair forces. For that reason, *full* is the default setting for GPUs. However, when running on CPUs, a *half* neighbor list is the default because it are often faster, just as it is for non-accelerated pair styles. Similarly, the *neigh/peq* keyword determines how neighbor lists are built for *fix qeq/reaxff/kk*.

If the *neigh/thread* keyword is set to *off*, then the KOKKOS package threads only over atoms. However, for small systems, this may not expose enough parallelism to keep a GPU busy. When this keyword is set to *on*, the KOKKOS package threads over both atoms and neighbors of atoms. When using *neigh/thread on*, a full neighbor list must also be used. Using *neigh/thread on* may be slower for large systems, so this this option is turned on by default only when there are 16K atoms or less owned by an MPI rank and when using a full neighbor list. Not all KOKKOS-enabled potentials support this keyword yet, and only thread over atoms. Many simple pairwise potentials such as Lennard-Jones do support threading over both atoms and neighbors.

If the *neigh/transpose* keyword is set to *off*, then the KOKKOS package will use the same memory layout for building the neighbor list on GPUs as used for the pair style. When this keyword is set to *on* it will use a different (transposed) memory layout to build the neighbor list on GPUs. This can be faster in some cases (e.g. ReaxFF HNS benchmark) but slower in others (e.g. Lennard Jones benchmark). The copy between different memory layouts is done out of place and therefore doubles the memory overhead of the neighbor list, which can be significant.

The *newton* keyword sets the Newton flags for pairwise and bonded interactions to *off* or *on*, the same as the *newton* command allows. The default for GPUs is *off* because this will almost always give better performance for the KOKKOS package. This means more computation is done, but less communication. However, when running on CPUs a value of *on* is the default since it can often be faster, just as it is for non-accelerated pair styles

The *binsize* keyword sets the size of bins used to bin atoms during neighbor list builds. The same value can be set by the *neigh\_modify binsize* command. Making it an option in the package kokkos command allows it to be set from the command line. The default value for CPUs is 0.0, which means the LAMMPS default will be used, which is bins = 1/2 the size of the pairwise cutoff + neighbor skin distance. This is fine when neighbor lists are built on the CPU. For GPU builds, a 2x larger binsize equal to the pairwise cutoff + neighbor skin is often faster, which is the default. Note that if you use a longer-than-usual pairwise cutoff, e.g. to allow for a smaller fraction of KSpace work with a *long-range Coulombic solver* because the GPU is faster at performing pairwise interactions, then this rule of thumb may give too large a binsize and the default should be overridden with a smaller value.

The *comm* and *comm/exchange* and *comm/forward* and *comm/pair/forward* and *comm/fix/forward* and *comm/reverse* and *comm/pair/reverse* keywords determine whether the host or device performs the packing and unpacking of data when communicating per-atom data between processors. “Exchange” communication happens only on timesteps that neighbor lists are rebuilt. The data is only for atoms that migrate to new processors. “Forward” communication happens every timestep. “Reverse” communication happens every timestep if the *newton* option is on. The data is for atom coordinates and any other atom properties that needs to be updated for ghost atoms owned by each processor. “Pair/comm” controls additional communication in pair styles, such as pair\_style EAM. “Fix/comm” controls additional communication in fixes, such as fix SHAKE.

The *comm* keyword is simply a short-cut to set the same value for all the comm keywords.

The value options for the keywords are *no* or *host* or *device*. A value of *no* means to use the standard non-KOKKOS method of packing/unpacking data for the communication. A value of *host* means to use the host, typically a multi-core



CPU, and perform the packing/unpacking in parallel with threads. A value of *device* means to use the device, typically a GPU, to perform the packing/unpacking operation.

For the *comm/pair/forward* or *comm/fix/forward* or *comm/pair/reverse* keywords, if a value of *host* is used it will be automatically be changed to *no* since these keywords don't support *host* mode. The value of *no* will also always be used when running on the CPU, i.e. setting the value to *device* will have no effect if the *pair/fix* style is running on the CPU. For the *comm/fix/forward* or *comm/pair/reverse* keywords, not all styles support *device* mode and in that case will run in *no* mode instead.

The optimal choice for these keywords depends on the input script and the hardware used. The *no* value is useful for verifying that the Kokkos-based *host* and *device* values are working correctly. It is the default when running on CPUs since it is usually the fastest.

When running on CPUs or Xeon Phi, the *host* and *device* values work identically. When using GPUs, the *device* value is the default since it will typically be optimal if all of your styles used in your input script are supported by the KOKKOS package. In this case data can stay on the GPU for many timesteps without being moved between the host and GPU, if you use the *device* value. If your script uses styles (e.g. *fixes*) which are not yet supported by the KOKKOS package, then data has to be moved between the host and device anyway, so it is typically faster to let the host handle communication, by using the *host* value. Using *host* instead of *no* will enable use of multiple threads to pack/unpack communicated data. When running small systems on a GPU, performing the exchange pack/unpack on the host CPU can give speedup since it reduces the number of CUDA kernel launches.

The *gpu/aware* keyword chooses whether GPU-aware MPI will be used. When this keyword is set to *on*, buffers in GPU memory are passed directly through MPI send/receive calls. This reduces overhead of first copying the data to the host CPU. However GPU-aware MPI is not supported on all systems, which can lead to segmentation faults and would require using a value of *off*. If LAMMPS can safely detect that GPU-aware MPI is not available (currently only possible with OpenMPI v2.0.0 or later), then the *gpu/aware* keyword is automatically set to *off* by default. When the *gpu/aware* keyword is set to *off* while any of the *comm* keywords are set to *device*, the value for these *comm* keywords will be automatically changed to *no*. This setting has no effect if not running on GPUs or if using only one MPI rank. GPU-aware MPI is available for OpenMPI 1.8 (or later versions), Mvapich2 1.9 (or later) when the "MV2\_USE\_CUDA" environment variable is set to "1", CrayMPI, and IBM Spectrum MPI when the "-gpu" flag is used.

The *pair/only* keyword can change how the KOKKOS suffix "kk" is applied when using an accelerator device. By default device acceleration is always used for all available styles. With *pair/only* set to *on* the suffix setting will choose device acceleration only for pair styles and run all other force computations on the host CPU. The *comm* flags will also automatically be changed to *no*. This can result in better performance for certain configurations and system sizes.

---

The *omp* style invokes settings associated with the use of the OPENMP package.

The *Nthreads* argument sets the number of OpenMP threads allocated for each MPI task. For example, if your system has nodes with dual quad-core processors, it has a total of 8 cores per node. You could use two MPI tasks per node (e.g. using the *-ppn* option of the *mpirun* command in MPICH or *-npernode* in OpenMPI), and set *Nthreads* = 4. This would use all 8 cores on each node. Note that the product of MPI tasks \* threads/task should not exceed the physical number of cores (on a node), otherwise performance will suffer.

Setting *Nthreads* = 0 instructs LAMMPS to use whatever value is the default for the given OpenMP environment. This is usually determined via the *OMP\_NUM\_THREADS* environment variable or the compiler runtime. Note that in most cases the default for OpenMP capable compilers is to use one thread for each available CPU core when *OMP\_NUM\_THREADS* is not explicitly set, which can lead to poor performance.

Here are examples of how to set the environment variable when launching LAMMPS:

```
env OMP_NUM_THREADS=4 lmp_machine -sf omp -in in.script
env OMP_NUM_THREADS=2 mpirun -np 2 lmp_machine -sf omp -in in.script
mpirun -x OMP_NUM_THREADS=2 -np 2 lmp_machine -sf omp -in in.script
```

or you can set it permanently in your shell's start-up script. All three of these examples use a total of 4 CPU cores.

Note that different MPI implementations have different ways of passing the `OMP_NUM_THREADS` environment variable to all MPI processes. The second example line above is for MPICH; the third example line with `-x` is for OpenMPI. Check your MPI documentation for additional details.

What combination of threads and MPI tasks gives the best performance is difficult to predict and can depend on many components of your input. Not all features of LAMMPS support OpenMP threading via the OPENMP package and the parallel efficiency can be very different, too.

---

**Note:** If you build LAMMPS with the GPU, INTEL, and / or OPENMP packages, be aware these packages all allow setting of the *Nthreads* value via their package commands, but there is only a single global *Nthreads* value used by OpenMP. Thus if multiple package commands are invoked, you should insure the values are consistent. If they are not, the last one invoked will take precedence, for all packages. Also note that if the *-sf hybrid intel omp command-line switch* is used, it invokes a “package intel” command, followed by a “package omp” command, both with a setting of *Nthreads* = 0. Likewise for a hybrid suffix for gpu and omp. Note that KOKKOS also supports setting the number of OpenMP threads from the command line using the “-k on” *command-line switch*. The default for KOKKOS is 1 thread per MPI task, so any other number of threads should be explicitly set using the “-k on” command-line switch (and this setting should be consistent with settings from any other packages used).

---

Optional keyword/value pairs can also be specified. Each has a default value as listed below.

The *neigh* keyword specifies whether neighbor list building will be multi-threaded in addition to force calculations. If *neigh* is set to *no* then neighbor list calculation is performed only by MPI tasks with no OpenMP threading. If *mode* is *yes* (the default), a multi-threaded neighbor list build is used. Using *neigh* = *yes* is almost always faster and should produce identical neighbor lists at the expense of using more memory. Specifically, neighbor list pages are allocated for all threads at the same time and each thread works within its own pages.

---

## 16.95.4 Restrictions

This command cannot be used after the simulation box is defined by a *read\_data* or *create\_box* command.

The *gpu* style of this command can only be invoked if LAMMPS was built with the GPU package. See the *Build package* doc page for more info.

The *intel* style of this command can only be invoked if LAMMPS was built with the INTEL package. See the *Build package* page for more info.

The *kk* style of this command can only be invoked if LAMMPS was built with the KOKKOS package. See the *Build package* doc page for more info.

The *omp* style of this command can only be invoked if LAMMPS was built with the OPENMP package. See the *Build package* doc page for more info.

## 16.95.5 Related commands

*suffix, -pk command-line switch*

## 16.95.6 Default

For the GPU package, the default is `Ngpu = 0` and the option defaults are `neigh = yes`, `newton = off`, `binsize = 0.0`, `split = 1.0`, `gpuID = 0` to `Ngpu-1`, `tpa = 1`, `omp = 0`, and `platform=-1`. These settings are made automatically if the “-sf gpu” *command-line switch* is used. If it is not used, you must invoke the package `gpu` command in your input script or via the “-pk gpu” *command-line switch*.

For the INTEL package, the default is `Nphi = 1` and the option defaults are `omp = 0`, `mode = mixed`, `lrt = no`, `balance = -1`, `tpc = 4`, `tptask = 240`. The default ghost option is determined by the pair style being used. This value is output to the screen in the offload report at the end of each run. Note that all of these settings, except “omp” and “mode”, are ignored if LAMMPS was not built with Xeon Phi co-processor support. These settings are made automatically if the “-sf intel” *command-line switch* is used. If it is not used, you must invoke the package `intel` command in your input script or via the “-pk intel” *command-line switch*.

For the KOKKOS package, the option defaults for GPUs are `neigh = full`, `neigh/req = full`, `newton = off`, `binsize` for GPUs = 2x LAMMPS default value, `comm = device`, `neigh/transpose = off`, `gpu/aware = on`. When LAMMPS can safely detect that GPU-aware MPI is not available, the default value of `gpu/aware` becomes “off”. For CPUs or Xeon Phis, the option defaults are `neigh = half`, `neigh/req = half`, `newton = on`, `binsize = 0.0`, and `comm = no`. The option `neigh/thread = on` when there are 16K atoms or less on an MPI rank, otherwise it is “off”. These settings are made automatically by the required “-k on” *command-line switch*. You can change them by using the package `kokkos` command in your input script or via the *-pk kokkos command-line switch*.

For the OMP package, the default is `Nthreads = 0` and the option defaults are `neigh = yes`. These settings are made automatically if the “-sf omp” *command-line switch* is used. If it is not used, you must invoke the package `omp` command in your input script or via the “-pk omp” *command-line switch*.

## 16.96 pair\_coeff command

### 16.96.1 Syntax

```
pair_coeff I J args
```

- I,J = numeric atom types (see asterisk form below), or type labels
- args = coefficients for one or more pairs of atom types

### 16.96.2 Examples

```
pair_coeff 1 2 1.0 1.0 2.5
pair_coeff 2 * 1.0 1.0
pair_coeff 3* 1*2 1.0 1.0 2.5
pair_coeff * * 1.0 1.0
pair_coeff * * nialhjea 1 1 2
pair_coeff * 3 morse.table ENTRY1
pair_coeff 1 2 lj/cut 1.0 1.0 2.5 # (for pair_style hybrid)

labelmap atom 1 C
```

(continues on next page)

```
labelmap atom 2 H
pair_coeff C H 1.0 1.0 2.5
```

### 16.96.3 Description

Specify the pairwise force field coefficients for one or more pairs of atom types. The number and meaning of the coefficients depends on the pair style. Pair coefficients can also be set in the data file read by the *read\_data* command or in a restart file.

I and J can be specified in one of several ways. Explicit numeric values can be used for each, as in the first example above. Or, one or both of the types in the I,J pair can be a type label, which is an alphanumeric string defined by the *labelmap* command or in a section of a data file read by the *read\_data* command, and which converts internally to a numeric type. Internally, LAMMPS will set coefficients for the symmetric J,I interaction to the same values as the I,J interaction.

For numeric values only, a wildcard asterisk can be used in place of or in conjunction with the I,J arguments to set the coefficients for multiple pairs of atom types. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If *N* is the number of atom types, then an asterisk with no numeric values means all types from 1 to *N*. A leading asterisk means all types from 1 to *n* (inclusive). A trailing asterisk means all types from *n* to *N* (inclusive). A middle asterisk means all types from *m* to *n* (inclusive). For the asterisk syntax, only type pairs with  $I \leq J$  are considered; if asterisks imply type pairs where  $J < I$ , they are ignored. Again internally, LAMMPS will set the coefficients for the symmetric J,I interactions to the same values as the  $I \leq J$  interactions.

Note that a *pair\_coeff* command can override a previous setting for the same I,J pair. For example, these commands set the coeffs for all I,J pairs, then overwrite the coeffs for just the I,J = 2,3 pair:

```
pair_coeff * * 1.0 1.0 2.5
pair_coeff 2 3 2.0 1.0 1.12
```

A line in a data file that specifies pair coefficients uses the exact same format as the arguments of the *pair\_coeff* command in an input script, with the exception of the I,J type arguments. In each line of the “Pair Coeffs” section of a data file, only a single type I is specified, which sets the coefficients for type I interacting with type I. This is because the section has exactly *N* lines, where *N* is the number of atom types. For this reason, the wild-card asterisk should also not be used as part of the I argument. Thus in a data file, the line corresponding to the first example above would be listed as

```
2 1.0 1.0 2.5
```

For many potentials, if coefficients for type pairs with  $I \neq J$  are not set explicitly by a *pair\_coeff* command, the values are inferred from the I,I and J,J settings by mixing rules; see the *pair\_modify* command for a discussion. Details on this option as it pertains to individual potentials are described on the page for the potential.

Many pair styles, typically for many-body potentials, use tabulated potential files as input, when specifying the *pair\_coeff* command. Potential files provided with LAMMPS are in the potentials directory of the distribution. For some potentials, such as EAM, other archives of suitable files can be found on the Web. They can be used with LAMMPS so long as they are in the format LAMMPS expects, as discussed on the individual doc pages. The first line of potential files may contain metadata with upper case tags followed their value. These may be parsed and used by LAMMPS. Currently supported are the “DATE:” tag and the UNITS: tag. For pair styles that have been programmed to support the metadata, the value of the “DATE:” tag is printed to the screen and logfile so that the version of a potential file can be later identified. The UNITS: tag indicates the *units* setting required for this particular potential file. If the potential file was created for a different sets of units, LAMMPS will terminate with an error. If the potential file does not contain the tag, no check will be made and it is the responsibility of the user to determine that the unit style is correct.

In some select cases and for specific combinations of unit styles, LAMMPS is capable of automatically converting potential parameters from a file. In those cases, a warning message signaling that an automatic conversion has happened is printed to the screen.

When a `pair_coeff` command using a potential file is specified, LAMMPS looks for the potential file in 2 places. First it looks in the location specified. E.g. if the file is specified as “niu3.eam”, it is looked for in the current working directory. If it is specified as “./potentials/niu3.eam”, then it is looked for in the potentials directory, assuming it is a sister directory of the current working directory. If the file is not found, it is then looked for in one of the directories specified by the `LAMMPS_POTENTIALS` environment variable. Thus if this is set to the potentials directory in the LAMMPS distribution, then you can use those files from anywhere on your system, without copying them into your working directory. Environment variables are set in different ways for different shells. Here are example settings for

csh, tcsh:

```
% setenv LAMMPS_POTENTIALS /path/to/lammps/potentials
```

bash:

```
% export LAMMPS_POTENTIALS=/path/to/lammps/potentials
```

Windows:

```
% set LAMMPS_POTENTIALS="C:\Path to LAMMPS\Potentials"
```

The `LAMMPS_POTENTIALS` environment variable may contain paths to multiple folders, if they are separated by “;” on Windows and “:” on all other operating systems, just like the `PATH` and similar environment variables.

---

The alphabetic list of pair styles defined in LAMMPS is given on the [pair\\_style](#) doc page. They are also listed in more compact form on the [Commands pair](#) doc page.

Click on the style to display the formula it computes and its coefficients as specified by the associated `pair_coeff` command.

---

## 16.96.4 Restrictions

This command must come after the simulation box is defined by a `read_data`, `read_restart`, or `create_box` command.

## 16.96.5 Related commands

[pair\\_style](#), [pair\\_modify](#), [read\\_data](#), [read\\_restart](#), [pair\\_write](#)

## 16.96.6 Default

none

## 16.97 pair\_modify command

### 16.97.1 Syntax

```
pair_modify keyword values ...
```

- one or more keyword/value pairs may be listed
- keyword = *pair* or *shift* or *mix* or *table* or *table/disp* or *tabinner* or *tabinner/disp* or *tail* or *compute* or *nofdotr* or *special* or *compute/tally* or *neigh/trim*

*pair* value = sub-style N

sub-style = sub-style of *pair hybrid*

N = which instance of sub-style (1 to M), only specify if sub-style is used.

↪multiple times

*mix* value = *geometric* or *arithmetic* or *sixthpower*

*shift* value = *yes* or *no*

*table* value = N

2^N = # of values in table

*table/disp* value = N

2^N = # of values in table

*tabinner* value = cutoff

cutoff = inner cutoff at which to begin table (distance units)

*tabinner/disp* value = cutoff

cutoff = inner cutoff at which to begin table (distance units)

*tail* value = *yes* or *no*

*compute* value = *yes* or *no*

*nofdotr* value = *none*

*special* values = which wt1 wt2 wt3

which = *lj/coul* or *lj* or *coul*

w1,w2,w3 = 1-2, 1-3, 1-4 weights from 0.0 to 1.0 inclusive

*compute/tally* value = *yes* or *no*

*neigh/trim* value = *yes* or *no*

### 16.97.2 Examples

```
pair_modify shift yes mix geometric
pair_modify tail yes
pair_modify table 12
pair_modify pair lj/cut compute no
pair_modify pair tersoff compute/tally no
pair_modify pair lj/cut/coul/long 1 special lj/coul 0.0 0.0 0.0
pair_modify pair lj/cut/coul/long special lj 0.0 0.0 0.5 special coul 0.0 0.0 0.8333333
```

### 16.97.3 Description

Modify the parameters of the currently defined pair style. If the pair style is *hybrid or hybrid/overlay*, then the specified parameters are by default modified for all the hybrid sub-styles.

---

**Note:** The behavior for hybrid pair styles can be changed by using the *pair* keyword, which allows selection of a specific sub-style to apply all remaining keywords to. The *special* and *compute/tally* keywords can **only** be used in conjunction with the *pair* keyword. See further details about these 3 keywords below.

---

The *mix* keyword affects pair coefficients for interactions between atoms of type I and J, when  $I \neq J$  and the coefficients are not explicitly set in the input script. Note that coefficients for  $I = J$  must be set explicitly, either in the input script via the *pair\_coeff* command or in the “Pair Coeffs” or “PairIJ Coeffs” sections of the *data file*. For some pair styles it is not necessary to specify coefficients when  $I \neq J$ , since a “mixing” rule will create them from the I,I and J,J settings. The *pair\_modify mix* value determines what formulas are used to compute the mixed coefficients. In each case, the cutoff distance is mixed the same way as sigma.

Note that not all pair styles support mixing and some mix options are not available for certain pair styles. Also, there are additional restrictions when using *pair style hybrid or hybrid/overlay*. See the page for individual pair styles for those restrictions. Note also that the *pair\_coeff* command also can be used to directly set coefficients for a specific  $I \neq J$  pairing, in which case no mixing is performed. If possible, LAMMPS will print an informational message about how many of the mixed pair coefficients were generated and which mixing rule was applied.

- mix *geometric*

$$\begin{aligned}\epsilon_{ij} &= \sqrt{\epsilon_i \epsilon_j} \\ \sigma_{ij} &= \sqrt{\sigma_i \sigma_j}\end{aligned}$$

- mix *arithmetic*

$$\begin{aligned}\epsilon_{ij} &= \sqrt{\epsilon_i \epsilon_j} \\ \sigma_{ij} &= \frac{1}{2}(\sigma_i + \sigma_j)\end{aligned}$$

- mix *sixthpower*

$$\begin{aligned}\epsilon_{ij} &= \frac{2\sqrt{\epsilon_i \epsilon_j} \sigma_i^3 \sigma_j^3}{\sigma_i^6 + \sigma_j^6} \\ \sigma_{ij} &= \left( \frac{1}{2}(\sigma_i^6 + \sigma_j^6) \right)^{\frac{1}{6}}\end{aligned}$$

The *shift* keyword determines whether a Lennard-Jones potential is shifted at its cutoff to 0.0. If so, this adds an energy term to each pairwise interaction which will be included in the thermodynamic output, but does not affect pair forces or atom trajectories. See the doc page for individual pair styles to see which ones support this option.

The *table* and *table/disp* keywords apply to pair styles with a long-range Coulombic term or long-range dispersion term respectively; see the page for individual styles to see which potentials support these options. If N is non-zero, a table of length  $2^N$  is pre-computed for forces and energies, which can shrink their computational cost by up to a factor of 2. The table is indexed via a bit-mapping technique (*Wolff*) and a linear interpolation is performed between adjacent table values. In our experiments with different table styles (lookup, linear, spline), this method typically gave the best performance in terms of speed and accuracy.

The choice of table length is a tradeoff in accuracy versus speed. A larger N yields more accurate force computations, but requires more memory which can slow down the computation due to cache misses. A reasonable value of N is between 8 and 16. The default value of 12 (table of length 4096) gives approximately the same accuracy as the no-table

( $N = 0$ ) option. For  $N = 0$ , forces and energies are computed directly, using a polynomial fit for the needed `erfc()` function evaluation, which is what earlier versions of LAMMPS did. Values greater than 16 typically slow down the simulation and will not improve accuracy; values from 1 to 8 give unreliable results.

The `tabinner` and `tabinner/disp` keywords set an inner cutoff above which the pairwise computation is done by table lookup (if tables are invoked), for the corresponding Coulombic and dispersion tables discussed with the `table` and `table/disp` keywords. The smaller the cutoff is set, the less accurate the table becomes (for a given number of table values), which can require use of larger tables. The default cutoff value is `sqrt(2.0)` distance units which means nearly all pairwise interactions are computed via table lookup for simulations with “real” units, but some close pairs may be computed directly (non-table) for simulations with “lj” units.

When the `tail` keyword is set to `yes`, certain pair styles will add a long-range VanderWaals tail “correction” to the energy and pressure. These corrections are bookkeeping terms which do not affect dynamics, unless a constant-pressure simulation is being performed. See the page for individual styles to see which support this option. These corrections are included in the calculation and printing of thermodynamic quantities (see the `thermo_style` command). Their effect will also be included in constant NPT or NPH simulations where the pressure influences the simulation box dimensions (e.g. the `fix npt` and `fix nph` commands). The formulas used for the long-range corrections come from equation 5 of (*Sun*).

---

**Note:** The tail correction terms are computed at the beginning of each run, using the current atom counts of each atom type. If atoms are deleted (or lost) or created during a simulation, e.g. via the `fix gcmc` command, the correction factors are not re-computed. If you expect the counts to change dramatically, you can break a run into a series of shorter runs so that the correction factors are re-computed more frequently.

---

Several additional assumptions are inherent in using tail corrections, including the following:

- The simulated system is a 3d bulk homogeneous liquid. This option should not be used for systems that are non-liquid, 2d, have a slab geometry (only 2d periodic), or inhomogeneous.
- $G(r)$ , the radial distribution function (rdf), is unity beyond the cutoff, so a fairly large cutoff should be used (i.e. 2.5 sigma for an LJ fluid), and it is probably a good idea to verify this assumption by checking the rdf. The rdf is not exactly unity beyond the cutoff for each pair of interaction types, so the tail correction is necessarily an approximation.

The tail corrections are computed at the beginning of each simulation run. If the number of atoms changes during the run, e.g. due to atoms leaving the simulation domain, or use of the `fix gcmc` command, then the corrections are not updated to reflect the changed atom count. If this is a large effect in your simulation, you should break the long run into several short runs, so that the correction factors are re-computed multiple times.

- Thermophysical properties obtained from calculations with this option enabled will not be thermodynamically consistent with the truncated force-field that was used. In other words, atoms do not feel any LJ pair interactions beyond the cutoff, but the energy and pressure reported by the simulation include an estimated contribution from those interactions.

The `compute` keyword allows pairwise computations to be turned off, even though a `pair_style` is defined. This is not useful for running a real simulation, but can be useful for debugging purposes or for performing a `rerun` simulation, when you only wish to compute partial forces that do not include the pairwise contribution.

Two examples are as follows. First, this option allows you to perform a simulation with `pair_style hybrid` with only a subset of the hybrid sub-styles enabled. Second, this option allows you to perform a simulation with only long-range interactions but no short-range pairwise interactions. Doing this by simply not defining a pair style will not work, because the `kspace_style` command requires a Kspace-compatible pair style be defined.

The `nofdotr` keyword allows to disable an optimization that computes the global stress tensor from the total forces and atom positions rather than from summing forces between individual pairs of atoms.

---



The *pair* keyword can only be used with the *hybrid and hybrid/overlay* pair styles. If used, it must appear first in the list of keywords.

Its meaning is that all the following parameters will only be modified for the specified sub-style. If the sub-style is defined multiple times, then an additional numeric argument *N* must also be specified, which is a number from 1 to *M* where *M* is the number of times the sub-style was listed in the *pair\_style hybrid* command. The extra number indicates which instance of the sub-style the remaining keywords will be applied to.

The *special* and *compute/tally* keywords can **only** be used in conjunction with the *pair* keyword and they must directly follow it. I.e. any other keyword, must appear after *pair*, *special*, and *compute/tally*.

The *special* keyword overrides the global *special\_bonds* 1-2, 1-3, 1-4 exclusion settings (weights) for the sub-style selected by the *pair* keyword.

Similar to the *special\_bonds* command, it takes 4 arguments. The *which* argument can be *lj* to change only the non-Coulomb weights (e.g. Lennard-Jones or Buckingham), *coul* to change only the Coulombic settings, or *lj/coul* to change both to the same values. The *wt1,wt2,wt3* values are numeric weights from 0.0 to 1.0 inclusive, for the 1-2, 1-3, and 1-4 bond topology neighbors, respectively. The *special* keyword can be used multiple times, e.g. to set the *lj* and *coul* settings to different values.

---

**Note:** The *special* keyword is not compatible with pair styles from the GPU or the INTEL package and attempting to use it will cause an error.

---

---

**Note:** Weights of exactly 0.0 or 1.0 in the *special\_bonds* command have implications on the neighbor list construction, which means that they cannot be overridden by using the *special* keyword. One workaround for this restriction is to use the *special\_bonds* command with weights like 1.0e-10 or 0.999999999 instead of 0.0 or 1.0, respectively, which enables to reset each them to any value between 0.0 and 1.0 inclusively. Otherwise you can set **all** global weights to an arbitrary number between 0.0 or 1.0, like 0.5, and then you have to override **all** *special* settings for **all** sub-styles which use the 1-2, 1-3, and 1-4 exclusion weights in their force/energy computation.

---

The *compute/tally* keyword disables or enables registering *compute \*/tally* computes for the sub-style specified by the *pair* keyword. Use *no* to disable, or *yes* to enable.

---

**Note:** The “pair\_modify pair compute/tally” command must be issued **before** the corresponding compute style is defined.

---

The *neigh/trim* keyword controls whether an explicit cutoff is set for each neighbor list request issued by individual pair sub-styles when using *pair hybrid/overlay*. When this keyword is set to *no*, then the cutoff of each pair sub-style neighbor list will be set equal to the largest cutoff, even if a shorter cutoff is specified for a particular sub-style. If possible the neighbor list will be copied directly from another list. When this keyword is set to *yes* then the cutoff of the neighbor list will be explicitly set to the value requested by the pair sub-style, and if possible the list will be created by trimming neighbors from another list with a longer cutoff, otherwise a new neighbor list will be created with the specified cutoff. The *yes* option can be faster when there are multiple pair styles with different cutoffs since the number of pair-wise distance checks between neighbors is reduced (but the time required to build the neighbor lists is increased). The *no* option could be faster when two or more neighbor lists have similar (but not exactly the same) cutoffs.

---

**Note:** The “pair\_modify neigh/trim” command *only* applies when there are multiple pair sub-styles for the same atoms with different cutoffs, i.e. when using pair style *hybrid/overlay*. If you have different cutoffs for different pairs for atoms type, the *neighbor style multi* should be used to create optimized neighbor lists.

---

## 16.97.4 Restrictions

You cannot use *shift* yes with *tail* yes, since those are conflicting options. You cannot use *tail* yes with 2d simulations. You cannot use *special* with pair styles from the GPU or INTEL package.

## 16.97.5 Related commands

*pair\_style*, *pair\_style hybrid*, *pair\_coeff*, *thermo\_style*, *compute \*/tally*, *neighbor multi*

## 16.97.6 Default

The option defaults are mix = geometric, shift = no, table = 12, tabinner = sqrt(2.0), tail = no, compute = yes, and neigh/trim yes.

Note that some pair styles perform mixing, but only a certain style of mixing. See the doc pages for individual pair styles for details.

---

(Wolff) Wolff and Rudd, Comp Phys Comm, 120, 200-32 (1999).

(Sun) Sun, J Phys Chem B, 102, 7338-7364 (1998).

## 16.98 pair\_style command

### 16.98.1 Syntax

```
pair_style style args
```

- style = one of the styles from the list below
- args = arguments used by a particular style

### 16.98.2 Examples

```
pair_style lj/cut 2.5
pair_style eam/alloy
pair_style hybrid lj/charmm/coul/long 10.0 eam
pair_style table linear 1000
pair_style none
```

### 16.98.3 Description

Set the formula(s) LAMMPS uses to compute pairwise interactions. In LAMMPS, pair potentials are defined between pairs of atoms that are within a cutoff distance and the set of active interactions typically changes over time. See the *bond\_style* command to define potentials between pairs of bonded atoms, which typically remain in place for the duration of a simulation.

In LAMMPS, pairwise force fields encompass a variety of interactions, some of which include many-body effects, e.g. EAM, Stillinger-Weber, Tersoff, REBO potentials. They are still classified as “pairwise” potentials because the set of

interacting atoms changes with time (unlike molecular bonds) and thus a neighbor list is used to find nearby interacting atoms.

Hybrid models where specified pairs of atom types interact via different pair potentials can be setup using the *hybrid* pair style.

The coefficients associated with a pair style are typically set for each pair of atom types, and are specified by the *pair\_coeff* command or read from a file by the *read\_data* or *read\_restart* commands.

The *pair\_modify* command sets options for mixing of type I-J interaction coefficients and adding energy offsets or tail corrections to Lennard-Jones potentials. Details on these options as they pertain to individual potentials are described on the doc page for the potential. Likewise, info on whether the potential information is stored in a *restart file* is listed on the potential doc page.

In the formulas listed for each pair style,  $E$  is the energy of a pairwise interaction between two atoms separated by a distance  $r$ . The force between the atoms is the negative derivative of this expression.

If the *pair\_style* command has a cutoff argument, it sets global cutoffs for all pairs of atom types. The distance(s) can be smaller or larger than the dimensions of the simulation box.

In many cases, the global cutoff value can be overridden for a specific pair of atom types by the *pair\_coeff* command.

If a new *pair\_style* command is specified with a new style, all previous *pair\_coeff* and *pair\_modify* command settings are erased; those commands must be re-specified if necessary.

If a new *pair\_style* command is specified with the same style, then only the global settings in that command are reset. Any previous *pair\_coeff* <*pair\_coeff*> and *pair\_modify* command settings are preserved. The only exception is that if the global cutoff in the *pair\_style* command is changed, it will override the corresponding cutoff in any of the previous *pair\_modify* commands.

Two pair styles which do not follow this rule are the *pair\_style table* and *hybrid* commands. A new *pair\_style* command for these styles will wipe out all previously specified *pair\_coeff* and *pair\_modify* settings, including for the sub-styles of the *hybrid* command.

Here is an alphabetic list of pair styles defined in LAMMPS. They are also listed in more compact form on the *Commands pair* doc page.

Click on the style to display the formula it computes, any additional arguments specified in the *pair\_style* command, and coefficients specified by the associated *pair\_coeff* command.

There are also additional accelerated pair styles included in the LAMMPS distribution for faster performance on CPUs, GPUs, and KNLs. The individual style names on the *Commands pair* doc page are followed by one or more of (g,i,k,o,t) to indicate which accelerated styles exist.

- *none* - turn off pairwise interactions
- *hybrid* - multiple styles of pairwise interactions
- *hybrid/overlay* - multiple styles of superposed pairwise interactions
- *hybrid/scaled* - multiple styles of scaled superposed pairwise interactions
- *zero* - neighbor list but no interactions
- *adp* - angular dependent potential (ADP) of Mishin
- *agni* - AGNI machine-learning potential
- *airebo* - AIREBO potential of Stuart
- *airebo/morse* - AIREBO with Morse instead of LJ
- *amoeba* -

- *atm* - Axilrod-Teller-Muto potential
- *awpmd/cut* - Antisymmetrized Wave Packet MD potential for atoms and electrons
- *beck* - Beck potential
- *body/nparticle* - interactions between body particles
- *body/rounded/polygon* - granular-style 2d polygon potential
- *body/rounded/polyhedron* - granular-style 3d polyhedron potential
- *bop* - BOP potential of Pettifor
- *born* - Born-Mayer-Huggins potential
- *born/coul/dsf* - Born with damped-shifted-force model
- *born/coul/dsf/cs* - Born with damped-shifted-force and core/shell model
- *born/coul/long* - Born with long-range Coulomb
- *born/coul/long/cs* - Born with long-range Coulomb and core/shell
- *born/coul/msm* - Born with long-range MSM Coulomb
- *born/coul/wolf* - Born with Wolf potential for Coulomb
- *born/coul/wolf/cs* - Born with Wolf potential for Coulomb and core/shell model
- *bpm/spring* - repulsive harmonic force with damping
- *brownian* - Brownian potential for Fast Lubrication Dynamics
- *brownian/poly* - Brownian potential for Fast Lubrication Dynamics with polydispersity
- *buck* - Buckingham potential
- *buck/coul/cut* - Buckingham with cutoff Coulomb
- *buck/coul/long* - Buckingham with long-range Coulomb
- *buck/coul/long/cs* - Buckingham with long-range Coulomb and core/shell
- *buck/coul/msm* - Buckingham with long-range MSM Coulomb
- *buck/long/coul/long* - long-range Buckingham with long-range Coulomb
- *buck/mdf* - Buckingham with a taper function
- *buck6d/coul/gauss/dsf* - dispersion-damped Buckingham with damped-shift-force model
- *buck6d/coul/gauss/long* - dispersion-damped Buckingham with long-range Coulomb
- *colloid* - integrated colloidal potential
- *comb* - charge-optimized many-body (COMB) potential
- *comb3* - charge-optimized many-body (COMB3) potential
- *cosine/squared* - Cooke-Kremer-Deserno membrane model potential
- *coul/cut* - cutoff Coulomb potential
- *coul/cut/dielectric* -
- *coul/cut/global* - cutoff Coulomb potential
- *coul/cut/soft* - Coulomb potential with a soft core
- *coul/debye* - cutoff Coulomb potential with Debye screening

- *coul/diel* - Coulomb potential with dielectric permittivity
- *coul/dsf* - Coulomb with damped-shifted-force model
- *coul/exclude* - subtract Coulomb potential for excluded pairs
- *coul/long* - long-range Coulomb potential
- *coul/long/cs* - long-range Coulomb potential and core/shell
- *coul/long/dielectric* -
- *coul/long/soft* - long-range Coulomb potential with a soft core
- *coul/msm* - long-range MSM Coulomb
- *coul/slater/cut* - smeared out Coulomb
- *coul/slater/long* - long-range smeared out Coulomb
- *coul/shield* - Coulomb for boron nitride for use with *ilp/graphene/hbn* potential
- *coul/streitz* - Coulomb via Streitz/Mintmire Slater orbitals
- *coul/tt* - damped charge-dipole Coulomb for Drude dipoles
- *coul/wolf* - Coulomb via Wolf potential
- *coul/wolf/cs* - Coulomb via Wolf potential with core/shell adjustments
- *dpd* - dissipative particle dynamics (DPD)
- *dpd/ext* - generalized force field for DPD
- *dpd/ext/tstat* - pairwise DPD thermostating with generalized force field
- *dpd/fdt* - DPD for constant temperature and pressure
- *dpd/fdt/energy* - DPD for constant energy and enthalpy
- *dpd/tstat* - pairwise DPD thermostating
- *dsmc* - Direct Simulation Monte Carlo (DSMC)
- *e3b* - Explicit-three body (E3B) water model
- *drip* - Dihedral-angle-corrected registry-dependent interlayer potential (DRIP)
- *eam* - embedded atom method (EAM)
- *eam/alloy* - alloy EAM
- *eam/cd* - concentration-dependent EAM
- *eam/cd/old* - older two-site model for concentration-dependent EAM
- *eam/fs* - Finnis-Sinclair EAM
- *eam/he* - Finnis-Sinclair EAM modified for Helium in metals
- *edip* - three-body EDIP potential
- *edip/multi* - multi-element EDIP potential
- *edpd* - eDPD particle interactions
- *eff/cut* - electron force field with a cutoff
- *eim* - embedded ion method (EIM)
- *exp6/rx* - reactive DPD potential

- *extep* - extended Tersoff potential
- *gauss* - Gaussian potential
- *gauss/cut* - generalized Gaussian potential
- *gayberne* - Gay-Berne ellipsoidal potential
- *granular* - Generalized granular potential
- *gran/hertz/history* - granular potential with Hertzian interactions
- *gran/hooke* - granular potential with history effects
- *gran/hooke/history* - granular potential without history effects
- *gw* - Gao-Weber potential
- *gw/zbl* - Gao-Weber potential with a repulsive ZBL core
- *harmonic/cut* - repulsive-only harmonic potential
- *hbond/dreiding/lj* - DREIDING hydrogen bonding LJ potential
- *hbond/dreiding/morse* - DREIDING hydrogen bonding Morse potential
- *hdnnp* - High-dimensional neural network potential
- *hippo* -
- *ilp/graphene/hbn* - registry-dependent interlayer potential (ILP)
- *ilp/tmd* - interlayer potential (ILP) potential for transition metal dichalcogenides (TMD)
- *kim* - interface to potentials provided by KIM project
- *kolmogorov/crespi/full* - Kolmogorov-Crespi (KC) potential with no simplifications
- *kolmogorov/crespi/z* - Kolmogorov-Crespi (KC) potential with normals along z-axis
- *lcbop* - long-range bond-order potential (LCBOP)
- *lebedeva/z* - Lebedeva interlayer potential for graphene with normals along z-axis
- *lennard/mdf* - LJ potential in A/B form with a taper function
- *line/lj* - LJ potential between line segments
- *list* - potential between pairs of atoms explicitly listed in an input file
- *lj/charmm/coul/charmm* - CHARMM potential with cutoff Coulomb
- *lj/charmm/coul/charmm/implicit* - CHARMM for implicit solvent
- *lj/charmm/coul/long* - CHARMM with long-range Coulomb
- *lj/charmm/coul/long/soft* - CHARMM with long-range Coulomb and a soft core
- *lj/charmm/coul/msm* - CHARMM with long-range MSM Coulomb
- *lj/charmmfsw/coul/charmmfsh* - CHARMM with force switching and shifting
- *lj/charmmfsw/coul/long* - CHARMM with force switching and long-range Coulomb
- *lj/class2* - COMPASS (class 2) force field without Coulomb
- *lj/class2/coul/cut* - COMPASS with cutoff Coulomb
- *lj/class2/coul/cut/soft* - COMPASS with cutoff Coulomb with a soft core
- *lj/class2/coul/long* - COMPASS with long-range Coulomb

- *lj/class2/coul/long/cs* - COMPASS with long-range Coulomb with core/shell adjustments
- *lj/class2/coul/long/soft* - COMPASS with long-range Coulomb with a soft core
- *lj/class2/soft* - COMPASS (class 2) force field with no Coulomb with a soft core
- *lj/cubic* - LJ with cubic after inflection point
- *lj/cut* - cutoff Lennard-Jones potential without Coulomb
- *lj/cut/coul/cut* - LJ with cutoff Coulomb
- *lj/cut/coul/cut/dielectric* -
- *lj/cut/coul/cut/soft* - LJ with cutoff Coulomb with a soft core
- *lj/cut/coul/debye* - LJ with Debye screening added to Coulomb
- *lj/cut/coul/debye/dielectric* -
- *lj/cut/coul/dsf* - LJ with Coulomb via damped shifted forces
- *lj/cut/coul/long* - LJ with long-range Coulomb
- *lj/cut/coul/long/cs* - LJ with long-range Coulomb with core/shell adjustments
- *lj/cut/coul/long/dielectric* -
- *lj/cut/coul/long/soft* - LJ with long-range Coulomb with a soft core
- *lj/cut/coul/msm* - LJ with long-range MSM Coulomb
- *lj/cut/coul/msm/dielectric* -
- *lj/cut/coul/wolf* - LJ with Coulomb via Wolf potential
- *lj/cut/dipole/cut* - point dipoles with cutoff
- *lj/cut/dipole/long* - point dipoles with long-range Ewald
- *lj/cut/soft* - LJ with a soft core
- *lj/cut/thole/long* - LJ with Coulomb with thole damping
- *lj/cut/tip4p/cut* - LJ with cutoff Coulomb for TIP4P water
- *lj/cut/tip4p/long* - LJ with long-range Coulomb for TIP4P water
- *lj/cut/tip4p/long/soft* - LJ with cutoff Coulomb for TIP4P water with a soft core
- *lj/expand* - Lennard-Jones for variable size particles
- *lj/expand/coul/long* - Lennard-Jones for variable size particles with long-range Coulomb
- *lj/gromacs* - GROMACS-style Lennard-Jones potential
- *lj/gromacs/coul/gromacs* - GROMACS-style LJ and Coulomb potential
- *lj/long/coul/long* - long-range LJ and long-range Coulomb
- *lj/long/coul/long/dielectric* -
- *lj/long/dipole/long* - long-range LJ and long-range point dipoles
- *lj/long/tip4p/long* - long-range LJ and long-range Coulomb for TIP4P water
- *lj/mdf* - LJ potential with a taper function
- *lj/relres* - LJ using multiscale Relative Resolution (RelRes) methodology (*Chaimovich*).
- *lj/spica* - LJ for SPICA coarse-graining

- *lj/spica/coul/long* - LJ for SPICA coarse-graining with long-range Coulomb
- *lj/spica/coul/msm* - LJ for SPICA coarse-graining with long-range Coulomb via MSM
- *lj/sf/dipole/sf* - LJ with dipole interaction with shifted forces
- *lj/smooth* - smoothed Lennard-Jones potential
- *lj/smooth/linear* - linear smoothed LJ potential
- *lj/switch3/coulgauss/long* - smoothed LJ vdW potential with Gaussian electrostatics
- *lj96/cut* - Lennard-Jones 9/6 potential
- *local/density* - generalized basic local density potential
- *lubricate* - hydrodynamic lubrication forces
- *lubricate/poly* - hydrodynamic lubrication forces with polydispersity
- *lubricateU* - hydrodynamic lubrication forces for Fast Lubrication Dynamics
- *lubricateU/poly* - hydrodynamic lubrication forces for Fast Lubrication with polydispersity
- *mdpd* - mDPD particle interactions
- *mdpd/rhosome* - mDPD particle interactions for mass density
- *meam* - modified embedded atom method (MEAM) in C
- *meam/spline* - splined version of MEAM
- *meam/sw/spline* - splined version of MEAM with a Stillinger-Weber term
- *mesocnt* - mesoscopic vdW potential for (carbon) nanotubes
- *mesocnt/viscous* - mesoscopic vdW potential for (carbon) nanotubes with friction
- *mgpt* - simplified model generalized pseudopotential theory (MGPT) potential
- *mesont/tpm* - nanotubes mesoscopic force field
- *mie/cut* - Mie potential
- *mm3/switch3/coulgauss/long* - smoothed MM3 vdW potential with Gaussian electrostatics
- *momb* - Many-Body Metal-Organic (MOMB) force field
- *morse* - Morse potential
- *morse/smooth/linear* - linear smoothed Morse potential
- *morse/soft* - Morse potential with a soft core
- *multi/lucy* - DPD potential with density-dependent force
- *multi/lucy/rx* - reactive DPD potential with density-dependent force
- *nb3b/harmonic* - non-bonded 3-body harmonic potential
- *nm/cut* - N-M potential
- *nm/cut/coul/cut* - N-M potential with cutoff Coulomb
- *nm/cut/coul/long* - N-M potential with long-range Coulomb
- *nm/cut/split* - Split 12-6 Lennard-Jones and N-M potential
- *oxdna/coaxstk* -
- *oxdna/excv* -



- *oxdna/hbond* -
- *oxdna/stk* -
- *oxdna/xstk* -
- *oxdna2/coaxstk* -
- *oxdna2/dh* -
- *oxdna2/excv* -
- *oxdna2/hbond* -
- *oxdna2/stk* -
- *oxdna2/xstk* -
- *oxrna2/coaxstk* -
- *oxrna2/dh* -
- *oxrna2/excv* -
- *oxrna2/hbond* -
- *oxrna2/stk* -
- *oxrna2/xstk* -
- *pace* - Atomic Cluster Expansion (ACE) machine-learning potential
- *peri/eps* - peridynamic EPS potential
- *peri/lps* - peridynamic LPS potential
- *peri/pmb* - peridynamic PMB potential
- *peri/ves* - peridynamic VES potential
- *polymorphic* - polymorphic 3-body potential
- *python* -
- *quip* -
- *rann* -
- *reaxff* - ReaxFF potential
- *rebo* - second generation REBO potential of Brenner
- *resquared* - Everaers RE-Squared ellipsoidal potential
- *saip/metal* - interlayer potential for hetero-junctions formed with hexagonal 2D materials and metal surfaces
- *sdpd/taitwater/isothermal* - smoothed dissipative particle dynamics for water at isothermal conditions
- *smatb* - Second Moment Approximation to the Tight Binding
- *smatb/single* - Second Moment Approximation to the Tight Binding for single-element systems
- *smd/hertz* -
- *smd/tlsph* -
- *smd/tri\_surface* -
- *smd/ulsph* -
- *smtbq* -

- *mliap* - Multiple styles of machine-learning potential
- *snap* - SNAP machine-learning potential
- *soft* - Soft (cosine) potential
- *sph/heatconduction* -
- *sph/idealgas* -
- *sph/lj* -
- *sph/rhosum* -
- *sph/taitwater* -
- *sph/taitwater/morris* -
- *spin/dipole/cut* -
- *spin/dipole/long* -
- *spin/dmi* -
- *spin/exchange* -
- *spin/exchange/biquadratic* -
- *spin/magelec* -
- *spin/neel* -
- *srp* -
- *srp/react* -
- *sw* - Stillinger-Weber 3-body potential
- *sw/angle/table* - Stillinger-Weber potential with tabulated angular term
- *sw/mod* - modified Stillinger-Weber 3-body potential
- *table* - tabulated pair potential
- *table/rx* -
- *tdpd* - tDPD particle interactions
- *tersoff* - Tersoff 3-body potential
- *tersoff/mod* - modified Tersoff 3-body potential
- *tersoff/mod/c* -
- *tersoff/table* -
- *tersoff/zbl* - Tersoff/ZBL 3-body potential
- *thole* - Coulomb interactions with thole damping
- *threebody/table* - generic tabulated three-body potential
- *tip4p/cut* - Coulomb for TIP4P water w/out LJ
- *tip4p/long* - long-range Coulomb for TIP4P water w/out LJ
- *tip4p/long/soft* -
- *tracker* - monitor information about pairwise interactions
- *tri/lj* - LJ potential between triangles

- *ufm* -
- *vashishta* - Vashishta 2-body and 3-body potential
- *vashishta/table* -
- *wf/cut* - Wang-Frenkel Potential for short-ranged interactions
- *yukawa* - Yukawa potential
- *yukawa/colloid* - screened Yukawa potential for finite-size particles
- *zbl* - Ziegler-Biersack-Littmark potential

## 16.98.4 Restrictions

This command must be used before any coefficients are set by the *pair\_coeff*, *read\_data*, or *read\_restart* commands.

Some pair styles are part of specific packages. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info. The doc pages for individual pair potentials tell if it is part of a package.

## 16.98.5 Related commands

*pair\_coeff*, *read\_data*, *pair\_modify*, *kpace\_style*, *dielectric*, *pair\_write*

## 16.98.6 Default

```
pair_style none
```

## 16.99 pair\_write command

### 16.99.1 Syntax

```
pair_write itype jtype N style inner outer file keyword Qi Qj
```

- *itype,jtype* = 2 atom types
- *N* = # of values
- *style* = *r* or *rsq* or *bitmap*
- *inner,outer* = inner and outer cutoff (distance units)
- *file* = name of file to write values to
- *keyword* = section name in file for this set of tabulated values
- *Qi,Qj* = 2 atom charges (charge units) (optional)

## 16.99.2 Examples

```
pair_write 1 3 500 r 1.0 10.0 table.txt LJ
pair_write 1 1 1000 rsq 2.0 8.0 table.txt Yukawa_1_1 -0.5 0.5
```

## 16.99.3 Description

Write energy and force values to a file as a function of distance for the currently defined pair potential. This is useful for plotting the potential function or otherwise debugging its values. If the file already exists, the table of values is appended to the end of the file to allow multiple tables of energy and force to be included in one file. In case a new file is created, the first line will be a comment containing a “DATE:” and “UNITS:” tag with the current date and the current *units* setting as argument. For subsequent invocations of the `pair_write` command, the current units setting is compared against the entry in the file, if present, and `pair_write` will refuse to add a table if the units are not the same.

The energy and force values are computed at distances from inner to outer for 2 interacting atoms of type *itype* and *jtype*, using the appropriate *pair\_coeff* coefficients. If the style is *r*, then *N* distances are used, evenly spaced in *r*; if the style is *rsq*, *N* distances are used, evenly spaced in  $r^2$ .

For example, for *N* = 7, style = *r*, inner = 1.0, and outer = 4.0, values are computed at  $r = 1.0, 1.5, 2.0, 2.5, 3.0, 3.5, 4.0$ .

If the style is *bitmap*, then  $2^N$  values are written to the file in a format and order consistent with how they are read in by the *pair\_coeff* command for pair style *table*. For reasonable accuracy in a bitmapped table, choose  $N \geq 12$ , an *inner* value that is smaller than the distance of closest approach of 2 atoms, and an *outer* value  $\leq$  cutoff of the potential.

If the pair potential is computed between charged atoms, the charges of the pair of interacting atoms can optionally be specified. If not specified, values of  $Q_i = Q_j = 1.0$  are used.

The file is written in the format used as input for the *pair\_style table* option with *keyword* as the section name. Each line written to the file lists an index number (1-*N*), a distance (in distance units), an energy (in energy units), and a force (in force units).

## 16.99.4 Restrictions

All force field coefficients for pair and other kinds of interactions must be set before this command can be invoked.

Due to how the pairwise force is computed, an inner value  $> 0.0$  must be specified even if the potential has a finite value at  $r = 0.0$ .

The *pair\_write* command can only be used for pairwise additive interactions for which a *Pair::single()* function can be and has been implemented. This excludes for example manybody potentials or TIP4P coulomb styles.

## 16.99.5 Related commands

*pair\_style table*, *pair\_style*, *pair\_coeff*

## 16.99.6 Default

none

## 16.100 partition command

### 16.100.1 Syntax

```
partition style N command ...
```

- style = *yes* or *no*
- N = partition number (see asterisk form below)
- command = any LAMMPS command

### 16.100.2 Examples

```
partition yes 1 processors 4 10 6
partition no 5 print "Active partition"
partition yes *5 fix all nve
partition yes 6* fix all nvt temp 1.0 1.0 0.1
```

### 16.100.3 Description

This command invokes the specified command on a subset of the partitions of processors you have defined via the *-partition command-line switch*.

Normally, every input script command in your script is invoked by every partition. This behavior can be modified by defining world- or universe-style *variables* that have different values for each partition. This mechanism can be used to cause your script to jump to different input script files on different partitions, if such a variable is used in a *jump* command.

The “partition” command is another mechanism for having an input script operate differently on different partitions. It is basically a prefix on any LAMMPS command. The command will only be invoked on the partition(s) specified by the *style* and *N* arguments.

If the *style* is *yes*, the command will be invoked on any partition which matches the *N* argument. If the *style* is *no* the command will be invoked on all the partitions which do not match the *Np* argument.

Partitions are numbered from 1 to *Np*, where *Np* is the number of partitions specified by the *-partition command-line switch*.

*N* can be specified in one of two ways. An explicit numeric value can be used, as in the first example above. Or a wild-card asterisk can be used to span a range of partition numbers. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. An asterisk with no numeric values means all partitions from 1 to *Np*. A leading asterisk means all partitions from 1 to *n* (inclusive). A trailing asterisk means all partitions from *n* to *Np* (inclusive). A middle asterisk means all partitions from *m* to *n* (inclusive).

This command can be useful for the “run\_style verlet/split” command which imposed requirements on how the *processors* command lays out a 3d grid of processors in each of 2 partitions.

## 16.100.4 Restrictions

none

## 16.100.5 Related commands

*run\_style verlet/split*

## 16.100.6 Default

none

# 16.101 plugin command

## 16.101.1 Syntax

```
plugin command args
```

- *command* = *load* or *unload* or *list* or *clear*
- *args* = list of arguments for a particular plugin command

*load file* = load plugin(s) from shared object in *file*

*unload style name* = unload plugin *name* of style *style*

*style* = *pair* or *bond* or *angle* or *dihedral* or *improper* or *kspace* or *compute* or *fix* or *region* or *command*

*list* = print a list of currently loaded plugins

*clear* = unload all currently loaded plugins

## 16.101.2 Examples

```
plugin load morse2plugin.so
plugin unload pair morse2/omp
plugin unload command hello
plugin list
plugin clear
```

## 16.101.3 Description

The `plugin` command allows to load (and unload) additional styles and commands into a LAMMPS binary from so-called dynamic shared object (DSO) files. This enables to add new functionality to an existing LAMMPS binary without having to recompile and link the entire executable.

The `load` command will load and initialize all plugins contained in the plugin DSO with the given filename. A message with information the plugin style and name and more will be printed. Individual DSO files may contain multiple plugins. More details about how to write and compile the plugin DSO is given in programmer's guide part of the manual under *Writing plugins*.

The *unload* command will remove the given style or the given name from the list of available styles. If the plugin style is currently in use, that style instance will be deleted.

The *list* command will print a list of the loaded plugins and their styles and names.

The *clear* command will unload all currently loaded plugins.

---

### Automatic loading of plugins

When the environment variable `LAMMPS_PLUGIN_PATH` is set, then LAMMPS will search the directory (or directories) listed in this path for files with names that end in `plugin.so` (e.g. `helloplugin.so`) and will try to load the contained plugins automatically at start-up.

---

## 16.101.4 Restrictions

The *plugin* command is part of the `PLUGIN` package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

If plugins access functions or classes from a package, LAMMPS must have been compiled with that package included.

Plugins are dependent on the LAMMPS binary interface (ABI) and particularly the MPI library used. So they are not guaranteed to work when the plugin was compiled with a different MPI library or different compilation settings or a different LAMMPS version. There are no checks, so if there is a mismatch the plugin object will either not load or data corruption and crashes may happen.

## 16.101.5 Related commands

none

## 16.101.6 Default

none

## 16.102 prd command

### 16.102.1 Syntax

```
prd N t_event n_dephase t_dephase t_correlate compute-ID seed keyword value ...
```

- `N` = # of timesteps to run (not including dephasing/quenching)
- `t_event` = timestep interval between event checks
- `n_dephase` = number of velocity randomizations to perform in each dephase run
- `t_dephase` = number of timesteps to run dynamics after each velocity randomization during dephase
- `t_correlate` = number of timesteps within which 2 consecutive events are considered to be correlated
- `compute-ID` = ID of the compute used for event detection
- `random_seed` = random # seed (positive integer)
- zero or more keyword/value pairs may be appended

- keyword = *min* or *temp* or *vel*

```
min values = etol ftol maxiter maxeval
  etol = stopping tolerance for energy, used in quenching
  ftol = stopping tolerance for force, used in quenching
  maxiter = max iterations of minimize, used in quenching
  maxeval = max number of force/energy evaluations, used in quenching
temp value = Tdephase
  Tdephase = target temperature for velocity randomization, used in dephasing
vel values = loop dist
  loop = all or local or geom, used in dephasing
  dist = uniform or gaussian, used in dephasing
time value = steps or clock
  steps = simulation runs for N timesteps on each replica (default)
  clock = simulation runs for N timesteps across all replicas
```

## 16.102.2 Examples

```
prd 5000 100 10 10 100 1 54982
prd 5000 100 10 10 100 1 54982 min 0.1 0.1 100 200
```

## 16.102.3 Description

Run a parallel replica dynamics (PRD) simulation using multiple replicas of a system. One or more replicas can be used. The total number of steps  $N$  to run can be interpreted in one of two ways; see discussion of the *time* keyword below.

PRD is described in ([Voter1998](#)) by Art Voter. Similar to global or local hyperdynamics (HD), PRD is a method for performing accelerated dynamics that is suitable for infrequent-event systems that obey first-order kinetics. A good overview of accelerated dynamics methods (AMD) for such systems is given in this review paper ([Voter2002](#)) from Art's group. To quote from the paper: "The dynamical evolution is characterized by vibrational excursions within a potential basin, punctuated by occasional transitions between basins. The transition probability is characterized by  $p(t) = k \cdot \exp(-kt)$  where  $k$  is the rate constant."

Both PRD and HD produce a time-accurate trajectory that effectively extends the timescale over which a system can be simulated, but they do it differently. PRD creates  $N_r$  replicas of the system and runs dynamics on each independently with a normal unbiased potential until an event occurs in one of the replicas. The time between events is reduced by a factor of  $N_r$  replicas. HD uses a single replica of the system and accelerates time by biasing the interaction potential in a manner such that each timestep is effectively longer. For both methods, per CPU second, more physical time elapses and more events occur. See the [hyper](#) page for more info about HD.

In PRD, each replica runs on a partition of one or more processors. Processor partitions are defined at run-time using the *-partition command-line switch*. Note that if you have MPI installed, you can run a multi-replica simulation with more replicas (partitions) than you have physical processors, e.g you can run a 10-replica simulation on one or two processors. However for PRD, this makes little sense, since running a replica on virtual instead of physical processors, offers no effective parallel speed-up in searching for infrequent events. See the [Howto replica](#) doc page for further discussion.

When a PRD simulation is performed, it is assumed that each replica is running the same model, though LAMMPS does not check for this. I.e. the simulation domain, the number of atoms, the interaction potentials, etc should be the same for every replica.

A PRD run has several stages, which are repeated each time an "event" occurs in one of the replicas, as explained below. The logic for a PRD run is as follows:



```

while (time remains):
  dephase for n_dephase*t_dephase steps
  until (event occurs on some replica):
    run dynamics for t_event steps
    quench
    check for uncorrelated event on any replica
  until (no correlated event occurs):
    run dynamics for t_correlate steps
    quench
    check for correlated event on this replica
  event replica shares state with all replicas

```

Before this loop begins, the state of the system on replica 0 is shared with all replicas, so that all replicas begin from the same initial state. The first potential energy basin is identified by quenching (an energy minimization, see below) the initial state and storing the resulting coordinates for reference.

In the first stage, dephasing is performed by each replica independently to eliminate correlations between replicas. This is done by choosing a random set of velocities, based on the *random\_seed* that is specified, and running *t\_dephase* timesteps of dynamics. This is repeated *n\_dephase* times. At each of the *n\_dephase* stages, if an event occurs during the *t\_dephase* steps of dynamics for a particular replica, the replica repeats the stage until no event occurs.

If the *temp* keyword is not specified, the target temperature for velocity randomization for each replica is the current temperature of that replica. Otherwise, it is the specified *Tdephase* temperature. The style of velocity randomization is controlled using the keyword *vel* with arguments that have the same meaning as their counterparts in the *velocity* command.

In the second stage, each replica runs dynamics continuously, stopping every *t\_event* steps to check if a transition event has occurred. This check is performed by quenching the system and comparing the resulting atom coordinates to the coordinates from the previous basin. The first time through the PRD loop, the “previous basin” is the set of quenched coordinates from the initial state of the system.

A quench is an energy minimization and is performed by whichever algorithm has been defined by the *min\_style* command. Minimization parameters may be set via the *min\_modify* command and by the *min* keyword of the PRD command. The latter are the settings that would be used with the *minimize* command. Note that typically, you do not need to perform a highly-converged minimization to detect a transition event, though you may need to in order to prevent a set of atoms in the system from relaxing to a saddle point.

The event check is performed by a compute with the specified *compute-ID*. Currently there is only one compute that works with the PRD command, which is the *compute event/displace* command. Other event-checking computes may be added. *Compute event/displace* checks whether any atom in the compute group has moved further than a specified threshold distance. If so, an “event” has occurred.

In the third stage, the replica on which the event occurred (event replica) continues to run dynamics to search for correlated events. This is done by running dynamics for *t\_correlate* steps, quenching every *t\_event* steps, and checking if another event has occurred.

The first time no correlated event occurs, the final state of the event replica is shared with all replicas, the new basin reference coordinates are updated with the quenched state, and the outer loop begins again. While the replica event is searching for correlated events, all the other replicas also run dynamics and event checking with the same schedule, but the final states are always overwritten by the state of the event replica.

The outer loop of the pseudo-code above continues until *N* steps of dynamics have been performed. Note that *N* only includes the dynamics of stages 2 and 3, not the steps taken during dephasing or the minimization iterations of quenching. The specified *N* is interpreted in one of two ways, depending on the *time* keyword. If the *time* value is *steps*, which is the default, then each replica runs for *N* timesteps. If the *time* value is *clock*, then the simulation runs until *N* aggregate timesteps across all replicas have elapsed. This aggregate time is the “clock” time defined below, which typically advances nearly *M* times faster than the timestepping on a single replica, where *M* is the number of replicas.

Four kinds of output can be generated during a PRD run: event statistics, thermodynamic output by each replica, dump files, and restart files.

When running with multiple partitions (each of which is a replica in this case), the print-out to the screen and master log.lammps file is limited to event statistics. Note that if a PRD run is performed on only a single replica then the event statistics will be intermixed with the usual thermodynamic output discussed below.

The quantities printed each time an event occurs are the timestep, CPU time, clock, event number, a correlation flag, the number of coincident events, and the replica number of the chosen event.

The timestep is the usual LAMMPS timestep, except that time does not advance during dephasing or quenches, but only during dynamics. Note that there are two kinds of dynamics in the PRD loop listed above that contribute to this timestepping. The first is when all replicas are performing independent dynamics, waiting for an event to occur. The second is when correlated events are being searched for, but only one replica is running dynamics.

The CPU time is the total elapsed time on each processor, since the start of the PRD run.

The clock is the same as the timestep except that it advances by  $M$  steps per timestep during the first kind of dynamics when the  $M$  replicas are running independently. The clock advances by only 1 step per timestep during the second kind of dynamics, when only a single replica is checking for a correlated event. Thus “clock” time represents the aggregate time (in steps) that has effectively elapsed during a PRD simulation on  $M$  replicas. If most of the PRD run is spent in the second stage of the loop above, searching for infrequent events, then the clock will advance nearly  $M$  times faster than it would if a single replica was running. Note the clock time between successive events should be drawn from  $p(t)$ .

The event number is a counter that increments with each event, whether it is uncorrelated or correlated.

The correlation flag will be 0 when an uncorrelated event occurs during the second stage of the loop listed above, i.e. when all replicas are running independently. The correlation flag will be 1 when a correlated event occurs during the third stage of the loop listed above, i.e. when only one replica is running dynamics.

When more than one replica detects an event at the end of the same event check (every  $t_{event}$  steps) during the second stage, then one of them is chosen at random. The number of coincident events is the number of replicas that detected an event. Normally, this value should be 1. If it is often greater than 1, then either the number of replicas is too large, or  $t_{event}$  is too large.

The replica number is the ID of the replica (from 0 to  $M-1$ ) in which the event occurred.

---

When running on multiple partitions, LAMMPS produces additional log files for each partition, e.g. log.lammps.0, log.lammps.1, etc. For the PRD command, these contain the thermodynamic output for each replica. You will see short runs and minimizations corresponding to the dynamics and quench operations of the loop listed above. The timestep will be reset appropriately depending on whether the operation advances time or not.

After the PRD command completes, timing statistics for the PRD run are printed in each replica’s log file, giving a breakdown of how much CPU time was spent in each stage (dephasing, dynamics, quenching, etc).

---

Any *dump files* defined in the input script, will be written to during a PRD run at timesteps corresponding to both uncorrelated and correlated events. This means the requested dump frequency in the *dump* command is ignored. There will be one dump file (per dump command) created for all partitions.

The atom coordinates of the dump snapshot are those of the minimum energy configuration resulting from quenching following a transition event. The timesteps written into the dump files correspond to the timestep at which the event occurred and NOT the clock. A dump snapshot corresponding to the initial minimum state used for event detection is written to the dump file at the beginning of each PRD run.

---

If the *restart* command is used, a single restart file for all the partitions is generated, which allows a PRD run to be continued by a new input script in the usual manner.

---

The restart file is generated at the end of the loop listed above. If no correlated events are found, this means it contains a snapshot of the system at time  $T + t\_correlate$ , where  $T$  is the time at which the uncorrelated event occurred. If correlated events were found, then it contains a snapshot of the system at time  $T + t\_correlate$ , where  $T$  is the time of the last correlated event.

The restart frequency specified in the *restart* command is interpreted differently when performing a PRD run. It does not mean the timestep interval between restart files. Instead it means an event interval for uncorrelated events. Thus a frequency of 1 means write a restart file every time an uncorrelated event occurs. A frequency of 10 means write a restart file every 10th uncorrelated event.

When an input script reads a restart file from a previous PRD run, the new script can be run on a different number of replicas or processors. However, it is assumed that  $t\_correlate$  in the new PRD command is the same as it was previously. If not, the calculation of the “clock” value for the first event in the new run will be slightly off.

---

### 16.102.4 Restrictions

This command can only be used if LAMMPS was built with the REPLICCA package. See the *Build package* doc page for more info.

The  $N$  and  $t\_correlate$  settings must be integer multiples of  $t\_event$ .

Runs restarted from restart file written during a PRD run will not produce identical results due to changes in the random numbers used for dephasing.

This command cannot be used when any fixes are defined that keep track of elapsed time to perform time-dependent operations. Examples include the “ave” fixes such as *fix ave/chunk*. Also *fix dt/reset* and *fix deposit*.

### 16.102.5 Related commands

*compute event/displace, min\_modify, min\_style, run\_style, minimize, velocity, temper, neb, tad, hyper*

### 16.102.6 Default

The option defaults are `min = 0.1 0.1 40 50`, no temp setting, `vel = geom gaussian`, and `time = steps`.

---

(Voter1998) Voter, Phys Rev B, 57, 13985 (1998).

(Voter2002) Voter, Montalenti, Germann, Annual Review of Materials Research 32, 321 (2002).

## 16.103 print command

### 16.103.1 Syntax

```
print string keyword value
```

- string = text string to print, which may contain variables
- zero or more keyword/value pairs may be appended
- keyword = *file* or *append* or *screen* or *universe*

```
file value = filename
append value = filename
screen value = yes or no
universe value = yes or no
```

### 16.103.2 Examples

```
print "Done with equilibration" file info.dat
print Vol=$v append info.dat screen no
print "The system volume is now $v"
print 'The system volume is now $v'
print "NEB calculation 1 complete" screen no universe yes
print ""
System volume = $v
System temperature = $t
""
```

### 16.103.3 Description

Print a text string to the screen and logfile. The text string must be a single argument, so if it is one line but more than one word, it should be enclosed in single or double quotes. To generate multiple lines of output, the string can be enclosed in triple quotes, as in the last example above. If the text string contains variables, they will be evaluated and their current values printed.

---

**Note:** As discussed on the [Commands parse](#) doc page, the text string can use “immediate” variables, specified as  $$(formula)$  with parenthesis, where the numeric formula has the same syntax as equal-style variables described on the [variable](#) doc page. This is a convenient way to evaluate a formula immediately without using the variable command to define a named variable and then use that variable in the text string. The formula can include a trailing colon and format string which determines the precision with which the numeric value is output. This is also explained on the [Commands parse](#) doc page.

---

If the *file* or *append* keyword is used, a filename is specified to which the output will be written. If *file* is used, then the filename is overwritten if it already exists. If *append* is used, then the filename is appended to if it already exists, or created if it does not exist.

If the *screen* keyword is used, output to the screen and logfile can be turned on or off as desired.

If the *universe* keyword is used, output to the global screen and logfile can be turned on or off as desired. In multi-partition calculations, the *screen* option and the corresponding output only apply to the screen and logfile of the individual partition.

If you want the print command to be executed multiple times (with changing variable values), there are 3 options. First, consider using the *fix print* command, which will print a string periodically during a simulation. Second, the print command can be used as an argument to the *every* option of the *run* command. Third, the print command could appear in a section of the input script that is looped over (see the *jump* and *next* commands).

See the [variable](#) command for a description of *equal* style variables which are typically the most useful ones to use with the print command. Equal-style variables can calculate formulas involving mathematical operations, atom properties, group properties, thermodynamic properties, global values calculated by a *compute* or *fix*, or references to other *variables*.

## 16.103.4 Restrictions

none

## 16.103.5 Related commands

*fix print, variable*

## 16.103.6 Default

The option defaults are no file output, screen = yes, and universe = no.

# 16.104 processors command

## 16.104.1 Syntax

```
processors Px Py Pz keyword args ...
```

- Px,Py,Pz = # of processors in each dimension of 3d grid overlaying the simulation domain
- zero or more keyword/arg pairs may be appended
- keyword = *grid* or *map* or *part* or *file*

```

grid arg = gstyle params ...
  gstyle = onelevel or twolevel or numa or custom
  onelevel params = none
  twolevel params = Nc Cx Cy Cz
    Nc = number of cores per node
    Cx,Cy,Cz = # of cores in each dimension of 3d sub-grid assigned to each node
  numa params = none
  custom params = infile
    infile = file containing grid layout
map arg = cart or cart/reorder or xyz or xzy or yxz or yzx or zxy or zyx
  cart = use MPI_Cart() methods to map processors to 3d grid with reorder = 0
  cart/reorder = use MPI_Cart() methods to map processors to 3d grid with reorder_
↳= 1
  xyz,xzy,yxz,yzx,zxy,zyx = map processors to 3d grid in IJK ordering
numa arg = none
part args = Psend Precv cstyle
  Psend = partition # (1 to Np) which will send its processor layout
  Precv = partition # (1 to Np) which will recv the processor layout
  cstyle = multiple
    multiple = Psend grid will be multiple of Precv grid in each dimension
file arg = outfile
  outfile = name of file to write 3d grid of processors to

```

## 16.104.2 Examples

```
processors * * 5
processors 2 4 4
processors * * 8 map xyz
processors * * * grid numa
processors * * * grid twolevel 4 * * 1
processors 4 8 16 grid custom myfile
processors * * * part 1 2 multiple
```

## 16.104.3 Description

Specify how processors are mapped as a regular 3d grid to the global simulation box. The mapping involves 2 steps. First if there are  $P$  processors it means choosing a factorization  $P = P_x$  by  $P_y$  by  $P_z$  so that there are  $P_x$  processors in the  $x$  dimension, and similarly for the  $y$  and  $z$  dimensions. Second, the  $P$  processors are mapped to the regular 3d grid. The arguments to this command control each of these 2 steps.

The  $P_x$ ,  $P_y$ ,  $P_z$  parameters affect the factorization. Any of the 3 parameters can be specified with an asterisk “\*”, which means LAMMPS will choose the number of processors in that dimension of the grid. It will do this based on the size and shape of the global simulation box so as to minimize the surface-to-volume ratio of each processor’s sub-domain.

Choosing explicit values for  $P_x$  or  $P_y$  or  $P_z$  can be used to override the default manner in which LAMMPS will create the regular 3d grid of processors, if it is known to be sub-optimal for a particular problem. E.g. a problem where the extent of atoms will change dramatically in a particular dimension over the course of the simulation.

The product of  $P_x$ ,  $P_y$ ,  $P_z$  must equal  $P$ , the total # of processors LAMMPS is running on. For a *2d simulation*,  $P_z$  must equal 1.

Note that if you run on a prime number of processors  $P$ , then a grid such as  $1 \times P \times 1$  will be required, which may incur extra communication costs due to the high surface area of each processor’s sub-domain.

Also note that if multiple partitions are being used then  $P$  is the number of processors in this partition; see the *partition command-line switch* page for details. Also note that you can prefix the processors command with the *partition* command to easily specify different  $P_x, P_y, P_z$  values for different partitions.

You can use the *partition* command to specify different processor grids for different partitions, e.g.

```
partition yes 1 processors 4 4 4
partition yes 2 processors 2 3 2
```

---

**Note:** This command only affects the initial regular 3d grid created when the simulation box is first specified via a *create\_box* or *read\_data* or *read\_restart* command. Or if the simulation box is re-created via the *replicate* command. The same regular grid is initially created, regardless of which *comm\_style* command is in effect.

---

If load-balancing is never invoked via the *balance* or *fix balance* commands, then the initial regular grid will persist for all simulations. If balancing is performed, some of the methods invoked by those commands retain the logical topology of the initial 3d grid, and the mapping of processors to the grid specified by the processors command. However the grid spacings in different dimensions may change, so that processors own sub-domains of different sizes. If the *comm\_style tiled* command is used, methods invoked by the balancing commands may discard the 3d grid of processors and tile the simulation domain with sub-domains of different sizes and shapes which no longer have a logical 3d connectivity. If that occurs, all the information specified by the processors command is ignored.

---

The *grid* keyword affects the factorization of  $P$  into  $P_x, P_y, P_z$  and it can also affect how the  $P$  processor IDs are mapped to the 3d grid of processors.

The *onelevel* style creates a 3d grid that is compatible with the  $P_x, P_y, P_z$  settings, and which minimizes the surface-to-volume ratio of each processor's sub-domain, as described above. The mapping of processors to the grid is determined by the *map* keyword setting.

The *twolevel* style can be used on machines with multicore nodes to minimize off-node communication. It insures that contiguous sub-sections of the 3d grid are assigned to all the cores of a node. For example if  $N_c$  is 4, then  $2 \times 2 \times 1$  or  $2 \times 1 \times 2$  or  $1 \times 2 \times 2$  sub-sections of the 3d grid will correspond to the cores of each node. This affects both the factorization and mapping steps.

The  $C_x, C_y, C_z$  settings are similar to the  $P_x, P_y, P_z$  settings, only their product should equal  $N_c$ . Any of the 3 parameters can be specified with an asterisk "\*", which means LAMMPS will choose the number of cores in that dimension of the node's sub-grid. As with  $P_x, P_y, P_z$ , it will do this based on the size and shape of the global simulation box so as to minimize the surface-to-volume ratio of each processor's sub-domain.

---

**Note:** For the *twolevel* style to work correctly, it assumes the MPI ranks of processors LAMMPS is running on are ordered by core and then by node. E.g. if you are running on 2 quad-core nodes, for a total of 8 processors, then it assumes processors 0,1,2,3 are on node 1, and processors 4,5,6,7 are on node 2. This is the default rank ordering for most MPI implementations, but some MPIs provide options for this ordering, e.g. via environment variable settings.

---

The *numa* style operates similar to the *twolevel* keyword except that it auto-detects which cores are running on which nodes. Currently, it does this in only 2 levels, but it may be extended in the future to account for socket topology and other non-uniform memory access (NUMA) costs. It also uses a different algorithm than the *twolevel* keyword for doing the two-level factorization of the simulation box into a 3d processor grid to minimize off-node communication, and it does its own MPI-based mapping of nodes and cores to the regular 3d grid. Thus it may produce a different layout of the processors than the *twolevel* options.

The *numa* style will give an error if the number of MPI processes is not divisible by the number of cores used per node, or any of the  $P_x$  or  $P_y$  or  $P_z$  values is greater than 1.

---

**Note:** Unlike the *twolevel* style, the *numa* style does not require any particular ordering of MPI ranks in order to work correctly. This is because it auto-detects which processes are running on which nodes.

---

The *custom* style uses the file *infile* to define both the 3d factorization and the mapping of processors to the grid.

The file should have the following format. Any number of initial blank or comment lines (starting with a "#" character) can be present. The first non-blank, non-comment line should have 3 values:

```
Px Py Pz
```

These must be compatible with the total number of processors and the  $P_x, P_y, P_z$  settings of the processors command.

This line should be immediately followed by  $P = P_x * P_y * P_z$  lines of the form:

```
ID I J K
```

where ID is a processor ID (from 0 to  $P-1$ ) and I,J,K are the processors location in the 3d grid. I must be a number from 1 to  $P_x$  (inclusive) and similarly for J and K. The  $P$  lines can be listed in any order, but no processor ID should appear more than once.

---

The *map* keyword affects how the  $P$  processor IDs (from 0 to  $P-1$ ) are mapped to the 3d grid of processors. It is only used by the *onelevel* and *twolevel* grid settings.

The *cart* style uses the family of MPI Cartesian functions to perform the mapping, namely `MPI_Cart_create()`, `MPI_Cart_get()`, `MPI_Cart_shift()`, and `MPI_Cart_rank()`. It invokes the `MPI_Cart_create()` function with its reorder flag = 0, so that MPI is not free to reorder the processors.

The *cart/reorder* style does the same thing as the *cart* style except it sets the reorder flag to 1, so that MPI can reorder processors if it desires.

The *xyz*, *xzy*, *yxz*, *yzx*, *zxy*, and *zyx* styles are all similar. If the style is IJK, then it maps the P processors to the grid so that the processor ID in the I direction varies fastest, the processor ID in the J direction varies next fastest, and the processor ID in the K direction varies slowest. For example, if you select style *xyz* and you have a 2x2x2 grid of 8 processors, the assignments of the 8 octants of the simulation domain will be:

```
proc 0 = lo x, lo y, lo z octant
proc 1 = hi x, lo y, lo z octant
proc 2 = lo x, hi y, lo z octant
proc 3 = hi x, hi y, lo z octant
proc 4 = lo x, lo y, hi z octant
proc 5 = hi x, lo y, hi z octant
proc 6 = lo x, hi y, hi z octant
proc 7 = hi x, hi y, hi z octant
```

Note that, in principle, an MPI implementation on a particular machine should be aware of both the machine's network topology and the specific subset of processors and nodes that were assigned to your simulation. Thus its `MPI_Cart` calls can optimize the assignment of MPI processes to the 3d grid to minimize communication costs. In practice, however, few if any MPI implementations actually do this. So it is likely that the *cart* and *cart/reorder* styles simply give the same result as one of the IJK styles.

Also note, that for the *twolevel* grid style, the *map* setting is used to first map the nodes to the 3d grid, then again to the cores within each node. For the latter step, the *cart* and *cart/reorder* styles are not supported, so an *xyz* style is used in their place.

---

The *part* keyword affects the factorization of P into  $P_x, P_y, P_z$ .

It can be useful when running in multi-partition mode, e.g. with the *run\_style verlet/split* command. It specifies a dependency between a sending partition *Psend* and a receiving partition *Precv* which is enforced when each is setting up their own mapping of their processors to the simulation box. Each of *Psend* and *Precv* must be integers from 1 to  $N_p$ , where  $N_p$  is the number of partitions you have defined via the *-partition command-line switch*.

A “dependency” means that the sending partition will create its regular 3d grid as  $P_x$  by  $P_y$  by  $P_z$  and after it has done this, it will send the  $P_x, P_y, P_z$  values to the receiving partition. The receiving partition will wait to receive these values before creating its own regular 3d grid and will use the sender's  $P_x, P_y, P_z$  values as a constraint. The nature of the constraint is determined by the *cstyle* argument.

For a *cstyle* of *multiple*, each dimension of the sender's processor grid is required to be an integer multiple of the corresponding dimension in the receiver's processor grid. This is a requirement of the *run\_style verlet/split* command.

For example, assume the sending partition creates a 4x6x10 grid = 240 processor grid. If the receiving partition is running on 80 processors, it could create a 4x2x10 grid, but it will not create a 2x4x10 grid, since in the y-dimension, 6 is not an integer multiple of 4.

---

**Note:** If you use the *partition* command to invoke different “processors” commands on different partitions, and you also use the *part* keyword, then you must insure that both the sending and receiving partitions invoke the “processors” command that connects the 2 partitions via the *part* keyword. LAMMPS cannot easily check for this, but your simulation will likely hang in its setup phase if this error has been made.

---



The *file* keyword writes the mapping of the factorization of P processors and their mapping to the 3d grid to the specified file *outfile*. This is useful to check that you assigned physical processors in the manner you desired, which can be tricky to figure out, especially when running on multiple partitions or on, a multicore machine or when the processor ranks were reordered by use of the *-reorder command-line switch* or due to use of MPI-specific launch options such as a config file.

If you have multiple partitions you should insure that each one writes to a different file, e.g. using a *world-style variable* for the filename. The file has a self-explanatory header, followed by one-line per processor in this format:

```
world-ID universe-ID original-ID: I J K: name
```

The IDs are the processor's rank in this simulation (the world), the universe (of multiple simulations), and the original MPI communicator used to instantiate LAMMPS, respectively. The world and universe IDs will only be different if you are running on more than one partition; see the *-partition command-line switch*. The universe and original IDs will only be different if you used the *-reorder command-line switch* to reorder the processors differently than their rank in the original communicator LAMMPS was instantiated with.

I,J,K are the indices of the processor in the regular 3d grid, each from 1 to Nd, where Nd is the number of processors in that dimension of the grid.

The *name* is what is returned by a call to `MPI_Get_processor_name()` and should represent an identifier relevant to the physical processors in your machine. Note that depending on the MPI implementation, multiple cores can have the same *name*.

#### 16.104.4 Restrictions

This command cannot be used after the simulation box is defined by a *read\_data* or *create\_box* command. It can be used before a restart file is read to change the 3d processor grid from what is specified in the restart file.

The *grid numa* keyword only currently works with the *map cart* option.

The *part* keyword (for the receiving partition) only works with the *grid onelevel* or *grid twolevel* options.

#### 16.104.5 Related commands

*partition*, *-reorder command-line switch*

#### 16.104.6 Default

The option defaults are Px Py Pz = \* \* \*, grid = onelevel, and map = cart.

### 16.105 python command

#### 16.105.1 Syntax

```
python func keyword args ...
```

- func = name of Python function
- one or more keyword/args pairs must be appended

keyword = *invoke* or *input* or *return* or *format* or *length* or *file* or *here* or *exists* ↵  
↵or *source*  
*invoke* arg = none = invoke the previously defined Python function  
*input* args = N i1 i2 ... iN  
N = # of inputs to function  
i1,...,iN = value, SELF, or LAMMPS variable name  
value = integer number, floating point number, or string  
SELF = reference to LAMMPS itself which can be accessed by Python function  
variable = v\_name, where name = name of LAMMPS variable, e.g. v\_abc  
*return* arg = varReturn  
varReturn = v\_name = LAMMPS variable name which return value of function will ↵  
↵be assigned to  
*format* arg = fstring with M characters  
M = N if no return value, where N = # of inputs  
M = N+1 if there is a return value  
fstring = each character (i,f,s,p) corresponds in order to an input or return ↵  
↵value  
'i' = integer, 'f' = floating point, 's' = string, 'p' = SELF  
*length* arg = Nlen  
Nlen = max length of string returned from Python function  
*file* arg = filename  
filename = file of Python code, which defines func  
*here* arg = inline  
inline = one or more lines of Python code which defines func  
must be a single argument, typically enclosed between triple quotes  
*exists* arg = none = Python code has been loaded by previous python command  
*source* arg = *filename* or *inline*  
filename = file of Python code which will be executed immediately  
inline = one or more lines of Python code which will be executed immediately  
must be a single argument, typically enclosed between triple quotes

## 16.105.2 Examples

```
python pForce input 2 v_x 20.0 return v_f format fff file force.py
python pForce invoke

python factorial input 1 myN return v_fac format ii here """
def factorial(n):
    if n == 1: return n
    return n * factorial(n-1)
"""

python loop input 1 SELF return v_value format pf here """
def loop(lmp_ptr,N,cut0):
    from lammps import lammps
    lmp = lammps(ptr=lmp_ptr)

    # loop N times, increasing cutoff each time

    for i in range(N):
        cut = cut0 + i*0.1
        lmp.set_variable("cut",cut)                # set a variable in LAMMPS
```

(continues on next page)

(continued from previous page)

```

    lmp.command("pair_style lj/cut ${cut}")    # LAMMPS commands
    lmp.command("pair_coeff * * 1.0 1.0")
    lmp.command("run 100")
    """

```

### 16.105.3 Description

Define a Python function or execute a previously defined function or execute some arbitrary python code. Arguments, including LAMMPS variables, can be passed to the function from the LAMMPS input script and a value returned by the Python function to a LAMMPS variable. The Python code for the function can be included directly in the input script or in a separate Python file. The function can be standard Python code or it can make “callbacks” to LAMMPS through its library interface to query or set internal values within LAMMPS. This is a powerful mechanism for performing complex operations in a LAMMPS input script that are not possible with the simple input script and variable syntax which LAMMPS defines. Thus your input script can operate more like a true programming language.

Use of this command requires building LAMMPS with the PYTHON package which links to the Python library so that the Python interpreter is embedded in LAMMPS. More details about this process are given below.

There are two ways to invoke a Python function once it has been defined. One is using the *invoke* keyword. The other is to assign the function to a *python-style variable* defined in your input script. Whenever the variable is evaluated, it will execute the Python function to assign a value to the variable. Note that variables can be evaluated in many different ways within LAMMPS. They can be substituted for directly in an input script. Or they can be passed to various commands as arguments, so that the variable is evaluated during a simulation run.

A broader overview of how Python can be used with LAMMPS is given on the [Python](#) doc page. There is an examples/python directory which illustrates use of the python command.

---

The *func* setting specifies the name of the Python function. The code for the function is defined using the *file* or *here* keywords as explained below. In case of the *source* keyword, the name of the function is ignored.

If the *invoke* keyword is used, no other keywords can be used, and a previous python command must have defined the Python function referenced by this command. This invokes the Python function with the previously defined arguments and return value processed as explained below. You can invoke the function as many times as you wish in your input script.

If the *source* keyword is used, no other keywords can be used. The argument can be a filename or a string with python commands, either on a single line enclosed in quotes, or as multiple lines enclosed in triple quotes. These python commands will be passed to the python interpreter and executed immediately without registering a python function for future execution.

The *input* keyword defines how many arguments *N* the Python function expects. If it takes no arguments, then the *input* keyword should not be used. Each argument can be specified directly as a value, e.g. 6 or 3.14159 or abc (a string of characters). The type of each argument is specified by the *format* keyword as explained below, so that Python will know how to interpret the value. If the word SELF is used for an argument it has a special meaning. A pointer is passed to the Python function which it converts into a reference to LAMMPS itself. This enables the function to call back to LAMMPS through its library interface as explained below. This allows the Python function to query or set values internal to LAMMPS which can affect the subsequent execution of the input script. A LAMMPS variable can also be used as an argument, specified as *v\_name*, where “name” is the name of the variable. Any style of LAMMPS variable can be used, as defined by the *variable* command. Each time the Python function is invoked, the LAMMPS variable is evaluated and its value is passed to the Python function.

The *return* keyword is only needed if the Python function returns a value. The specified *varReturn* must be of the form *v\_name*, where “name” is the name of a python-style LAMMPS variable, defined by the *variable* command. The Python function can return a numeric or string value, as specified by the *format* keyword.

As explained on the [variable](#) doc page, the definition of a python-style variable associates a Python function name with the variable. This must match the *func* setting for this command. For example these two commands would be self-consistent:

```
variable foo python myMultiply
python myMultiply return v_foo format f file funcs.py
```

The two commands can appear in either order in the input script so long as both are specified before the Python function is invoked for the first time.

The *format* keyword must be used if the *input* or *return* keyword is used. It defines an *fstring* with M characters, where M = sum of number of inputs and outputs. The order of characters corresponds to the N inputs, followed by the return value (if it exists). Each character must be one of the following: “i” for integer, “f” for floating point, “s” for string, or “p” for SELF. Each character defines the type of the corresponding input or output value of the Python function and affects the type conversion that is performed internally as data is passed back and forth between LAMMPS and Python. Note that it is permissible to use a *python-style variable* in a LAMMPS command that allows for an equal-style variable as an argument, but only if the output of the Python function is flagged as a numeric value (“i” or “f”) via the *format* keyword.

If the *return* keyword is used and the *format* keyword specifies the output as a string, then the default maximum length of that string is 63 characters (64-1 for the string terminator). If you want to return a longer string, the *length* keyword can be specified with its *Nlen* value set to a larger number (the code allocates space for Nlen+1 to include the string terminator). If the Python function generates a string longer than the default 63 or the specified *Nlen*, it will be truncated.

---

Either the *file*, *here*, or *exists* keyword must be used, but only one of them. These keywords specify what Python code to load into the Python interpreter. The *file* keyword gives the name of a file, which should end with a “.py” suffix, which contains Python code. The code will be immediately loaded into and run in the “main” module of the Python interpreter. Note that Python code which contains a function definition does not “execute” the function when it is run; it simply defines the function so that it can be invoked later.

The *here* keyword does the same thing, except that the Python code follows as a single argument to the *here* keyword. This can be done using triple quotes as delimiters, as in the examples above. This allows Python code to be listed verbatim in your input script, with proper indentation, blank lines, and comments, as desired. See the [Commands parse](#) doc page, for an explanation of how triple quotes can be used as part of input script syntax.

The *exists* keyword takes no argument. It means that Python code containing the required Python function defined by the *func* setting, is assumed to have been previously loaded by another python command.

Note that the Python code that is loaded and run must contain a function with the specified *func* name. To operate properly when later invoked, the function code must match the *input* and *return* and *format* keywords specified by the python command. Otherwise Python will generate an error.

---

This section describes how Python code can be written to work with LAMMPS.

Whether you load Python code from a file or directly from your input script, via the *file* and *here* keywords, the code can be identical. It must be indented properly as Python requires. It can contain comments or blank lines. If the code is in your input script, it cannot however contain triple-quoted Python strings, since that will conflict with the triple-quote parsing that the LAMMPS input script performs.

All the Python code you specify via one or more python commands is loaded into the Python “main” module, i.e. `__main__`. The code can define global variables or statements that are outside of function definitions. It can contain multiple functions, only one of which matches the *func* setting in the python command. This means you can use the *file* keyword once to load several functions, and the *exists* keyword thereafter in subsequent python commands to access the other functions previously loaded.

A Python function you define (or more generally, the code you load) can import other Python modules or classes, it can make calls to other system functions or functions you define, and it can access or modify global variables (in the “main” module) which will persist between successive function calls. The latter can be useful, for example, to prevent a function from being invoked multiple times per timestep by different commands in a LAMMPS input script that access the returned python-style variable associated with the function. For example, consider this function loaded with two global variables defined outside the function:

```
nsteplast = -1
nvaluelast = 0

def expensive(nstep):
    global nsteplast, nvaluelast
    if nstep == nsteplast: return nvaluelast
    nsteplast = nstep
    # perform complicated calculation
    nvalue = ...
    nvaluelast = nvalue
    return nvalue
```

Nsteplast stores the previous timestep the function was invoked (passed as an argument to the function). Nvaluelast stores the return value computed on the last function invocation. If the function is invoked again on the same timestep, the previous value is simply returned, without re-computing it. The “global” statement inside the Python function allows it to overwrite the global variables.

Note that if you load Python code multiple times (via multiple python commands), you can overwrite previously loaded variables and functions if you are not careful. E.g. if the code above were loaded twice, the global variables would be re-initialized, which might not be what you want. Likewise, if a function with the same name exists in two chunks of Python code you load, the function loaded second will override the function loaded first.

It’s important to realize that if you are running LAMMPS in parallel, each MPI task will load the Python interpreter and execute a local copy of the Python function(s) you define. There is no connection between the Python interpreters running on different processors. This implies three important things.

First, if you put a print statement in your Python function, you will see P copies of the output, when running on P processors. If the prints occur at (nearly) the same time, the P copies of the output may be mixed together. Welcome to the world of parallel programming and debugging.

Second, if your Python code loads modules that are not pre-loaded by the Python library, then it will load the module from disk. This may be a bottleneck if 1000s of processors try to load a module at the same time. On some large supercomputers, loading of modules from disk by Python may be disabled. In this case you would need to pre-build a Python library that has the required modules pre-loaded and link LAMMPS with that library.

Third, if your Python code calls back to LAMMPS (discussed in the next section) and causes LAMMPS to perform an MPI operation requires global communication (e.g. via MPI\_Allreduce), such as computing the global temperature of the system, then you must insure all your Python functions (running independently on different processors) call back to LAMMPS. Otherwise the code may hang.

Your Python function can “call back” to LAMMPS through its library interface, if you use the SELF input to pass Python a pointer to LAMMPS. The mechanism for doing this in your Python function is as follows:

```
def foo(lmp_ptr, ...):
    from lammpls import lammpls
    lmp = lammpls(ptr=lmp_ptr)
    lmp.command('print "Hello from inside Python"')
    ...
```

The function definition must include a variable (lmpptr in this case) which corresponds to SELF in the python command. The first line of the function imports the “*lammps*” Python module. The second line creates a Python object lmp which wraps the instance of LAMMPS that called the function. The “ptr=lmpptr” argument is what makes that happen. The third line invokes the command() function in the LAMMPS library interface. It takes a single string argument which is a LAMMPS input script command for LAMMPS to execute, the same as if it appeared in your input script. In this case, LAMMPS should output

```
Hello from inside Python
```

to the screen and log file. Note that since the LAMMPS print command itself takes a string in quotes as its argument, the Python string must be delimited with a different style of quotes.

The *Use Python with LAMMPS* page describes the syntax for how Python wraps the various functions included in the LAMMPS library interface.

A more interesting example is in the examples/python/in.python script which loads and runs the following function from examples/python/funcs.py:

```
def loop(N,cut0,thresh,lmpptr):
    print("LOOP ARGS", N, cut0, thresh, lmpptr)
    from lammps import lammps
    lmp = lammps(ptr=lmpptr)
    natoms = lmp.get_natoms()

    for i in range(N):
        cut = cut0 + i*0.1

        lmp.set_variable("cut",cut)           # set a variable in LAMMPS
        lmp.command("pair_style lj/cut ${cut}") # LAMMPS command
        #lmp.command("pair_style lj/cut %d" % cut) # LAMMPS command option

        lmp.command("pair_coeff * * 1.0 1.0") # ditto
        lmp.command("run 10")                # ditto
        pe = lmp.extract_compute("thermo_pe",0,0) # extract total PE from LAMMPS
        print("PE", pe/natoms, thresh)
        if pe/natoms < thresh: return
```

with these input script commands:

```
python      loop input 4 10 1.0 -4.0 SELF format iffp file funcs.py
python      loop invoke
```

This has the effect of looping over a series of 10 short runs (10 timesteps each) where the pair style cutoff is increased from a value of 1.0 in distance units, in increments of 0.1. The looping stops when the per-atom potential energy falls below a threshold of -4.0 in energy units. More generally, Python can be used to implement a loop with complex logic, much more so than can be created using the LAMMPS *jump* and *if* commands.

Several LAMMPS library functions are called from the loop function. Get\_natoms() returns the number of atoms in the simulation, so that it can be used to normalize the potential energy that is returned by extract\_compute() for the “thermo\_pe” compute that is defined by default for LAMMPS thermodynamic output. Set\_variable() sets the value of a string variable defined in LAMMPS. This library function is a useful way for a Python function to return multiple values to LAMMPS, more than the single value that can be passed back via a return statement. This cutoff value in the “cut” variable is then substituted (by LAMMPS) in the pair\_style command that is executed next. Alternatively, the “LAMMPS command option” line could be used in place of the 2 preceding lines, to have Python insert the value into the LAMMPS command string.

---

**Note:** When using the callback mechanism just described, recognize that there are some operations you should not attempt because LAMMPS cannot execute them correctly. If the Python function is invoked between runs in the LAMMPS input script, then it should be OK to invoke any LAMMPS input script command via the library interface `command()` or `file()` functions, so long as the command would work if it were executed in the LAMMPS input script directly at the same point.

---

However, a Python function can also be invoked during a run, whenever an associated LAMMPS variable it is assigned to is evaluated. If the variable is an input argument to another LAMMPS command (e.g. *fix setforce*), then the Python function will be invoked inside the class for that command, in one of its methods that is invoked in the middle of a timestep. You cannot execute arbitrary input script commands from the Python function (again, via the `command()` or `file()` functions) at that point in the run and expect it to work. Other library functions such as those that invoke `compute` or other variables may have hidden side effects as well. In these cases, LAMMPS has no simple way to check that something illogical is being attempted.

The same applies to Python functions called during a simulation run at each time step using *fix python/invoke*.

---

If you run Python code directly on your workstation, either interactively or by using Python to launch a Python script stored in a file, and your code has an error, you will typically see informative error messages. That is not the case when you run Python code from LAMMPS using an embedded Python interpreter. The code will typically fail silently. LAMMPS will catch some errors but cannot tell you where in the Python code the problem occurred. For example, if the Python code cannot be loaded and run because it has syntax or other logic errors, you may get an error from Python pointing to the offending line, or you may get one of these generic errors from LAMMPS:

```
Could not process Python file
Could not process Python string
```

When the Python function is invoked, if it does not return properly, you will typically get this generic error from LAMMPS:

```
Python function evaluation failed
```

Here are three suggestions for debugging your Python code while running it under LAMMPS.

First, don't run it under LAMMPS, at least to start with! Debug it using plain Python. Load and invoke your function, pass it arguments, check return values, etc.

Second, add Python print statements to the function to check how far it gets and intermediate values it calculates. See the discussion above about printing from Python when running in parallel.

Third, use Python exception handling. For example, say this statement in your Python function is failing, because you have not initialized the variable `foo`:

```
foo += 1
```

If you put one (or more) statements inside a "try" statement, like this:

```
import exceptions
print("Inside simple function")
try:
    foo += 1      # one or more statements here
except Exception as e:
    print("FOO error:", e)
```

then you will get this message printed to the screen:

```
F00 error: local variable 'foo' referenced before assignment
```

If there is no error in the try statements, then nothing is printed. Either way the function continues on (unless you put a return or sys.exit() in the except clause).

---

### 16.105.4 Restrictions

This command is part of the PYTHON package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Building LAMMPS with the PYTHON package will link LAMMPS with the Python library on your system. Settings to enable this are in the lib/python/Makefile.lammps file. See the lib/python/README file for information on those settings.

If you use Python code which calls back to LAMMPS, via the SELF input argument explained above, there is an extra step required when building LAMMPS. LAMMPS must also be built as a shared library and your Python function must be able to load the “*lammps*” Python module that wraps the LAMMPS library interface. These are the same steps required to use Python by itself to wrap LAMMPS. Details on these steps are explained on the *Python* doc page. Note that it is important that the stand-alone LAMMPS executable and the LAMMPS shared library be consistent (built from the same source code files) in order for this to work. If the two have been built at different times using different source files, problems may occur.

### 16.105.5 Related commands

*shell, variable, fix python/invoke*

### 16.105.6 Default

none

## 16.106 quit command

### 16.106.1 Syntax

```
quit status
```

status = numerical exit status (optional)



## 16.106.2 Examples

```
quit
if "$n > 10000" then "quit 1"
```

## 16.106.3 Description

This command causes LAMMPS to exit, after shutting down all output cleanly.

It can be used as a debug statement in an input script, to terminate the script at some intermediate point.

It can also be used as an invoked command inside the “then” or “else” portion of an *if* command.

The optional status argument is an integer which signals the return status to a program calling LAMMPS. A return status of 0 usually indicates success. A status != 0 is failure, where the specified value can be used to distinguish the kind of error, e.g. where in the input script the quit was invoked. If not specified, a status of 0 is returned.

## 16.106.4 Restrictions

none

## 16.106.5 Related commands

*if*

## 16.106.6 Default

none

## 16.107 read\_data command

### 16.107.1 Syntax

```
read_data file keyword args ...
```

- file = name of data file to read in
- zero or more keyword/arg pairs may be appended
- keyword = *add* or *offset* or *shift* or *extra/atom/types* or *extra/bond/types* or *extra/angle/types* or *extra/dihedral/types* or *extra/improper/types* or *extra/bond/per/atom* or *extra/angle/per/atom* or *extra/dihedral/per/atom* or *extra/improper/per/atom* or *group* or *nocoeff* or *fix*

*add* arg = *append* or *IDoffset* or *IDoffset MOLOffset* or *merge*

*append* = add new atoms with atom IDs appended to current IDs

*IDoffset* = add new atoms with atom IDs having IDoffset added

*MOLOffset* = add new atoms with molecule IDs having MOLOffset added (only when [molecule IDs are enabled](#))

*merge* = add new atoms with their atom IDs (and molecule IDs) unchanged

*offset* args = toff boff aoff doff ioff

```
toff = offset to add to atom types
boff = offset to add to bond types
aoff = offset to add to angle types
doff = offset to add to dihedral types
ioff = offset to add to improper types
shift args = Sx Sy Sz
  Sx,Sy,Sz = distance to shift atoms when adding to system (distance units)
extra/atom/types arg = # of extra atom types
extra/bond/types arg = # of extra bond types
extra/angle/types arg = # of extra angle types
extra/dihedral/types arg = # of extra dihedral types
extra/improper/types arg = # of extra improper types
extra/bond/per/atom arg = leave space for this many new bonds per atom
extra/angle/per/atom arg = leave space for this many new angles per atom
extra/dihedral/per/atom arg = leave space for this many new dihedrals per atom
extra/improper/per/atom arg = leave space for this many new impropers per atom
extra/special/per/atom arg = leave space for extra 1-2,1-3,1-4 interactions per atom
group args = groupID
  groupID = add atoms in data file to this group
nocoeff = ignore force field parameters
fix args = fix-ID header-string section-string
  fix-ID = ID of fix to process header lines and sections of data file
  header-string = header lines containing this string will be passed to fix
  section-string = section names with this string will be passed to fix
```

## 16.107.2 Examples

```
read_data data.lj
read_data ../run7/data.polymer.gz
read_data data.protein fix mycmap crossterm CMAP
read_data data.water add append offset 3 1 1 1 1 shift 0.0 0.0 50.0
read_data data.water add merge group solvent
```

## 16.107.3 Description

Read in a data file containing information LAMMPS needs to run a simulation. The file can be ASCII text or a gzipped text file (detected by a `.gz` suffix). This is one of 3 ways to specify initial atom coordinates; see the `read_restart` and `create_atoms` commands for alternative methods. Also see the explanation of the `-restart command-line switch` which can convert a restart file to a data file.

This command can be used multiple times to add new atoms and their properties to an existing system by using the `add`, `offset`, and `shift` keywords. See more details below, which includes the use case for the `extra` keywords.

The `group` keyword adds all the atoms in the data file to the specified group-ID. The group will be created if it does not already exist. This is useful if you are reading multiple data files and wish to put sets of atoms into different groups so they can be operated on later. E.g. a group of added atoms can be moved to new positions via the `displace_atoms` command. Note that atoms read from the data file are also always added to the “all” group. The `group` command discusses atom groups, as used in LAMMPS.

The `nocoeff` keyword tells `read_data` to ignore force field parameters. The various `Coeff` sections are still read and have to have the correct number of lines, but they are not applied. This also allows to read a data file without having any pair, bond, angle, dihedral or improper styles defined, or to read a data file for a different force field.

The use of the *fix* keyword is discussed below.

---

### 16.107.4 Reading multiple data files

The `read_data` command can be used multiple times with the same or different data files to build up a complex system from components contained in individual data files. For example one data file could contain fluid in a confined domain; a second could contain wall atoms, and the second file could be read a third time to create a wall on the other side of the fluid. The third set of atoms could be rotated to an opposing direction using the `displace_atoms` command, after the third `read_data` command is used.

The `add`, `offset`, `shift`, `extra`, and `group` keywords are useful in this context.

If a simulation box does not yet exist, the `add` keyword cannot be used; the `read_data` command is being used for the first time. If a simulation box does exist, due to using the `create_box` command, or a previous `read_data` command, then the `add` keyword must be used.

---

**Note:** The simulation box size (xlo to xhi, ylo to yhi, zlo to zhi) in the new data file will be merged with the existing simulation box to create a large enough box in each dimension to contain both the existing and new atoms. Each box dimension never shrinks due to this merge operation, it only stays the same or grows. Care must be used if you are growing the existing simulation box in a periodic dimension. If there are existing atoms with bonds that straddle that periodic boundary, then the atoms may become far apart if the box size grows. This will separate the atoms in the bond, which can lead to “lost” bond atoms or bad dynamics.

---

The three choices for the `add` argument affect how the atom IDs and molecule IDs of atoms in the data file are treated. If `append` is specified, atoms in the data file are added to the current system, with their atom IDs reset so that an atom-ID = M in the data file becomes atom-ID = N+M, where N is the largest atom ID in the current system. This rule is applied to all occurrences of atom IDs in the data file, e.g. in the Velocity or Bonds section. This is also done for molecule IDs, if the atom style does support molecule IDs or they are enabled via `fix property/atom`. If `IDoffset` is specified, then `IDoffset` is a numeric value is given, e.g. 1000, so that an atom-ID = M in the data file becomes atom-ID = 1000+M. For systems with enabled molecule IDs, another numerical argument `MOLoffset` is required representing the equivalent offset for molecule IDs. If `merge` is specified, the data file atoms are added to the current system without changing their IDs. They are assumed to merge (without duplication) with the currently defined atoms. It is up to you to insure there are no multiply defined atom IDs, as LAMMPS only performs an incomplete check that this is the case by insuring the resulting max atom-ID  $\geq$  the number of atoms. For molecule IDs, there is no check done at all.

The `offset` and `shift` keywords can only be used if the `add` keyword is also specified.

The `offset` keyword adds the specified offset values to the atom types, bond types, angle types, dihedral types, and improper types as they are read from the data file. E.g. if `toff` = 2, and the file uses atom types 1,2,3, then the added atoms will have atom types 3,4,5. These offsets apply to all occurrences of types in the data file, e.g. for the Atoms or Masses or Pair Coeffs or Bond Coeffs sections. This makes it easy to use atoms and molecules and their attributes from a data file in different simulations, where you want their types (atom, bond, angle, etc) to be different depending on what other types already exist. All five offset values must be specified, but individual values will be ignored if the data file does not use that attribute (e.g. no bonds).

---

**Note:** Offsets are **ignored** on lines using type labels, as the type labels will determine the actual types directly depending on the current `labelmap` settings.

---

The `shift` keyword can be used to specify an (Sx, Sy, Sz) displacement applied to the coordinates of each atom. Sz must be 0.0 for a 2d simulation. This is a mechanism for adding structured collections of atoms at different locations within the simulation box, to build up a complex geometry. It is up to you to insure atoms do not end up overlapping

unphysically which would lead to bad dynamics. Note that the *displace\_atoms* command can be used to move a subset of atoms after they have been read from a data file. Likewise, the *delete\_atoms* command can be used to remove overlapping atoms. Note that the shift values (Sx, Sy, Sz) are also added to the simulation box information (xlo, xhi, ylo, yhi, zlo, zhi) in the data file to shift its boundaries. E.g.  $xlo\_new = xlo + Sx$ ,  $xhi\_new = xhi + Sx$ .

The *extra* keywords can only be used the first time the *read\_data* command is used. They are useful if you intend to add new atom, bond, angle, etc types later with additional *read\_data* commands. This is because the maximum number of allowed atom, bond, angle, etc types is set by LAMMPS when the system is first initialized. If you do not use the *extra* keywords, then the number of these types will be limited to what appears in the first data file you read. For example, if the first data file is a solid substrate of Si, it will likely specify a single atom type. If you read a second data file with a different material (water molecules) that sit on top of the substrate, you will want to use different atom types for those atoms. You can only do this if you set the *extra/atom/types* keyword to a sufficiently large value when reading the substrate data file. Note that use of the *extra* keywords also allows each data file to contain sections like Masses or Pair Coeffs or Bond Coeffs which are sized appropriately for the number of types in that data file. If the *offset* keyword is used appropriately when each data file is read, the values in those sections will be stored correctly in the larger data structures allocated by the use of the *extra* keywords. E.g. the substrate file can list mass and pair coefficients for type 1 silicon atoms. The water file can list mass and pair coefficients for type 1 and type 2 hydrogen and oxygen atoms. Use of the *extra* and *offset* keywords will store those mass and pair coefficient values appropriately in data structures that allow for 3 atom types (Si, H, O). Of course, you would still need to specify coefficients for H/Si and O/Si interactions in your input script to have a complete pairwise interaction model.

An alternative to using the *extra* keywords with the *read\_data* command, is to use the *create\_box* command to initialize the simulation box and all the various type limits you need via its *extra* keywords. Then use the *read\_data* command one or more times to populate the system with atoms, bonds, angles, etc, using the *offset* keyword if desired to alter types used in the various data files you read.

---

### 16.107.5 Format of a data file

The structure of the data file is important, though many settings and sections are optional or can come in any order. See the examples directory for sample data files for different problems.

The file will be read line by line, but there is a limit of 254 characters per line and characters beyond that limit will be ignored.

A data file has a header and a body. The header appears first. The first line of the header and thus of the data file is *always* skipped; it typically contains a description of the file or a comment from the software that created the file.

Then lines are read one line at a time. Lines can have a trailing comment starting with '#' that is ignored. There *must* be at least one blank between any valid content and the comment. If a line is blank (i.e. contains only white-space after comments are deleted), it is skipped. If the line contains a header keyword, the corresponding value(s) is/are read from the line. A line that is *not* blank and does *not* contain a header keyword begins the body of the file.

The body of the file contains zero or more sections. The first line of a section has only a keyword. This line can have a trailing comment starting with '#' that is either ignored or can be used to check for a style match, as described below. There must be a blank between the keyword and any comment. The *next* line is *always* skipped. The remaining lines of the section contain values. The number of lines depends on the section keyword as described below. Zero or more blank lines can be used *between* sections. Sections can appear in any order, with a few exceptions as noted below.

The keyword *fix* can be used one or more times. Each usage specifies a fix that will be used to process a specific portion of the data file. Any header line containing *header-string* and any section that is an exact match to *section-string* will be passed to the specified fix. See the *fix property/atom* command for an example of a fix that operates in this manner. The doc page for the fix defines the syntax of the header line(s) and section that it reads from the data file. Note that the *header-string* can be specified as NULL, in which case no header lines are passed to the fix. This means the fix can infer the length of its Section from standard header settings, such as the number of atoms. Also the *section-string* may be specified as NULL, and in that case the fix ID is used as section name.

The formatting of individual lines in the data file (indentation, spacing between words and numbers) is not important except that header and section keywords (e.g. *atoms*, *xlo xhi*, *Masses*, *Bond Coeffs*) must be capitalized as shown and cannot have extra white-space between their words - e.g. two spaces or a tab between the 2 words in “*xlo xhi*” or the 2 words in “*Bond Coeffs*”, is not valid.

### 16.107.6 Format of the header of a data file

These are the recognized header keywords. Header lines can come in any order. The value(s) are read from the beginning of the line. Thus the keyword *atoms* should be in a line like “1000 *atoms*”; the keyword *ylo yhi* should be in a line like “-10.0 10.0 *ylo yhi*”; the keyword *xy xz yz* should be in a line like “0.0 5.0 6.0 *xy xz yz*”. All these settings have a default value of 0, except the *lo/hi* box size defaults are -0.5 and 0.5. A line need only appear if the value is different than the default.

- *atoms* = # of atoms in system
- *bonds* = # of bonds in system
- *angles* = # of angles in system
- *dihedrals* = # of dihedrals in system
- *impropers* = # of impropers in system
- *atom types* = # of atom types in system
- *bond types* = # of bond types in system
- *angle types* = # of angle types in system
- *dihedral types* = # of dihedral types in system
- *improper types* = # of improper types in system
- *extra bond per atom* = leave space for this many new bonds per atom (deprecated, use *extra/bond/per/atom* keyword)
- *extra angle per atom* = leave space for this many new angles per atom (deprecated, use *extra/angle/per/atom* keyword)
- *extra dihedral per atom* = leave space for this many new dihedrals per atom (deprecated, use *extra/dihedral/per/atom* keyword)
- *extra improper per atom* = leave space for this many new impropers per atom (deprecated, use *extra/improper/per/atom* keyword)
- *extra special per atom* = leave space for this many new special bonds per atom (deprecated, use *extra/special/per/atom* keyword)
- *ellipsoids* = # of ellipsoids in system
- *lines* = # of line segments in system
- *triangles* = # of triangles in system
- *bodies* = # of bodies in system
- *xlo xhi* = simulation box boundaries in x dimension
- *ylo yhi* = simulation box boundaries in y dimension
- *zlo zhi* = simulation box boundaries in z dimension
- *xy xz yz* = simulation box tilt factors for triclinic system

The initial simulation box size is determined by the *lo/hi* settings. In any dimension, the system may be periodic or non-periodic; see the *boundary* command. When the simulation box is created it is also partitioned into a regular 3d grid of rectangular bricks, one per processor, based on the number of processors being used and the settings of the *processors* command. The partitioning can later be changed by the *balance* or *fix balance* commands.

If the *xy xz yz* line does not appear, LAMMPS will set up an axis-aligned (orthogonal) simulation box. If the line does appear, LAMMPS creates a non-orthogonal simulation domain shaped as a parallelepiped with triclinic symmetry. The parallelepiped has its “origin” at (*xlo,ylo,zlo*) and is defined by 3 edge vectors starting from the origin given by  $A = (xhi-xlo,0,0)$ ;  $B = (xy,yhi-ylo,0)$ ;  $C = (xz,yz,zhi-zlo)$ . *Xy,xz,yz* can be 0.0 or positive or negative values and are called “tilt factors” because they are the amount of displacement applied to faces of an originally orthogonal box to transform it into the parallelepiped.

By default, the tilt factors (*xy,xz,yz*) can not skew the box more than half the distance of the corresponding parallel box length. For example, if *xlo* = 2 and *xhi* = 12, then the x box length is 10 and the xy tilt factor must be between -5 and 5. Similarly, both *xz* and *yz* must be between  $-(xhi-xlo)/2$  and  $+(yhi-ylo)/2$ . Note that this is not a limitation, since if the maximum tilt factor is 5 (as in this example), then configurations with tilt = ..., -15, -5, 5, 15, 25, ... are all geometrically equivalent. If you wish to define a box with tilt factors that exceed these limits, you can use the *box tilt* command, with a setting of *large*; a setting of *small* is the default.

See the *Howto triclinic* page for a geometric description of triclinic boxes, as defined by LAMMPS, and how to transform these parameters to and from other commonly used triclinic representations.

When a triclinic system is used, the simulation domain should normally be periodic in the dimension that the tilt is applied to, which is given by the second dimension of the tilt factor (e.g. y for xy tilt). This is so that pairs of atoms interacting across that boundary will have one of them shifted by the tilt factor. Periodicity is set by the *boundary* command. For example, if the xy tilt factor is non-zero, then the y dimension should be periodic. Similarly, the z dimension should be periodic if *xz* or *yz* is non-zero. LAMMPS does not require this periodicity, but you may lose atoms if this is not the case.

Also note that if your simulation will tilt the box, e.g. via the *fix deform* command, the simulation box must be setup to be triclinic, even if the tilt factors are initially 0.0. You can also change an orthogonal box to a triclinic box or vice versa by using the *change box* command with its *ortho* and *triclinic* options.

For 2d simulations, the *zlo zhi* values should be set to bound the z coords for atoms that appear in the file; the default of -0.5 0.5 is valid if all z coords are 0.0. For 2d triclinic simulations, the *xz* and *yz* tilt factors must be 0.0.

If the system is periodic (in a dimension), then atom coordinates can be outside the bounds (in that dimension); they will be remapped (in a periodic sense) back inside the box. Note that if the *add* option is being used to add atoms to a simulation box that already exists, this periodic remapping will be performed using simulation box bounds that are the union of the existing box and the box boundaries in the new data file.

If the system is non-periodic (in a dimension), then an image flag for that direction has no meaning, since there cannot be periodic images without periodicity and the data file is therefore - technically speaking - invalid. This situation would happen when a data file was written with periodic boundaries and then read back for non-periodic boundaries. Accepting a non-zero image flag can lead to unexpected results for any operations and computations in LAMMPS that internally use unwrapped coordinates (for example computing the center of mass of a group of atoms). Thus all non-zero image flags for non-periodic dimensions will be reset to zero on reading the data file and LAMMPS will print a warning message, if that happens. This is equivalent to wrapping atoms individually back into the principal unit cell in that direction. This operation is equivalent to the behavior of the *change\_box command* when used to change periodicity.

If those atoms with non-zero image flags are involved in bonded interactions, this reset can lead to undesired changes, when the image flag values differ between the atoms, i.e. the bonded interaction straddles domain boundaries. For example a bond can become stretched across the unit cell if one of its atoms is wrapped to one side of the cell and the second atom to the other. In those cases the data file needs to be pre-processed externally to become valid again. This can be done by first unwrapping coordinates and then wrapping entire molecules instead of individual atoms back into the principal simulation cell and finally expanding the cell dimensions in the non-periodic direction as needed, so that the image flag would be zero.

**Note:** If the system is non-periodic (in a dimension), then all atoms in the data file must have coordinates (in that dimension) that are “greater than or equal to” the lo value and “less than or equal to” the hi value. If the non-periodic dimension is of style “fixed” (see the *boundary* command), then the atom coords must be strictly “less than” the hi value, due to the way LAMMPS assign atoms to processors. Note that you should not make the lo/hi values radically smaller/larger than the extent of the atoms. For example, if your atoms extend from 0 to 50, you should not specify the box bounds as -10000 and 10000 unless you also use the *processors* command. This is because LAMMPS uses the specified box size to layout the 3d grid of processors. A huge (mostly empty) box will be sub-optimal for performance when using “fixed” boundary conditions (see the *boundary* command). When using “shrink-wrap” boundary conditions (see the *boundary* command), a huge (mostly empty) box may cause a parallel simulation to lose atoms when LAMMPS shrink-wraps the box around the atoms. The *read\_data* command will generate an error in this case.

---

The “extra bond per atom” setting (angle, dihedral, improper) is only needed if new bonds (angles, dihedrals, impropers) will be added to the system when a simulation runs, e.g. by using the *fix bond/create* command. Using this header flag is deprecated; please use the *extra/bond/per/atom* keyword (and correspondingly for angles, dihedrals and impropers) in the *read\_data* command instead. Either will pre-allocate space in LAMMPS data structures for storing the new bonds (angles, dihedrals, impropers).

The “extra special per atom” setting is typically only needed if new bonds/angles/etc will be added to the system, e.g. by using the *fix bond/create* command. Or if entire new molecules will be added to the system, e.g. by using the *fix deposit* or *fix pour* commands, which will have more special 1-2,1-3,1-4 neighbors than any other molecules defined in the data file. Using this header flag is deprecated; please use the *extra/special/per/atom* keyword instead. Using this setting will pre-allocate space in the LAMMPS data structures for storing these neighbors. See the *special\_bonds* and *molecule* doc pages for more discussion of 1-2,1-3,1-4 neighbors.

---

**Note:** All of the “extra” settings are only applied in the first data file read and when no simulation box has yet been created; as soon as the simulation box is created (and *read\_data* implies that), these settings are *locked* and cannot be changed anymore. Please see the description of the *add* keyword above for reading multiple data files. If they appear in later data files, they are ignored.

---

The “ellipsoids” and “lines” and “triangles” and “bodies” settings are only used with *atom\_style ellipsoid or line or tri or body* and specify how many of the atoms are finite-size ellipsoids or lines or triangles or bodies; the remainder are point particles. See the discussion of *ellipsoidflag* and the *Ellipsoids* section below. See the discussion of *lineflag* and the *Lines* section below. See the discussion of *triangleflag* and the *Triangles* section below. See the discussion of *bodyflag* and the *Bodies* section below.

---

**Note:** For *atom\_style template*, the molecular topology (bonds,angles,etc) is contained in the molecule templates read-in by the *molecule* command. This means you cannot set the *bonds*, *angles*, etc header keywords in the data file, nor can you define *Bonds*, *Angles*, etc sections as discussed below. You can set the *bond types*, *angle types*, etc header keywords, though it is not necessary. If specified, they must match the maximum values defined in any of the template molecules.

---

### 16.107.7 Format of the body of a data file

These are the section keywords for the body of the file.

- *Atoms, Velocities, Masses, Ellipsoids, Lines, Triangles, Bodies* = atom-property sections
- *Bonds, Angles, Dihedrals, Improvers* = molecular topology sections
- *Atom Type Labels, Bond Type Labels, Angle Type Labels, Dihedral Type Labels, Improper Type Labels* = type label maps
- *Pair Coeffs, PairIJ Coeffs, Bond Coeffs, Angle Coeffs, Dihedral Coeffs, Improper Coeffs* = force field sections
- *BondBond Coeffs, BondAngle Coeffs, MiddleBondTorsion Coeffs, EndBondTorsion Coeffs, AngleTorsion Coeffs, AngleAngleTorsion Coeffs, BondBond13 Coeffs, AngleAngle Coeffs* = class 2 force field sections

These keywords will check an appended comment for a match with the currently defined style:

- *Atoms, Pair Coeffs, PairIJ Coeffs, Bond Coeffs, Angle Coeffs, Dihedral Coeffs, Improper Coeffs*

For example, these lines:

```
Atoms # sphere
Pair Coeffs # lj/cut
```

will check if the currently-defined *atom\_style* is *sphere*, and the current *pair\_style* is *lj/cut*. If not, LAMMPS will issue a warning to indicate that the data file section likely does not contain the correct number or type of parameters expected for the currently-defined style.

Each section is listed below in alphabetic order. The format of each section is described including the number of lines it must contain and rules (if any) for where it can appear in the data file.

Any individual line in the various sections can have a trailing comment starting with “#” for annotation purposes. There must be at least one blank between valid content and the comment. E.g. in the Atoms section:

```
10 1 17 -1.0 10.0 5.0 6.0 # salt ion
```

---

*Angle Coeffs* section:

- one line per angle type
- line syntax: ID coeffs

```
ID = angle type (1-N)
coeffs = list of coeffs
```

- example:

```
6 70 108.5 0 0
```

The number and meaning of the coefficients are specific to the defined angle style. See the *angle\_style* and *angle\_coeff* commands for details. Coefficients can also be set via the *angle\_coeff* command in the input script.

---

*Angle Type Labels* section:

- one line per angle type
- line syntax: ID label



```
ID = angle type (1-N)
label = alphanumeric type label
```

Define alphanumeric type labels for each numeric angle type. These can be used in the Angles section in place of a numeric type, but only if this section appears before the Angles section.

See the [Howto type labels](#) doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

*AngleAngle Coeffs* section:

- one line per improper type
  - line syntax: ID coeffs
- ```
ID = improper type (1-N)
coeffs = list of coeffs (see improper\_coeff)
```

*AngleAngleTorsion Coeffs* section:

- one line per dihedral type
  - line syntax: ID coeffs
- ```
ID = dihedral type (1-N)
coeffs = list of coeffs (see dihedral\_coeff)
```

*Angles* section:

- one line per angle
- line syntax: ID type atom1 atom2 atom3

```
ID = number of angle (1-Nangles)
type = angle type (1-Nangletype, or type label)
atom1,atom2,atom3 = IDs of 1st,2nd,3rd atom in angle
```

example:

```
2 2 17 29 430
```

The 3 atoms are ordered linearly within the angle. Thus the central atom (around which the angle is computed) is the atom2 in the list. E.g. H,O,H for a water molecule. The *Angles* section must appear after the *Atoms* section.

All values in this section must be integers (1, not 1.0). However, the type can be a numeric value or an alphanumeric label. The latter is only allowed if the type label has been defined by the *labelmap* command or an Angle Type Labels section earlier in the data file. See the [Howto type labels](#) doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

*AngleTorsion Coeffs* section:

- one line per dihedral type
- line syntax: ID coeffs

```
ID = dihedral type (1-N)
coeffs = list of coeffs (see dihedral_coeff)
```

---

*Atom Type Labels* section:

- one line per atom type
- line syntax: ID label

```
ID = numeric atom type (1-N)
label = alphanumeric type label
```

Define alphanumeric type labels for each numeric atom type. These can be used in the Atoms section in place of a numeric type, but only if the Atom Type Labels section appears before the Atoms section.

See the [Howto type labels](#) doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

---

*Atoms* section:

- one line per atom
- line syntax: depends on atom style

An *Atoms* section must appear in the data file if `natoms > 0` in the header section. The atoms can be listed in any order. These are the line formats for each *atom style* in LAMMPS. As discussed below, each line can optionally have 3 flags (`nx,ny,nz`) appended to it, which indicate which image of a periodic simulation box the atom is in. These may be important to include for some kinds of analysis.

angle	atom-ID molecule-ID atom-type x y z
atomic	atom-ID atom-type x y z
body	atom-ID atom-type bodyflag mass x y z
bond	atom-ID molecule-ID atom-type x y z
charge	atom-ID atom-type q x y z
dielectric	atom-ID atom-type q x y z normx normy normz area ed em epsilon curvature
dipole	atom-ID atom-type q x y z mux muy muz
dpd	atom-ID atom-type theta x y z
edpd	atom-ID atom-type edpd_temp edpd_cv x y z
electron	atom-ID atom-type q espin eradius x y z
ellipsoid	atom-ID atom-type ellipsoidflag density x y z
full	atom-ID molecule-ID atom-type q x y z
line	atom-ID molecule-ID atom-type lineflag density x y z
mdpd	atom-ID atom-type rho x y z
mesont	atom-ID molecule-ID atom-type bond_nt mass mradius mlength buckling x y z
molecular	atom-ID molecule-ID atom-type x y z
peri	atom-ID atom-type volume density x y z
smd	atom-ID atom-type molecule volume mass kradius cradius x0 y0 z0 x y z
sph	atom-ID atom-type rho esph cv x y z
sphere	atom-ID atom-type diameter density x y z
bpm/sphere	atom-ID molecule-ID atom-type diameter density x y z
spin	atom-ID atom-type x y z spx spy spz sp
tdpd	atom-ID atom-type x y z cc1 cc2 ... ccNspecies
template	atom-ID atom-type molecule-ID template-index template-atom x y z
tri	atom-ID molecule-ID atom-type triangleflag density x y z
wavepacket	atom-ID atom-type charge espin eradius etag cs_re cs_im x y z
hybrid	atom-ID atom-type x y z sub-style1 sub-style2 ...

The per-atom values have these meanings and units, listed alphabetically:

- atom-ID = integer ID of atom
- atom-type = type of atom (1-Ntype, or type label)
- bodyflag = 1 for body particles, 0 for point particles
- bond\_nt = bond NT factor for MESONT particles (?? units)
- buckling = buckling factor for MESONT particles (?? units)
- ccN = chemical concentration for tDPD particles for each species (mole/volume units)
- cradius = contact radius for SMD particles (distance units)
- cs\_re,cs\_im = real/imaginary parts of wave packet coefficients
- cv = heat capacity (need units) for SPH particles
- density = density of particle (mass/distance<sup>3</sup> or mass/distance<sup>2</sup> or mass/distance units, depending on dimensionality of particle)
- diameter = diameter of spherical atom (distance units)
- edpd\_temp = temperature for eDPD particles (temperature units)
- edpd\_cv = volumetric heat capacity for eDPD particles (energy/temperature/volume units)
- ellipsoidflag = 1 for ellipsoidal particles, 0 for point particles
- eradius = electron radius (or fixed-core radius)

- `esph` = energy (need units) for SPH particles
- `espin` = electron spin (+1/-1), 0 = nuclei, 2 = fixed-core, 3 = pseudo-cores (i.e. ECP)
- `etag` = integer ID of electron that each wave packet belongs to
- `kradius` = kernel radius for SMD particles (distance units)
- `lineflag` = 1 for line segment particles, 0 for point or spherical particles
- `mass` = mass of particle (mass units)
- `mlength` = ?? length for MESONT particles (distance units)
- `molecule-ID` = integer ID of molecule the atom belongs to
- `mradius` = ?? radius for MESONT particles (distance units)
- `mux,muy,muz` = components of dipole moment of atom (dipole units)
- `q` = charge on atom (charge units)
- `rho` = density (need units) for SPH particles
- `sp` = magnitude of magnetic spin of atom (Bohr magnetons)
- `spx,spy,spz` = components of magnetic spin of atom (unit vector)
- `template-atom` = which atom within a template molecule the atom is
- `template-index` = which molecule within the molecule template the atom is part of
- `theta` = internal temperature of a DPD particle
- `triangleflag` = 1 for triangular particles, 0 for point or spherical particles
- `volume` = volume of Peridynamic particle (distance<sup>3</sup> units)
- `x,y,z` = coordinates of atom (distance units)
- `x0,y0,z0` = original (strain-free) coordinates of atom (distance units)

The units for these quantities depend on the unit style; see the *units* command for details.

For 2d simulations specify `z` as 0.0, or a value within the *zlo zhi* setting in the data file header.

The atom-ID is used to identify the atom throughout the simulation and in dump files. Normally, it is a unique value from 1 to `Natoms` for each atom. Unique values larger than `Natoms` can be used, but they will cause extra memory to be allocated on each processor, if an atom map array is used, but not if an atom map hash is used; see the *atom\_modify* command for details. If an atom map is not used (e.g. an atomic system with no bonds), and you don't care if unique atom IDs appear in dump files, then the atom-IDs can all be set to 0.

The atom-type can be a numeric value or an alphanumeric label. The latter is only allowed if the type label has been defined by the *labelmap* command or an Atom Type Labels section earlier in the data file. See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

The molecule ID is a second identifier attached to an atom. Normally, it is a number from 1 to `N`, identifying which molecule the atom belongs to. It can be 0 if it is a non-bonded atom or if you don't care to keep track of molecule assignments.

The diameter specifies the size of a finite-size spherical particle. It can be set to 0.0, which means that atom is a point particle.

The *ellipsoidflag*, *lineflag*, *triangleflag*, and *bodyflag* determine whether the particle is a finite-size ellipsoid or line or triangle or body of finite size, or whether the particle is a point particle. Additional attributes must be defined for each ellipsoid, line, triangle, or body in the corresponding *Ellipsoids*, *Lines*, *Triangles*, or *Bodies* section.

The *template-index* and *template-atom* are only defined used by *atom\_style template*. In this case the *molecule* command is used to define a molecule template which contains one or more molecules (as separate files). If an atom belongs to one of those molecules, its *template-index* and *template-atom* are both set to positive integers; if not the values are both 0. The *template-index* is which molecule (1 to Nmols) the atom belongs to. The *template-atom* is which atom (1 to Natoms) within the molecule the atom is.

Some pair styles and fixes and computes that operate on finite-size particles allow for a mixture of finite-size and point particles. See the doc pages of individual commands for details.

For finite-size particles, the density is used in conjunction with the particle volume to set the mass of each particle as  $\text{mass} = \text{density} * \text{volume}$ . In this context, volume can be a 3d quantity (for spheres or ellipsoids), a 2d quantity (for triangles), or a 1d quantity (for line segments). If the volume is 0.0, meaning a point particle, then the density value is used as the mass. One exception is for the body atom style, in which case the mass of each particle (body or point particle) is specified explicitly. This is because the volume of the body is unknown.

Note that for 2d simulations of spheres, this command will treat them as spheres when converting density to mass. However, they can also be modeled as 2d discs (circles) if the *set density/disc* command is used to reset their mass after the *read\_data* command is used. A *disc* keyword can also be used with time integration fixes, such as *fix nve/sphere* and *fix nvt/sphere* to time integrate their motion as 2d discs (not 3d spheres), by changing their moment of inertia.

For *atom\_style hybrid*, following the 5 initial values (ID,type,x,y,z), specific values for each sub-style must be listed. The order of the sub-styles is the same as they were listed in the *atom\_style* command. The specific values for each sub-style are those that are not the 5 standard ones (ID,type,x,y,z). For example, for the “charge” sub-style, a “q” value would appear. For the “full” sub-style, a “molecule-ID” and “q” would appear. These are listed in the same order they appear as listed above. Thus if

```
atom_style hybrid charge sphere
```

were used in the input script, each atom line would have these fields:

```
atom-ID atom-type x y z q diameter density
```

Note that if a non-standard value is defined by multiple sub-styles, it only appears once in the atom line. E.g. the atom line for *atom\_style hybrid dipole full* would list “q” only once, with the dipole sub-style fields; “q” does not appear with the full sub-style fields.

```
atom-ID atom-type x y z q mux muy myz molecule-ID
```

Atom lines specify the (x,y,z) coordinates of atoms. These can be inside or outside the simulation box. When the data file is read, LAMMPS wraps coordinates outside the box back into the box for dimensions that are periodic. As discussed above, if an atom is outside the box in a non-periodic dimension, it will be lost.

LAMMPS always stores atom coordinates as values which are inside the simulation box. It also stores 3 flags which indicate which image of the simulation box (in each dimension) the atom would be in if its coordinates were unwrapped across periodic boundaries. An image flag of 0 means the atom is still inside the box when unwrapped. A value of 2 means add 2 box lengths to get the unwrapped coordinate. A value of -1 means subtract 1 box length to get the unwrapped coordinate. LAMMPS updates these flags as atoms cross periodic boundaries during the simulation. The *dump* command can output atom coordinates in wrapped or unwrapped form, as well as the 3 image flags.

In the data file, atom lines (all lines or none of them) can optionally list 3 trailing integer values (nx,ny,nz), which are used to initialize the atom’s image flags. If nx,ny,nz values are not listed in the data file, LAMMPS initializes them to 0. Note that the image flags are immediately updated if an atom’s coordinates need to wrapped back into the simulation box.

It is only important to set image flags correctly in a data file if a simulation model relies on unwrapped coordinates for some calculation; otherwise they can be left unspecified. Examples of LAMMPS commands that use unwrapped coordinates internally are as follows:

- Atoms in a rigid body (see *fix rigid*, *fix rigid/small*) must have consistent image flags, so that when the atoms are unwrapped, they are near each other, i.e. as a single body.
- If the *replicate* command is used to generate a larger system, image flags must be consistent for bonded atoms when the bond crosses a periodic boundary. I.e. the values of the image flags should be different by 1 (in the appropriate dimension) for the two atoms in such a bond.
- If you plan to *dump* image flags and perform post-analysis that will unwrap atom coordinates, it may be important that a continued run (restarted from a data file) begins with image flags that are consistent with the previous run.

---

**Note:** If your system is an infinite periodic crystal with bonds then it is impossible to have fully consistent image flags. This is because some bonds will cross periodic boundaries and connect two atoms with the same image flag.

---

Atom velocities and other atom quantities not defined above are set to 0.0 when the *Atoms* section is read. Velocities can be set later by a *Velocities* section in the data file or by a *velocity* or *set* command in the input script.

---

*Bodies* section:

- one or more lines per body
- first line syntax: atom-ID Ninteger Ndouble

```
Ninteger = # of integer quantities for this particle
Ndouble = # of floating-point quantities for this particle
```

- 0 or more integer lines with total of Ninteger values
- 0 or more double lines with total of Ndouble values
- example:

```
12 3 6
2 3 2
1.0 2.0 3.0 1.0 2.0 4.0
```

- example:

```
12 0 14
1.0 2.0 3.0 1.0 2.0 4.0 1.0
2.0 3.0 1.0 2.0 4.0 4.0 2.0
```

The *Bodies* section must appear if *atom\_style body* is used and any atoms listed in the *Atoms* section have a bodyflag = 1. The number of bodies should be specified in the header section via the “bodies” keyword.

Each body can have a variable number of integer and/or floating-point values. The number and meaning of the values is defined by the body style, as described in the *Howto body* doc page. The body style is given as an argument to the *atom\_style body* command.

The Ninteger and Ndouble values determine how many integer and floating-point values are specified for this particle. Ninteger and Ndouble can be as large as needed and can be different for every body. Integer values are then listed next on subsequent lines. Lines are read one at a time until Ninteger values are read. Floating-point values follow on subsequent lines, Again lines are read one at a time until Ndouble values are read. Note that if there are no values of a particular type, no lines appear for that type.

The *Bodies* section must appear after the *Atoms* section.

---

*Bond Coeffs* section:

- one line per bond type
- line syntax: ID coeffs

```
ID = bond type (1-N)
coeffs = list of coeffs
```

- example:

```
4 250 1.49
```

The number and meaning of the coefficients are specific to the defined bond style. See the *bond\_style* and *bond\_coeff* commands for details. Coefficients can also be set via the *bond\_coeff* command in the input script.

---

*Bond Type Labels* section:

- one line per bond type
- line syntax: ID label

```
ID = bond type (1-N)
label = alphanumeric type label
```

Define alphanumeric type labels for each numeric bond type. These can be used in the Bonds section in place of a numeric type, but only if this section appears before the Angles section.

See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

---

*BondAngle Coeffs* section:

- one line per angle type
- line syntax: ID coeffs

```
ID = angle type (1-N)
coeffs = list of coeffs (see class 2 section of angle_coeff)
```

*BondBond Coeffs* section:

- one line per angle type
- line syntax: ID coeffs

```
ID = angle type (1-N)
coeffs = list of coeffs (see class 2 section of angle_coeff)
```

*BondBond13 Coeffs* section:

- one line per dihedral type
- line syntax: ID coeffs

```
ID = dihedral type (1-N)
coeffs = list of coeffs (see class 2 section of dihedral_coeff)
```

*Bonds* section:

- one line per bond
- line syntax: ID type atom1 atom2

```
ID = bond number (1-Nbonds)
type = bond type (1-Nbondtype, or type label)
atom1,atom2 = IDs of 1st,2nd atom in bond
```

- example:

```
12 3 17 29
```

The *Bonds* section must appear after the *Atoms* section.

All values in this section must be integers (1, not 1.0). However, the type can be a numeric value or an alphanumeric label. The latter is only allowed if the type label has been defined by the *labelmap* command or a Bond Type Labels section earlier in the data file. See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

---

*Dihedral Coeffs* section:

- one line per dihedral type
- line syntax: ID coeffs

```
ID = dihedral type (1-N)
coeffs = list of coeffs
```

- example:

```
3 0.6 1 0 1
```

The number and meaning of the coefficients are specific to the defined dihedral style. See the *dihedral\_style* and *dihedral\_coeff* commands for details. Coefficients can also be set via the *dihedral\_coeff* command in the input script.

---

*Dihedral Type Labels* section:

- one line per dihedral type
- line syntax: ID label

```
ID = dihedral type (1-N)
label = alphanumeric type label
```

Define alphanumeric type labels for each numeric dihedral type. These can be used in the Dihedrals section in place of a numeric type, but only if this section appears before the Dihedrals section.

See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

---

*Dihedrals* section:



- one line per dihedral
- line syntax: ID type atom1 atom2 atom3 atom4

```
ID = number of dihedral (1-Ndihedrals)
type = dihedral type (1-Ndihedraltype, or type label)
atom1,atom2,atom3,atom4 = IDs of 1st,2nd,3rd,4th atom in dihedral
```

- example:

```
12 4 17 29 30 21
```

The 4 atoms are ordered linearly within the dihedral. The *Dihedrals* section must appear after the *Atoms* section.

All values in this section must be integers (1, not 1.0). However, the type can be a numeric value or an alphanumeric label. The latter is only allowed if the type label has been defined by the *labelmap* command or a Dihedral Type Labels section earlier in the data file. See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

*Ellipsoids* section:

- one line per ellipsoid
- line syntax: atom-ID shapex shapey shapez quatw quati quatj quatk

```
atom-ID = ID of atom which is an ellipsoid
shapex,shapey,shapez = 3 diameters of ellipsoid (distance units)
quatw,quati,quatj,quatk = quaternion components for orientation of atom
```

- example:

```
12 1 2 1 1 0 0 0
```

The *Ellipsoids* section must appear if *atom\_style ellipsoid* is used and any atoms are listed in the *Atoms* section with an ellipsoidflag = 1. The number of ellipsoids should be specified in the header section via the “ellipsoids” keyword.

The 3 shape values specify the 3 diameters or aspect ratios of a finite-size ellipsoidal particle, when it is oriented along the 3 coordinate axes. They must all be non-zero values.

The values *quatw*, *quati*, *quatj*, and *quatk* set the orientation of the atom as a quaternion (4-vector). Note that the shape attributes specify the aspect ratios of an ellipsoidal particle, which is oriented by default with its x-axis along the simulation box’s x-axis, and similarly for y and z. If this body is rotated (via the right-hand rule) by an angle theta around a unit vector (a,b,c), then the quaternion that represents its new orientation is given by (cos(theta/2), a\*sin(theta/2), b\*sin(theta/2), c\*sin(theta/2)). These 4 components are quatw, quati, quatj, and quatk as specified above. LAMMPS normalizes each atom’s quaternion in case (a,b,c) is not specified as a unit vector.

The *Ellipsoids* section must appear after the *Atoms* section.

*EndBondTorsion Coeffs* section:

- one line per dihedral type
- line syntax: ID coeffs

```
ID = dihedral type (1-N)
coeffs = list of coeffs (see class 2 section of dihedral_coeff)
```

*Improper Coeffs* section:

- one line per improper type
- line syntax: ID coeffs

```
ID = improper type (1-N)
coeffs = list of coeffs
```

- example:

```
2 20 0.0548311
```

The number and meaning of the coefficients are specific to the defined improper style. See the *improper\_style* and *improper\_coeff* commands for details. Coefficients can also be set via the *improper\_coeff* command in the input script.

---

*Improper Type Labels* section:

- one line per improper type
- line syntax: ID label

```
ID = improper type (1-N)
label = alphanumeric type label
```

Define alphanumeric type labels for each numeric improper type. These can be used in the Improvers section in place of a numeric type, but only if the this section appears before the Improvers section.

See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

---

*Improvers* section:

- one line per improper
- line syntax: ID type atom1 atom2 atom3 atom4

```
ID = number of improper (1-Nimprovers)
type = improper type (1-Nimproptype, or type label)
atom1,atom2,atom3,atom4 = IDs of 1st,2nd,3rd,4th atom in improper
```

- example:

```
12 3 17 29 13 100
```

The ordering of the 4 atoms determines the definition of the improper angle used in the formula for each *improper style*. See the doc pages for individual styles for details.

The *Improvers* section must appear after the *Atoms* section.

All values in this section must be integers (1, not 1.0). However, the type can be a numeric value or an alphanumeric label. The latter is only allowed if the type label has been defined by the *labelmap* command or a Improper Type Labels section earlier in the data file. See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

---

*Lines* section:

- one line per line segment
- line syntax: atom-ID x1 y1 x2 y2

```
atom-ID = ID of atom which is a line segment
x1,y1 = 1st end point
x2,y2 = 2nd end point
```

- example:

```
12 1.0 0.0 2.0 0.0
```

The *Lines* section must appear if *atom\_style line* is used and any atoms are listed in the *Atoms* section with a lineflag = 1. The number of lines should be specified in the header section via the “lines” keyword.

The 2 end points are the end points of the line segment. The ordering of the 2 points should be such that using a right-hand rule to cross the line segment with a unit vector in the +z direction, gives an “outward” normal vector perpendicular to the line segment. I.e. normal = (c2-c1) x (0,0,1). This orientation may be important for defining some interactions.

The *Lines* section must appear after the *Atoms* section.

---

*Masses* section:

- one line per atom type
- line syntax: ID mass

```
ID = atom type (1-N or atom type label)
mass = mass value
```

- example:

```
3 1.01
```

This defines the mass of each atom type. This can also be set via the *mass* command in the input script. This section cannot be used for atom styles that define a mass for individual atoms - e.g. *atom\_style sphere*.

Using type labels instead of atom type numbers is only allowed if the type label has been defined by the *labelmap* command or a Atom Type Labels section earlier in the data file. See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

---

*MiddleBondTorsion Coeffs* section:

- one line per dihedral type
- line syntax: ID coeffs

```
ID = dihedral type (1-N)
coeffs = list of coeffs (see class 2 section of dihedral_coeff)
```

*Pair Coeffs* section:

---

- one line per atom type
- line syntax: ID coeffs

```
ID = atom type (1-N)
coeffs = list of coeffs
```

- example:

```
3 0.022 2.35197 0.022 2.35197
```

The number and meaning of the coefficients are specific to the defined pair style. See the *pair\_style* and *pair\_coeff* commands for details. Since pair coefficients for types  $I \neq J$  are not specified, these will be generated automatically by the pair style's mixing rule. See the individual *pair\_style* doc pages and the *pair\_modify mix* command for details. Pair coefficients can also be set via the *pair\_coeff* command in the input script.

---

*PairIJ Coeffs* section:

- one line per pair of atom types for all I,J with  $I \leq J$
- line syntax: ID1 ID2 coeffs

```
ID1 = atom type I = 1-N
ID2 = atom type J = I-N, with I <= J
coeffs = list of coeffs
```

- examples:

```
3 3 0.022 2.35197 0.022 2.35197
3 5 0.022 2.35197 0.022 2.35197
```

This section must have  $N*(N+1)/2$  lines where  $N = \#$  of atom types. The number and meaning of the coefficients are specific to the defined pair style. See the *pair\_style* and *pair\_coeff* commands for details. Since pair coefficients for types  $I \neq J$  are all specified, these values will turn off the default mixing rule defined by the pair style. See the individual *pair\_style* doc pages and the *pair\_modify mix* command for details. Pair coefficients can also be set via the *pair\_coeff* command in the input script.

---

*Triangles* section:

- one line per triangle
- line syntax: atom-ID x1 y1 z1 x2 y2 z2 x3 y3 z3

```
atom-ID = ID of atom which is a line segment
x1,y1,z1 = 1st corner point
x2,y2,z2 = 2nd corner point
x3,y3,z3 = 3rd corner point
```

- example:

```
12 0.0 0.0 0.0 2.0 0.0 1.0 0.0 2.0 1.0
```

The *Triangles* section must appear if *atom\_style tri* is used and any atoms are listed in the *Atoms* section with a triangleflag = 1. The number of lines should be specified in the header section via the “triangles” keyword.

The 3 corner points are the corner points of the triangle. The ordering of the 3 points should be such that using a right-hand rule to go from point1 to point2 to point3 gives an “outward” normal vector to the face of the triangle. I.e.  $\text{normal} = (c_2 - c_1) \times (c_3 - c_1)$ . This orientation may be important for defining some interactions.

The *Triangles* section must appear after the *Atoms* section.

*Velocities* section:

- one line per atom
- line syntax: depends on atom style

all styles except those listed	atom-ID vx vy vz
electron	atom-ID vx vy vz ervel
ellipsoid	atom-ID vx vy vz lx ly lz
sphere	atom-ID vx vy vz wx wy wz
hybrid	atom-ID vx vy vz sub-style1 sub-style2 ...

where the keywords have these meanings:

vx,vy,vz = translational velocity of atom lx,ly,lz = angular momentum of aspherical atom wx,wy,wz = angular velocity of spherical atom ervel = electron radial velocity (0 for fixed-core):ul

The velocity lines can appear in any order. This section can only be used after an *Atoms* section. This is because the *Atoms* section must have assigned a unique atom ID to each atom so that velocities can be assigned to them.

Vx, vy, vz, and ervel are in *units* of velocity. Lx, ly, lz are in units of angular momentum (distance-velocity-mass). Wx, Wy, Wz are in units of angular velocity (radians/time).

For atom\_style hybrid, following the 4 initial values (ID,vx,vy,vz), specific values for each sub-style must be listed. The order of the sub-styles is the same as they were listed in the *atom\_style* command. The sub-style specific values are those that are not the 5 standard ones (ID,vx,vy,vz). For example, for the “sphere” sub-style, “wx”, “wy”, “wz” values would appear. These are listed in the same order they appear as listed above. Thus if

```
atom_style hybrid electron sphere
```

were used in the input script, each velocity line would have these fields:

```
atom-ID vx vy vz ervel wx wy wz
```

Translational velocities can also be set by the *velocity* command in the input script.

## 16.107.8 Restrictions

To read gzipped data files, you must compile LAMMPS with the -DLAMMPS\_GZIP option. See the *Build settings* doc page for details.

Label maps are currently not supported when using the KOKKOS package.

## 16.107.9 Related commands

*read\_dump*, *read\_restart*, *create\_atoms*, *write\_data*, *labelmap*

### 16.107.10 Default

The default for all the *extra* keywords is 0.

## 16.108 read\_dump command

### 16.108.1 Syntax

```
read_dump file Nstep field1 field2 ... keyword values ...
```

- file = name of dump file to read
- Nstep = snapshot timestep to read from file
- one or more fields may be appended
- zero or more keyword/value pairs may be appended
- keyword = *nfile* or *box* or *timestep* or *replace* or *purge* or *trim* or *add* or *label* or *scaled* or *wrapped* or *format*

*nfile* value = Nfiles = how many parallel dump files exist

*box* value = *yes* or *no* = replace simulation box with dump box

*timestep* value = *yes* or *no* = reset simulation timestep with dump timestep

*replace* value = *yes* or *no* = overwrite atoms with dump atoms

*purge* value = *yes* or *no* = delete all atoms before adding dump atoms

*trim* value = *yes* or *no* = trim atoms not in dump snapshot

*add* value = *yes* or *keep* or *no* = add new dump atoms to system

*label* value = field column

field = one of the listed fields or *id* or *type*

column = label on corresponding column in dump file

*scaled* value = *yes* or *no* = coords in dump file are scaled/unscaled

*wrapped* value = *yes* or *no* = coords in dump file are wrapped/unwrapped

*format* values = format of dump file, must be last keyword if used

*native* = native LAMMPS dump file

*xyz* = XYZ file

*adios* [timeout value] = dump file written by the *dump adios* command

timeout = specify waiting time for the arrival of the timestep when running  
↔concurrently.

The value is a float number and is interpreted in seconds.

*molfile* style path = VMD molfile plugin interface

style = *dcd* or *xyz* or others supported by molfile plugins

path = optional path for location of molfile plugins

## 16.108.2 Examples

```

read_dump dump.file 5000 x y z
read_dump dump.xyz 5 x y z box no format xyz
read_dump dump.xyz 10 x y z box no format molfile xyz "../plugins"
read_dump dump.dcd 0 x y z box yes format molfile dcd
read_dump dump.file 1000 x y z vx vy vz box yes format molfile lammstrj /usr/local/lib/
↳vmd/plugins/LINUXAMD64/plugins/molfile
read_dump dump.file 5000 x y vx vy trim yes
read_dump dump.file 5000 x y vx vy add yes box no timestep no
read_dump ../run7/dump.file.gz 10000 x y z box yes
read_dump dump.xyz 10 x y z box no format molfile xyz ../plugins
read_dump dump.dcd 0 x y z format molfile dcd
read_dump dump.file 1000 x y z vx vy vz format molfile lammstrj /usr/local/lib/vmd/
↳plugins/LINUXAMD64/plugins/molfile
read_dump dump.bp 5000 x y z vx vy vz format adios
read_dump dump.bp 5000 x y z vx vy vz format adios timeout 60.0

```

## 16.108.3 Description

Read atom information from a dump file to overwrite the current atom coordinates, and optionally the atom velocities and image flags, the simulation timestep, and the simulation box dimensions. This is useful for restarting a run from a particular snapshot in a dump file. See the [read\\_restart](#) and [read\\_data](#) commands for alternative methods to do this. Also see the [rerun](#) command for a means of reading multiple snapshots from a dump file.

Note that a simulation box must already be defined before using the `read_dump` command. This can be done by the [create\\_box](#), [read\\_data](#), or [read\\_restart](#) commands. The `read_dump` command can reset the simulation box dimensions, as explained below.

Also note that reading per-atom information from a dump snapshot is limited to the atom coordinates, velocities and image flags, as explained below. Other atom properties, which may be necessary to run a valid simulation, such as atom charge, or bond topology information for a molecular system, are not read from (or may not even be contained in) dump files. Thus this auxiliary information should be defined in the usual way, e.g. in a data file read in by a [read\\_data](#) command, before using the `read_dump` command, or by the [set](#) command, after the dump snapshot is read.

If the dump filename specified as *file* ends with “.gz”, the dump file is read in gzipped format.

You can read dump files that were written (in parallel) to multiple files via the “%” wild-card character in the dump file name. If any specified dump file name contains a “%”, they must all contain it. See the [dump](#) command for details. The “%” wild-card character is only supported by the *native* format for dump files, described next.

If reading parallel dump files, you must also use the *nfile* keyword to tell LAMMPS how many parallel files exist, via its specified *Nfiles* value.

The format of the dump file is selected through the *format* keyword. If specified, it must be the last keyword used, since all remaining arguments are passed on to the dump reader. The *native* format is for native LAMMPS dump files, written with a [dump\\_atom](#) or [dump\\_custom](#) command. The *xyz* format is for generic XYZ formatted dump files. These formats take no additional values.

The *molfile* format supports reading data through using the [VMD](#) molfile plugin interface. This dump reader format is only available, if the [MOLFILE](#) package has been installed when compiling LAMMPS.

The *molfile* format takes one or two additional values. The *style* value determines the file format to be used and can be any format that the molfile plugins support, such as DCD or XYZ. Note that DCD dump files can be written by LAMMPS via the [dump\\_dcd](#) command. The *path* value specifies a list of directories which LAMMPS will search

for the molfile plugins appropriate to the specified *style*. The syntax of the *path* value is like other search paths: it can contain multiple directories separated by a colon (or semi-colon on windows). The *path* keyword is optional and defaults to “.”, i.e. the current directory.

The *adios* format supports reading data that was written by the *dump adios* command. The entire dump is read in parallel across all the processes, dividing the atoms evenly among the processes. The number of writers that has written the dump file does not matter. Using the *adios* style for *dump* and *read\_dump* is a convenient way to dump all atoms from *N* writers and read it back by *M* readers. If one is running two LAMMPS instances concurrently where one dumps data and the other is reading it with the *rerun* command, the *timeout* option can be specified to wait on the reader side for the arrival of the requested step.

Support for other dump format readers may be added in the future.

---

Global information is first read from the dump file, namely timestep and box information.

The dump file is scanned for a snapshot with a timestamp that matches the specified *Nstep*. This means the LAMMPS timestep the dump file snapshot was written on for the *native* or *adios* formats.

The list of timestamps available in an *adios* .bp file is stored in the variable *ntimestep*:

```
$ bpls dump.bp -d ntimestep
uint64_t ntimestep 5*scalar
(0)      0 50 100 150 200
```

Note that the *xyz* and *molfile* formats do not store the timestep. For these formats, timesteps are numbered logically, in a sequential manner, starting from 0. Thus to access the 10th snapshot in an *xyz* or *molfile* formatted dump file, use *Nstep* = 9.

The dimensions of the simulation box for the selected snapshot are also read; see the *box* keyword discussion below. For the *native* format, an error is generated if the snapshot is for a triclinic box and the current simulation box is orthogonal or vice versa. A warning will be generated if the snapshot box boundary conditions (periodic, shrink-wrapped, etc) do not match the current simulation boundary conditions, but the boundary condition information in the snapshot is otherwise ignored. See the “boundary” command for more details. The *adios* reader does the same as the *native* format reader.

For the *xyz* format, no information about the box is available, so you must set the *box* flag to *no*. See details below.

For the *molfile* format, reading simulation box information is typically supported, but the location of the simulation box origin is lost and no explicit information about periodicity or orthogonal/triclinic box shape is available. The MOLFILE package makes a best effort to guess based on heuristics, but this may not always work perfectly.

---

Per-atom information from the dump file snapshot is then read from the dump file snapshot. This corresponds to the specified *fields* listed in the *read\_dump* command. It is an error to specify a z-dimension field, namely *z*, *vz*, or *iz*, for a 2d simulation.

For dump files in *native* format, each column of per-atom data has a text label listed in the file. A matching label for each field must appear, e.g. the label “vy” for the field *vy*. For the *x*, *y*, *z* fields any of the following labels are considered a match:

```
x, xs, xu, xsu for field x
y, ys, yu, ysu for field y
z, zs, zu, zsu for field z
```

The meaning of *xs* (scaled), *xu* (unwrapped), and *xsu* (scaled and unwrapped) is explained on the *dump* command doc page. These labels are searched for in the list of column labels in the dump file, in order, until a match is found.

The dump file must also contain atom IDs, with a column label of “id”.

---



If the *add* keyword is specified with a value of *yes* or *keep*, as discussed below, the dump file must contain atom types, with a column label of “type”.

If a column label you want to read from the dump file is not a match to a specified field, the *label* keyword can be used to specify the specific column label from the dump file to associate with that field. An example is if a time-averaged coordinate is written to the dump file via the *fix ave/atom* command. The column will then have a label corresponding to the fix-ID rather than “x” or “xs”. The *label* keyword can also be used to specify new column labels for fields *id* and *type*.

For dump files in *xyz* format, only the *x*, *y*, and *z* fields are supported. The dump file does not store atom IDs, so these are assigned consecutively to the atoms as they appear in the dump file, starting from 1. Thus you should insure that order of atoms is consistent from snapshot to snapshot in the XYZ dump file. See the *dump\_modify sort* command if the XYZ dump file was written by LAMMPS.

For dump files in *molfile* format, the *x*, *y*, *z*, *vx*, *vy*, and *vz* fields can be specified. However, not all molfile formats store velocities, or their respective plugins may not support reading of velocities. The molfile dump files do not store atom IDs, so these are assigned consecutively to the atoms as they appear in the dump file, starting from 1. Thus you should insure that order of atoms are consistent from snapshot to snapshot in the molfile dump file. See the *dump\_modify sort* command if the dump file was written by LAMMPS.

The *adios* format supports all fields that the *native* format supports except for the *q* charge field. The list of fields stored in an adios .bp file is recorded in the attributes *columns* (array of short strings) and *columnstr* (space-separated single string).

```
$ bpls -la dump.bp column*
  string    columns          attr = {"id", "type", "x", "y", "z", "vx", "vy", "vz"}
  string    columnstr       attr = "id type x y z vx vy vz "
```

Information from the dump file snapshot is used to overwrite or replace properties of the current system. There are various options for how this is done, determined by the specified fields and optional keywords.

The timestep of the snapshot becomes the current timestep for the simulation unless the *timestep* keyword is specified with a *no* value (default setting is *yes*). See the *reset\_timestep* command if you wish to change this to a different value after the dump snapshot is read.

If the *box* keyword is specified with a *yes* value, then the current simulation box dimensions are replaced by the dump snapshot box dimensions. If the *box* keyword is specified with a *no* value, the current simulation box is unchanged.

If the *purge* keyword is specified with a *yes* value, then all current atoms in the system are deleted before any of the operations invoked by the *replace*, *trim*, or *add* keywords take place.

If the *replace* keyword is specified with a *yes* value, then atoms with IDs that are in both the current system and the dump snapshot have their properties overwritten by field values. If the *replace* keyword is specified with a *no* value, atoms with IDs that are in both the current system and the dump snapshot are not modified.

If the *trim* keyword is specified with a *yes* value, then atoms with IDs that are in the current system but not in the dump snapshot are deleted. These atoms are unaffected if the *trim* keyword is specified with a *no* value.

If the *add* keyword is specified with a *no* value (default), then dump file atoms with IDs that are not in the current system are not added to the system. They are simply ignored.

If a *yes* value is specified, the atoms with new IDs are added to the system but their atom IDs are not preserved. Instead, after all the atoms are added, new IDs are assigned to them in the same manner as is described for the *create\_atoms* command. Basically the largest existing atom ID in the system is identified, and all the added atoms are assigned IDs that consecutively follow the largest ID.

If a *keep* value is specified, the atoms with new IDs are added to the system and their atom IDs are preserved. This may lead to non-contiguous IDs for the combined system.

Note that atoms added via the *add* keyword will only have the attributes read from the dump file due to the *field* arguments. For example, if *x* or *y* or *z* or *q* is not specified as a field, a value of 0.0 is used for added atoms. Added atoms must have an atom type, so this value must appear in the dump file.

Any other attributes (e.g. charge or particle diameter for spherical particles) will be set to default values, the same as if the *create\_atoms* command were used.

---

Atom coordinates read from the dump file are first converted into unscaled coordinates, relative to the box dimensions of the snapshot. These coordinates are then be assigned to an existing or new atom in the current simulation. The coordinates will then be remapped to the simulation box, whether it is the original box or the dump snapshot box. If periodic boundary conditions apply, this means the atom will be remapped back into the simulation box if necessary. If shrink-wrap boundary conditions apply, the new coordinates may change the simulation box dimensions. If fixed boundary conditions apply, the atom will be lost if it is outside the simulation box.

For *native* format dump files, the 3 xyz image flags for an atom in the dump file are set to the corresponding values appearing in the dump file if the *ix*, *iy*, *iz* fields are specified. If not specified, the image flags for replaced atoms are not changed and image flags for new atoms are set to default values. If coordinates read from the dump file are in unwrapped format (e.g. *xu*) then the image flags for read-in atoms are also set to default values. The remapping procedure described in the previous paragraph will then change images flags for all atoms (old and new) if periodic boundary conditions are applied to remap an atom back into the simulation box.

---

**Note:** If you get a warning about inconsistent image flags after reading in a dump snapshot, it means one or more pairs of bonded atoms now have inconsistent image flags. As discussed on the [Errors common](#) page this may or may not cause problems for subsequent simulations. One way this can happen is if you read image flag fields from the dump file but do not also use the dump file box parameters.

---

LAMMPS knows how to compute unscaled and remapped coordinates for the snapshot column labels discussed above, e.g. *x*, *xs*, *xu*, *xsu*. If another column label is assigned to the *x* or *y* or *z* field via the *label* keyword, e.g. for coordinates output by the *fix ave/atom* command, then LAMMPS needs to know whether the coordinate information in the dump file is scaled and/or wrapped. This can be set via the *scaled* and *wrapped* keywords. Note that the value of the *scaled* and *wrapped* keywords is ignored for fields *x* or *y* or *z* if the *label* keyword is not used to assign a column label to that field.

The scaled/unscaled and wrapped/unwrapped setting must be identical for any of the *x*, *y*, *z* fields that are specified. Thus you cannot read *xs* and *yu* from the dump file. Also, if the dump file coordinates are scaled and the simulation box is triclinic, then all 3 of the *x*, *y*, *z* fields must be specified, since they are all needed to generate absolute, unscaled coordinates.

---

### 16.108.4 Restrictions

To read gzipped dump files, you must compile LAMMPS with the `-DLAMMPS_GZIP` option. See the [Build settings](#) doc page for details.

The *molfile* dump file formats are part of the MOLFILE package. They are only enabled if LAMMPS was built with that packages. See the [Build package](#) page for more info.

To write and read adios .bp files, you must compile LAMMPS with the [ADIOS](#) package.

## 16.108.5 Related commands

*dump*, *dump molfile*, *dump adios*, *read\_data*, *read\_restart*, *rerun*

## 16.108.6 Default

The option defaults are box = yes, timestep = yes, replace = yes, purge = no, trim = no, add = no, scaled = no, wrapped = yes, and format = native.

## 16.109 read\_restart command

### 16.109.1 Syntax

```
read_restart file flag
```

- file = name of binary restart file to read in

### 16.109.2 Examples

```
read_restart save.10000
read_restart restart.*
read_restart restart.*.mpio
```

### 16.109.3 Description

Read in a previously saved system configuration from a restart file. This allows continuation of a previous run. Details about what information is stored (and not stored) in a restart file is given below. Basically this operation will re-create the simulation box with all its atoms and their attributes as well as some related global settings, at the point in time it was written to the restart file by a previous simulation. The simulation box will be partitioned into a regular 3d grid of rectangular bricks, one per processor, based on the number of processors in the current simulation and the settings of the *processors* command. The partitioning can later be changed by the *balance* or *fix balance* commands.

Restart files are saved in binary format to enable exact restarts, meaning that the trajectories of a restarted run will precisely match those produced by the original run had it continued on.

The binary restart file format was not designed with backward, forward, or cross-platform compatibility in mind, so the files are only expected to be read correctly by the same LAMMPS executable on the same platform. Changes to the architecture, compilation settings, or LAMMPS version can render a restart file unreadable or it may read the data incorrectly. If you want a more portable format, you can use the data file format as created by the *write\_data* command. Binary restart files can also be converted into a data file from the command line by the LAMMPS executable that wrote them using the *-restart2data* command line flag.

Several things can prevent exact restarts due to round-off effects, in which case the trajectories in the 2 runs will slowly diverge. These include running on a different number of processors or changing certain settings such as those set by the *newton* or *processors* commands. LAMMPS will issue a warning in these cases.

Certain fixes will not restart exactly, though they should provide statistically similar results. These include *fix shake* and *fix langevin*.

Certain pair styles will not restart exactly, though they should provide statistically similar results. This is because the forces they compute depend on atom velocities, which are used at half-step values every timestep when forces are

computed. When a run restarts, forces are initially evaluated with a full-step velocity, which is different than if the run had continued. These pair styles include *granular pair styles*, *pair dpd*, and *pair lubricate*.

If a restarted run is immediately different than the run which produced the restart file, it could be a LAMMPS bug, so consider *reporting it* if you think the behavior is a bug.

Because restart files are binary, they may not be portable to other machines. In this case, you can use the *-restart command-line switch* to convert a restart file to a data file.

Similar to how restart files are written (see the *write\_restart* and *restart* commands), the restart filename can contain two wild-card characters. If a “\*” appears in the filename, the directory is searched for all filenames that match the pattern where “\*” is replaced with a timestep value. The file with the largest timestep value is read in. Thus, this effectively means, read the latest restart file. It’s useful if you want your script to continue a run from where it left off. See the *run* command and its “upto” option for how to specify the run command so it does not need to be changed either.

If a “%” character appears in the restart filename, LAMMPS expects a set of multiple files to exist. The *restart* and *write\_restart* commands explain how such sets are created. *read\_restart* will first read a filename where “%” is replaced by “base”. This file tells LAMMPS how many processors created the set and how many files are in it. *read\_restart* then reads the additional files. For example, if the restart file was specified as *save.%* when it was written, then *read\_restart* reads the files *save.base*, *save.0*, *save.1*, ... *save.P-1*, where P is the number of processors that created the restart file.

Note that P could be the total number of processors in the previous simulation, or some subset of those processors, if the *fileper* or *nfile* options were used when the restart file was written; see the *restart* and *write\_restart* commands for details. The processors in the current LAMMPS simulation share the work of reading these files; each reads a roughly equal subset of the files. The number of processors which created the set can be different the number of processors in the current LAMMPS simulation. This can be a fast mode of input on parallel machines that support parallel I/O.

A restart file can also be read in parallel as one large binary file via the MPI-IO library, assuming it was also written with MPI-IO. MPI-IO is part of the MPI standard for versions 2.0 and above. Using MPI-IO requires two steps. First, build LAMMPS with its MPIIO package installed, e.g.

```
make yes-mpiio    # installs the MPIIO package
make mpi         # build LAMMPS for your platform
```

Second, use a restart filename which contains “.mpiio”. Note that it does not have to end in “.mpiio”, just contain those characters. Unlike MPI-IO dump files, a particular restart file must be both written and read using MPI-IO.

---

Here is the list of information included in a restart file, which means these quantities do not need to be re-specified in the input script that reads the restart file, though you can redefine many of these settings after the restart file is read.

- *units*
- *newton bond* (see discussion of *newton* command below)
- *atom style* and *atom\_modify* settings *id*, *map*, *sort*
- *comm style* and *comm\_modify* settings *mode*, *cutoff*, *vel*
- *timestep*
- simulation box size and shape and *boundary* settings
- atom *group* definitions
- per-type atom settings such as *mass*
- per-atom attributes including their group assignments and molecular topology attributes (bonds, angles, etc)
- force field styles (*pair*, *bond*, *angle*, etc)
- force field coefficients (*pair*, *bond*, *angle*, etc) in some cases (see below)

- *pair\_modify* settings, except the compute option
- *special\_bonds* settings

Here is a list of information not stored in a restart file, which means you must re-issue these commands in your input script, after reading the restart file.

- *newton pair* (see discussion of newton command below)
- *fix* commands (see below)
- *compute* commands (see below)
- *variable* commands
- *region* commands
- *neighbor list* criteria including *neigh\_modify* settings
- *kpace\_style* and *kpace\_modify* settings
- info for *thermodynamic*, *dump*, or *restart* output

The *newton* command has two settings, one for pairwise interactions, the other for bonded. Both settings are stored in the restart file. For the bond setting, the value in the file will overwrite the current value (at the time the read\_restart command is issued) and warn if the two values are not the same and the current value is not the default. For the pair setting, the value in the file will not overwrite the current value (so that you can override the previous run's value), but a warning is issued if the two values are not the same and the current value is not the default.

Note that some force field styles (pair, bond, angle, etc) do not store their coefficient info in restart files. Typically these are many-body or tabulated potentials which read their parameters from separate files. In these cases you will need to re-specify the *pair\_coeff*, *bond\_coeff*, etc commands in your restart input script. The doc pages for individual force field styles mention if this is the case. This is also true of *pair\_style hybrid* (bond hybrid, angle hybrid, etc) commands; they do not store coefficient info.

As indicated in the above list, the *fixes* used for a simulation are not stored in the restart file. This means the new input script should specify all fixes it will use. However, note that some fixes store an internal "state" which is written to the restart file. This allows the fix to continue on with its calculations in a restarted simulation. To re-enable such a fix, the fix command in the new input script must be of the same style and use the same fix-ID as was used in the input script that wrote the restart file.

If a match is found, LAMMPS prints a message indicating that the fix is being re-enabled. If no match is found before the first run or minimization is performed by the new script, the "state" information for the saved fix is discarded. At the time the discard occurs, LAMMPS will also print a list of fixes for which the information is being discarded. See the doc pages for individual fixes for info on which ones can be restarted in this manner. Note that fixes which are created internally by other LAMMPS commands (computes, fixes, etc) will have style names which are all-capitalized, and IDs which are generated internally.

Likewise, the *computes* used for a simulation are not stored in the restart file. This means the new input script should specify all computes it will use. However, some computes create a fix internally to store "state" information that persists from timestep to timestep. An example is the *compute msd* command which uses a fix to store a reference coordinate for each atom, so that a displacement can be calculated at any later time. If the compute command in the new input script uses the same compute-ID and group-ID as was used in the input script that wrote the restart file, then it will create the same fix in the restarted run. This means the re-created fix will be re-enabled with the stored state information as described in the previous paragraph, so that the compute can continue its calculations in a consistent manner.

---

**Note:** There are a handful of commands which can be used before or between runs which may require a system initialization. Examples include the "balance", "displace\_atoms", "delete\_atoms", "set" (some options), and "velocity" (some options) commands. This is because they can migrate atoms to new processors. Thus they will also discard unused "state" information from fixes. You will know the discard has occurred because a list of discarded fixes will be

printed to the screen and log file, as explained above. This means that if you wish to retain that info in a restarted run, you must re-specify the relevant fixes and computes (which create fixes) before those commands are used.

---

Some pair styles, like the *granular pair styles*, also use a fix to store “state” information that persists from timestep to timestep. In the case of granular potentials, it is contact information between pairs of touching particles. This info will also be re-enabled in the restart script, assuming you re-use the same granular pair style.

LAMMPS allows bond interactions (angle, etc) to be turned off or deleted in various ways, which can affect how their info is stored in a restart file.

If bonds (angles, etc) have been turned off by the *fix shake* or *delete\_bonds* command, their info will be written to a restart file as if they are turned on. This means they will need to be turned off again in a new run after the restart file is read.

Bonds that are broken (e.g. by a bond-breaking potential) are written to the restart file as broken bonds with a type of 0. Thus these bonds will still be broken when the restart file is read.

Bonds that have been broken by the *fix bond/break* command have disappeared from the system. No information about these bonds is written to the restart file.

---

## 16.109.4 Restrictions

To write and read restart files in parallel with MPI-IO, the MPIIO package must be installed.

## 16.109.5 Related commands

*read\_data, read\_dump, write\_restart, restart*

## 16.109.6 Default

none

## 16.110 region command

### 16.110.1 Syntax

```
region ID style args keyword arg ...
```

- ID = user-assigned name for the region
- style = *delete* or *block* or *cone* or *cylinder* or *ellipsoid* or *plane* or *prism* or *sphere* or *union* or *intersect*

*delete* = no args

*block* args = xlo xhi ylo yhi zlo zhi

xlo,xhi,ylo,yhi,zlo,zhi = bounds of block in all dimensions (distance units)

*cone* args = dim c1 c2 radlo radhi lo hi

dim = x or y or z = axis of cone

c1,c2 = coords of cone axis in other 2 dimensions (distance units)

radlo,radhi = cone radii at lo and hi end (distance units)

lo,hi = bounds of cone in dim (distance units)

```

cylinder args = dim c1 c2 radius lo hi
  dim = x or y or z = axis of cylinder
  c1,c2 = coords of cylinder axis in other 2 dimensions (distance units)
  radius = cylinder radius (distance units)
  c1,c2, and radius can be a variable (see below)
  lo,hi = bounds of cylinder in dim (distance units)
ellipsoid args = x y z a b c
  x,y,z = center of ellipsoid (distance units)
  a,b,c = half the length of the principal axes of the ellipsoid (distance units)
  x,y,z,a,b and c can be a variable (see below)
plane args = px py pz nx ny nz
  px,py,pz = point on the plane (distance units)
  nx,ny,nz = direction normal to plane (distance units)
prism args = xlo xhi ylo yhi zlo zhi xy xz yz
  xlo,xhi,ylo,yhi,zlo,zhi = bounds of untilted prism (distance units)
  xy = distance to tilt y in x direction (distance units)
  xz = distance to tilt z in x direction (distance units)
  yz = distance to tilt z in y direction (distance units)
sphere args = x y z radius
  x,y,z = center of sphere (distance units)
  radius = radius of sphere (distance units)
  x,y,z, and radius can be a variable (see below)
union args = N reg-ID1 reg-ID2 ...
  N = # of regions to follow, must be 2 or greater
  reg-ID1,reg-ID2, ... = IDs of regions to join together
intersect args = N reg-ID1 reg-ID2 ...
  N = # of regions to follow, must be 2 or greater
  reg-ID1,reg-ID2, ... = IDs of regions to intersect

```

- zero or more keyword/arg pairs may be appended
- keyword = *side* or *units* or *move* or *rotate* or *open*

```

side value = in or out
  in = the region is inside the specified geometry
  out = the region is outside the specified geometry
units value = lattice or box
  lattice = the geometry is defined in lattice units
  box = the geometry is defined in simulation box units
move args = v_x v_y v_z
  v_x,v_y,v_z = equal-style variables for x,y,z displacement of region over time.
  ↪(distance units)
rotate args = v_theta Px Py Pz Rx Ry Rz
  v_theta = equal-style variable for rotaton of region over time (in radians)
  Px,Py,Pz = origin for axis of rotation (distance units)
  Rx,Ry,Rz = axis of rotation vector
open value = integer from 1-6 corresponding to face index (see below)

```

- accelerated styles (with same args) = *block/kk*

## 16.110.2 Examples

```
region 1 block -3.0 5.0 INF 10.0 INF INF
region 2 sphere 0.0 0.0 0.0 5 side out
region void cylinder y 2 3 5 -5.0 EDGE units box
region 1 prism 0 10 0 10 0 10 2 0 0
region outside union 4 side1 side2 side3 side4
region 2 sphere 0.0 0.0 0.0 5 side out move v_left v_up NULL
region openbox block 0 10 0 10 0 10 open 5 open 6 units box
region funnel cone z 10 10 2 5 0 10 open 1 units box
```

## 16.110.3 Description

This command defines a geometric region of space. Various other commands use regions. For example, the region can be filled with atoms via the *create\_atoms* command. Or a bounding box around the region, can be used to define the simulation box via the *create\_box* command. Or the atoms in the region can be identified as a group via the *group* command, or deleted via the *delete\_atoms* command. Or the surface of the region can be used as a boundary wall via the *fix wall/region* command.

Commands which use regions typically test whether an atom's position is contained in the region or not. For this purpose, coordinates exactly on the region boundary are considered to be interior to the region. This means, for example, for a spherical region, an atom on the sphere surface would be part of the region if the sphere were defined with the *side in* keyword, but would not be part of the region if it were defined using the *side out* keyword. See more details on the *side* keyword below.

Normally, regions in LAMMPS are “static”, meaning their geometric extent does not change with time. If the *move* or *rotate* keyword is used, as described below, the region becomes “dynamic”, meaning it's location or orientation changes with time. This may be useful, for example, when thermostating a region, via the *compute temp/region* command, or when the *fix wall/region* command uses a region surface as a bounding wall on particle motion, i.e. a rotating container.

The *delete* style removes the named region. Since there is little overhead to defining extra regions, there is normally no need to do this, unless you are defining and discarding large numbers of regions in your input script.

The lo/hi values for *block* or *cone* or *cylinder* or *prism* styles can be specified as EDGE or INF. EDGE means they extend all the way to the global simulation box boundary. Note that this is the current box boundary; if the box changes size during a simulation, the region does not. INF means a large negative or positive number (1.0e20), so it should encompass the simulation box even if it changes size. If a region is defined before the simulation box has been created (via *create\_box* or *read\_data* or *read\_restart* commands), then an EDGE or INF parameter cannot be used. For a *prism* region, a non-zero tilt factor in any pair of dimensions cannot be used if both the lo/hi values in either of those dimensions are INF. E.g. if the xy tilt is non-zero, then xlo and xhi cannot both be INF, nor can ylo and yhi.

---

**Note:** Regions in LAMMPS do not get wrapped across periodic boundaries, as specified by the *boundary* command. For example, a spherical region that is defined so that it overlaps a periodic boundary is not treated as 2 half-spheres, one on either side of the simulation box.

---

---

**Note:** Regions in LAMMPS are always 3d geometric objects, regardless of whether the *dimension* of a simulation is 2d or 3d. Thus when using regions in a 2d simulation, you should be careful to define the region so that its intersection with the 2d x-y plane of the simulation has the 2d geometric extent you want.

---

For style *cone*, an axis-aligned cone is defined which is like a *cylinder* except that two different radii (one at each end) can be defined. Either of the radii (but not both) can be 0.0.



For style *cone* and *cylinder*, the  $c1,c2$  params are coordinates in the 2 other dimensions besides the cylinder axis dimension. For  $\text{dim} = x$ ,  $c1/c2 = y/z$ ; for  $\text{dim} = y$ ,  $c1/c2 = x/z$ ; for  $\text{dim} = z$ ,  $c1/c2 = x/y$ . Thus the third example above specifies a cylinder with its axis in the  $y$ -direction located at  $x = 2.0$  and  $z = 3.0$ , with a radius of 5.0, and extending in the  $y$ -direction from -5.0 to the upper box boundary.

For style *ellipsoid*, an axis-aligned ellipsoid is defined. The ellipsoid has its center at  $(x,y,z)$  and is defined by 3 axis-aligned vectors given by  $A = (a,0,0)$ ;  $B = (0,b,0)$ ;  $C = (0,0,c)$ . Note that although the ellipsoid is specified as axis-aligned it can be rotated via the optional *rotate* keyword.

For style *plane*, a plane is defined which contain the point  $(px,py,pz)$  and has a normal vector  $(nx,ny,nz)$ . The normal vector does not have to be of unit length. The “inside” of the plane is the half-space in the direction of the normal vector; see the discussion of the *side* option below.

For style *prism*, a parallelepiped is defined (it’s too hard to spell parallelepiped in an input script!). The parallelepiped has its “origin” at  $(xlo,ylo,zlo)$  and is defined by 3 edge vectors starting from the origin given by  $A = (xhi-xlo,0,0)$ ;  $B = (xy,yhi-ylo,0)$ ;  $C = (xz,yz,zhi-zlo)$ .  $Xy,xz,yz$  can be 0.0 or positive or negative values and are called “tilt factors” because they are the amount of displacement applied to faces of an originally orthogonal box to transform it into the parallelepiped.

A prism region that will be used with the *create\_box* command to define a triclinic simulation box must have tilt factors  $(xy,xz,yz)$  that do not skew the box more than half the distance of corresponding the parallel box length. For example, if  $xlo = 2$  and  $xhi = 12$ , then the  $x$  box length is 10 and the  $xy$  tilt factor must be between -5 and 5. Similarly, both  $xz$  and  $yz$  must be between  $-(xhi-xlo)/2$  and  $+(yhi-ylo)/2$ . Note that this is not a limitation, since if the maximum tilt factor is 5 (as in this example), then configurations with tilt =  $\dots, -15, -5, 5, 15, 25, \dots$  are all geometrically equivalent.

For style *sphere*, a sphere is defined with its center at  $(x,y,z)$  and with radius as its radius.

The *radius* value for styles *sphere* and *cylinder*, and the parameters  $a,b,c$  for style *ellipsoid*, can each be specified as an equal-style *variable*. Likewise, for style *sphere* and *ellipsoid* the  $x$ -,  $y$ -, and  $z$ - coordinates of the center of the sphere/ellipsoid can be specified as an equal-style variable. And for style *cylinder* the two center positions  $c1$  and  $c2$  for the location of the cylinder axes can be specified as a equal-style variable.

If the value is a variable, it should be specified as  $v\_name$ , where  $name$  is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the radius of the region.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent radius or have a time dependent position of the sphere or cylinder region.

See the [Howto tricilinc](#) page for a geometric description of triclinic boxes, as defined by LAMMPS, and how to transform these parameters to and from other commonly used triclinic representations.

The *union* style creates a region consisting of the volume of all the listed regions combined. The *intersect* style creates a region consisting of the volume that is common to all the listed regions.

---

**Note:** The *union* and *intersect* regions operate by invoking methods from their list of sub-regions. Thus you cannot delete the sub-regions after defining a *union* or *intersection* region.

---

The *side* keyword determines whether the region is considered to be inside or outside of the specified geometry. Using this keyword in conjunction with *union* and *intersect* regions, complex geometries can be built up. For example, if the interior of two spheres were each defined as regions, and a *union* style with *side* = out was constructed listing the region-IDs of the 2 spheres, the resulting region would be all the volume in the simulation box that was outside both of the spheres.

The *units* keyword determines the meaning of the distance units used to define the region for any argument above listed as having distance units. It also affects the scaling of the velocity vector specified with the *vel* keyword, the amplitude

vector specified with the *wiggle* keyword, and the rotation point specified with the *rotate* keyword, since they each involve a distance metric.

A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacings which are used as follows:

- For style *block*, the lattice spacing in dimension x is applied to xlo and xhi, similarly the spacings in dimensions y,z are applied to ylo/yhi and zlo/zhi.
- For style *cone*, the lattice spacing in argument *dim* is applied to lo and hi. The spacings in the two radial dimensions are applied to c1 and c2. The two cone radii are scaled by the lattice spacing in the dimension corresponding to c1.
- For style *cylinder*, the lattice spacing in argument *dim* is applied to lo and hi. The spacings in the two radial dimensions are applied to c1 and c2. The cylinder radius is scaled by the lattice spacing in the dimension corresponding to c1.
- For style *ellipsoid*, the lattice spacing in dimensions x,y,z are applied to the ellipsoid center x,y,z. The spacing in dimensions x,y,z are applied to the ellipsoid radii a,b,c respectively.
- For style *plane*, the lattice spacing in dimension x is applied to px and nx, similarly the spacings in dimensions y,z are applied to py/ny and pz/nz.
- For style *prism*, the lattice spacing in dimension x is applied to xlo and xhi, similarly for ylo/yhi and zlo/zhi. The lattice spacing in dimension x is applied to xy and xz, and the spacing in dimension y to yz.
- For style *sphere*, the lattice spacing in dimensions x,y,z are applied to the sphere center x,y,z. The spacing in dimension x is applied to the sphere radius.

---

If the *move* or *rotate* keywords are used, the region is “dynamic”, meaning its location or orientation changes with time. These keywords cannot be used with a *union* or *intersect* style region. Instead, the keywords should be used to make the individual sub-regions of the *union* or *intersect* region dynamic. Normally, each sub-region should be “dynamic” in the same manner (e.g. rotate around the same point), though this is not a requirement.

The *move* keyword allows one or more *equal-style variables* to be used to specify the x,y,z displacement of the region, typically as a function of time. A variable is specified as v\_name, where name is the variable name. Any of the three variables can be specified as NULL, in which case no displacement is calculated in that dimension.

Note that equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a region displacement that change as a function of time or spans consecutive runs in a continuous fashion. For the latter, see the *start* and *stop* keywords of the *run* command and the *elaplong* keyword of *thermo\_style custom* for details.

For example, these commands would displace a region from its initial position, in the positive x direction, effectively at a constant velocity:

```
variable dx equal ramp(0,10)
region 2 sphere 10.0 10.0 0.0 5 move v_dx NULL NULL
```

Note that the initial displacement is 0.0, though that is not required.

Either of these variables would “wiggle” the region back and forth in the y direction:

```
variable dy equal swiggle(0,5,100)
variable dysame equal 5*sin(2*PI*elaplong*dt/100)
region 2 sphere 10.0 10.0 0.0 5 move NULL v_dy NULL
```

The *rotate* keyword rotates the region around a rotation axis  $R = (R_x, R_y, R_z)$  that goes through a point  $P = (P_x, P_y, P_z)$ . The rotation angle is calculated, presumably as a function of time, by a variable specified as *v\_theta*, where *theta* is the variable name. The variable should generate its result in radians. The direction of rotation for the region around the rotation axis is consistent with the right-hand rule: if your right-hand thumb points along  $R$ , then your fingers wrap around the axis in the direction of rotation.

The *move* and *rotate* keywords can be used together. In this case, the displacement specified by the *move* keyword is applied to the  $P$  point of the *rotate* keyword.

---

The *open* keyword can be used (multiple times) to indicate that one or more faces of the region are ignored for purposes of particle/wall interactions. This keyword is only relevant for regions used by the *fix wall/region* and *fix wall/gran/region* commands. It can be used to create “open” containers where only some of the region faces are walls. For example, a funnel can be created with a *cone* style region that has an open face at the smaller radius for particles to flow out, or at the larger radius for pouring particles into the cone, or both.

Note that using the *open* keyword partly overrides the *side* keyword, since both exterior and interior surfaces of an open region are tested for particle contacts. The exception to this is a *union* or *intersect* region which includes an open sub-region. In that case the *side* keyword is still used to define the union/intersect region volume, and the *open* settings are only applied to the individual sub-regions that use them.

The indices specified as part of the *open* keyword have the following meanings:

For style *block*, indices 1-6 correspond to the *xlo*, *xhi*, *ylo*, *yhi*, *zlo*, *zhi* surfaces of the block. I.e. 1 is the *yz* plane at  $x = xlo$ , 2 is the *yz*-plane at  $x = xhi$ , 3 is the *xz* plane at  $y = ylo$ , 4 is the *xz* plane at  $y = yhi$ , 5 is the *xy* plane at  $z = zlo$ , 6 is the *xy* plane at  $z = zhi$ ). In the second-to-last example above, the region is a box open at both *xy* planes.

For style *prism*, values 1-6 have the same mapping as for style *block*. I.e. in an untilted *prism*, *open* indices correspond to the *xlo*, *xhi*, *ylo*, *yhi*, *zlo*, *zhi* surfaces.

For style *cylinder*, index 1 corresponds to the flat end cap at the low coordinate along the cylinder axis, index 2 corresponds to the high-coordinate flat end cap along the cylinder axis, and index 3 is the curved cylinder surface. For example, a *cylinder* region with *open 1 open 2* keywords will be open at both ends (e.g. a section of pipe), regardless of the cylinder orientation.

For style *cone*, the mapping is the same as for style *cylinder*. Index 1 is the low-coordinate flat end cap, index 2 is the high-coordinate flat end cap, and index 3 is the curved cone surface. In the last example above, a *cone* region is defined along the *z*-axis that is open at the *zlo* value (e.g. for use as a funnel).

For all other styles, the *open* keyword is ignored. As indicated above, this includes the *intersect* and *union* regions, though their sub-regions can be defined with the *open* keyword.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

**Note:** Currently, only *block* style regions are supported by Kokkos. The code using the region (such as a *fix* or *compute*) must also be supported by Kokkos or no acceleration will occur.

---

#### 16.110.4 Restrictions

A prism cannot be of 0.0 thickness in any dimension; use a small z thickness for 2d simulations. For 2d simulations, the xz and yz parameters must be 0.0.

#### 16.110.5 Related commands

*lattice, create\_atoms, delete\_atoms, group*

#### 16.110.6 Default

The option defaults are side = in, units = lattice, and no move or rotation.

### 16.111 replicate command

#### 16.111.1 Syntax

```
replicate nx ny nz keyword
```

nx,ny,nz = replication factors in each dimension

- optional *keyword* = *bbox*

*bbox* = only check atoms in replicas that overlap with a processor's sub-domain

#### 16.111.2 Examples

```
replicate 2 3 2
```

#### 16.111.3 Description

Replicate the current simulation one or more times in each dimension. For example, replication factors of 2,2,2 will create a simulation with 8x as many atoms by doubling the simulation domain in each dimension. A replication factor of 1 in a dimension leaves the simulation domain unchanged. When the new simulation box is created it is also partitioned into a regular 3d grid of rectangular bricks, one per processor, based on the number of processors being used and the settings of the *processors* command. The partitioning can later be changed by the *balance* or *fix balance* commands.

All properties of the atoms are replicated, including their velocities, which may or may not be desirable. New atom IDs are assigned to new atoms, as are molecule IDs. Bonds and other topology interactions are created between pairs of new atoms as well as between old and new atoms. This is done by using the image flag for each atom to “unwrap” it out of the periodic box before replicating it.

This means that any molecular bond you specify in the original data file that crosses a periodic boundary should be between two atoms with image flags that differ by 1. This will allow the bond to be unwrapped appropriately.

The optional keyword *bbox* uses a bounding box to only check atoms in replicas that overlap with a processor's sub-domain when assigning atoms to processors. It typically results in a substantial speedup when using the replicate

command on a large number of processors. It does require temporary use of more memory, specifically that each processor can store all atoms in the entire system before it is replicated.

### 16.111.4 Restrictions

A 2d simulation cannot be replicated in the z dimension.

If a simulation is non-periodic in a dimension, care should be used when replicating it in that dimension, as it may put atoms nearly on top of each other.

---

**Note:** You cannot use the replicate command on a system which has a molecule that spans the box and is bonded to itself across a periodic boundary, so that the molecule is effectively a loop. A simple example would be a linear polymer chain that spans the simulation box and bonds back to itself across the periodic boundary. More realistic examples would be a CNT (meant to be an infinitely long CNT) or a graphene sheet or a bulk periodic crystal where there are explicit bonds specified between near neighbors. (Note that this only applies to systems that have permanent bonds as specified in the data file. A CNT that is just atoms modeled with the *AIREBO potential* has no such permanent bonds, so it can be replicated.) The reason replication does not work with those systems is that the image flag settings described above cannot be made consistent. I.e. it is not possible to define images flags so that when every pair of bonded atoms is unwrapped (using the image flags), they will be close to each other. The only way the replicate command could work in this scenario is for it to break a bond, insert more atoms, and re-connect the loop for the larger simulation box. But it is not clever enough to do this. So you will have to construct a larger version of your molecule as a pre-processing step and input a new data file to LAMMPS.

---

If the current simulation was read in from a restart file (before a run is performed), there must not be any fix information stored in the file for individual atoms. Similarly, no fixes can be defined at the time the replicate command is used that require vectors of atom information to be stored. This is because the replicate command does not know how to replicate that information for new atoms it creates. To work around this restriction, restart files may be converted into data files and fixes may be undefined via the *unfix* command before and redefined after the replicate command.

### 16.111.5 Related commands

none

### 16.111.6 Default

none

## 16.112 rerun command

### 16.112.1 Syntax

```
rerun file1 file2 ... keyword args ...
```

- file1,file2,... = dump file(s) to read
- one or more keywords may be appended, keyword *dump* must appear and be last

keyword = *first* or *last* or *every* or *skip* or *start* or *stop* or *post* or *dump*

*first* args = Nfirst

Nfirst = dump timestep to start on

```
last args = Nlast
  Nlast = dumptimestep to stop on
every args = Nevery
  Nevery = read snapshots matching every this many timesteps
skip args = Nskip
  Nskip = read one out of every Nskip snapshots
start args = Nstart
  Nstart = timestep on which pseudo run will start
stop args = Nstop
  Nstop = timestep to which pseudo run will end
post value = yes or no
dump args = same as read_dump command starting with its field arguments
```

### 16.112.2 Examples

```
rerun dump.file dump x y z vx vy vz
rerun dump1.txt dump2.txt first 10000 every 1000 dump x y z
rerun dump.vels dump x y z vx vy vz box yes format molfile lammprj
rerun dump.dcd dump x y z box no format molfile dcd
rerun ../run7/dump.file.gz skip 2 dump x y z box yes
rerun dump.bp dump x y z box no format adios
rerun dump.bp dump x y z vx vy vz format adios timeout 10.0
```

### 16.112.3 Description

Perform a pseudo simulation run where atom information is read one snapshot at a time from a dump file(s), and energies and forces are computed on the snapshot to produce thermodynamic or other output.

This can be useful in the following kinds of scenarios, after an initial simulation produced the dump file:

- Compute the energy and forces of snapshots using a different potential.
- Calculate one or more diagnostic quantities on the snapshots that were not computed in the initial run. These can also be computed with settings not used in the initial run, e.g. computing an RDF via the *compute rdf* command with a longer cutoff than was used initially.
- Calculate the portion of per-atom forces resulting from a subset of the potential. E.g. compute only Coulombic forces. This can be done by only defining only a Coulombic pair style in the rerun script. Doing this in the original script would result in different (bad) dynamics.

Conceptually, using the rerun command is like running an input script that has a loop in it (see the *next* and *jump* commands). Each iteration of the loop reads one snapshot from the dump file via the *read\_dump* command, sets the timestep to the appropriate value, and then invokes a *run* command for zero timesteps to simply compute energy and forces, and any other *thermodynamic output* or diagnostic info you have defined. This computation also invokes any fixes you have defined that apply constraints to the system, such as *fix shake* or *fix indent*.

Note that a simulation box must already be defined before using the rerun command. This can be done by the *create\_box*, *read\_data*, or *read\_restart* commands.

Also note that reading per-atom information from dump snapshots is limited to the atom coordinates, velocities and image flags as explained in the *read\_dump* command. Other atom properties, which may be necessary to compute energies and forces, such as atom charge, or bond topology information for a molecular system, are not read from (or even contained in) dump files. Thus this auxiliary information should be defined in the usual way, e.g. in a data file read in by a *read\_data* command, before using the rerun command.

Also note that the frequency of thermodynamic or dump output from the rerun simulation will depend on settings made in the rerun script, the same as for output from any LAMMPS simulation. See further info below as to what that means if the timesteps for snapshots read from dump files do not match the specified output frequency.

---

If more than one dump file is specified, the dump files are read one after the other in the order specified. It is assumed that snapshot timesteps will be in ascending order. If a snapshot is encountered that is not in ascending order, it will skip the snapshot until it reads one that is. This allows skipping of a duplicate snapshot (same timestep), e.g. that appeared at the end of one file and beginning of the next. However if you specify a series of dump files in an incorrect order (with respect to the timesteps they contain), you may skip large numbers of snapshots.

Note that the dump files specified as part of the *dump* keyword can be parallel files, i.e. written as multiple files either per processor and/or per snapshot. If that is the case they will also be read in parallel which can make the rerun command operate dramatically faster for large systems. See the page for the *read\_dump* and *dump* commands which describe how to read and write parallel dump files.

The *first*, *last*, *every*, *skip* keywords determine which snapshots are read from the dump file(s). Snapshots are skipped until they have a timestep  $\geq N_{first}$ . When a snapshot with a timestep  $> N_{last}$  is encountered, the rerun command finishes. Note that the defaults for *first* and *last* are to read all snapshots. If the *every* keyword is set to a value  $> 0$ , then only snapshots with timesteps that are a multiple of *Nevery* are read (the first snapshot is always read). If *Nevery* = 0, then this criterion is ignored, i.e. every snapshot is read that meets the other criteria. If the *skip* keyword is used, then after the first snapshot is read, every Nth snapshot is read, where  $N = N_{skip}$ . E.g. if *Nskip* = 3, then only 1 out of every 3 snapshots is read, assuming the snapshot timestep is also consistent with the other criteria.

---

**Note:** Not all dump formats contain the timestep and not all dump readers support reading it. In that case individual snapshots are assigned consecutive timestep numbers starting at 1.

---

The *start* and *stop* keywords do not affect which snapshots are read from the dump file(s). Rather, they have the same meaning that they do for the *run* command. They only need to be defined if (a) you are using a *fix* command that changes some value over time, and (b) you want the reference point for elapsed time (from start to stop) to be different than the *first* and *last* settings. See the page for individual fixes to see which ones can be used with the *start/stop* keywords. Note that if you define neither of the *start/stop* or *first/last* keywords, then LAMMPS treats the pseudo run as going from 0 to a huge value (effectively infinity). This means that any quantity that a fix scales as a fraction of elapsed time in the run, will essentially remain at its initial value. Also note that an error will occur if you read a snapshot from the dump file with a timestep value larger than the *stop* setting you have specified.

The *post* keyword can be used to minimize the output to the screen that happens after a *rerun* command, similar to the *post* keyword of the *run* command. It is set to *no* by default.

The *dump* keyword is required and must be the last keyword specified. Its arguments are passed internally to the *read\_dump* command. The first argument following the *dump* keyword should be the *field1* argument of the *read\_dump* command. See the *read\_dump* page for details on the various options it allows for extracting information from the dump file snapshots, and for using that information to alter the LAMMPS simulation.

---

In general, a LAMMPS input script that uses a rerun command can include and perform all the usual operations of an input script that uses the *run* command. There are a few exceptions and points to consider, as discussed here.

Fixes that perform time integration, such as *fix nve* or *fix npt* are not invoked, since no time integration is performed. Fixes that perturb or constrain the forces on atoms will be invoked, just as they would during a normal run. Examples are *fix indent* and *fix langevin*. So you should think carefully as to whether that makes sense for the manner in which you are reprocessing the dump snapshots.

If you only want the rerun script to perform an analysis that does not involve pair interactions, such as use compute msd to calculated displacements over time, you do not need to define a *pair style*, which may also mean neighbor lists

will not need to be calculated which saves time. The *comm\_modify cutoff* command can also be used to insure ghost atoms are acquired from far enough away for operations like bond and angle evaluations, if no pair style is being used.

Every time a snapshot is read, the timestep for the simulation is reset, as if the *reset\_timestep* command were used. This command has some restrictions as to what fixes can be defined. See its page for details. For example, the *fix deposit* and *fix dt/reset* fixes are in this category. They also make no sense to use with a rerun command.

If time-averaging fixes like *fix ave/time* are used, they are invoked on timesteps that are a function of their *Nevery*, *Nrepeat*, and *Nfreq* settings. As an example, see the *fix ave/time* page for details. You must insure those settings are consistent with the snapshot timestamps that are read from the dump file(s). If an averaging fix is not invoked on a timestep it expects to be, LAMMPS will flag an error.

The various forms of LAMMPS output, as defined by the *thermo\_style*, *thermo*, *dump*, and *restart* commands occur with specified frequency, e.g. every N steps. If the timestep for a dump snapshot is not a multiple of N, then it will be read and processed, but no output will be produced. If you want output for every dump snapshot, you can simply use N=1 for an output frequency, e.g. for thermodynamic output or new dump file output.

---

## 16.112.4 Restrictions

The *rerun* command is subject to all restrictions of the *read\_dump* command.

## 16.112.5 Related commands

*read\_dump*

## 16.112.6 Default

The option defaults are first = 0, last = a huge value (effectively infinity), start = same as first, stop = same as last, every = 0, skip = 1, post = no;

## 16.113 reset\_atom\_ids command

### 16.113.1 Syntax

```
reset_atom_ids keyword values ...  
  
* zero or more keyword/value pairs may be appended  
* keyword = *sort*
```

sort value = yes or no



## 16.113.2 Examples

```
reset_atom_ids
reset_atom_ids sort yes
```

## 16.113.3 Description

Reset atom IDs for the system, including all the global IDs stored for bond, angle, dihedral, improper topology data. This will create a set of IDs that are numbered contiguously from 1 to N for a N atoms system.

This can be useful to do after performing a “delete\_atoms” command for a molecular system. The delete\_atoms compress yes option will not perform this operation due to the existence of bond topology. It can also be useful to do after any simulation which has lost atoms, e.g. due to atoms moving outside a simulation box with fixed boundaries (see the “boundary command”), or due to evaporation (see the “fix evaporate” command).

If the *sort* keyword is used with a setting of *yes*, then the assignment of new atom IDs will be the same no matter how many processors LAMMPS is running on. This is done by first doing a spatial sort of all the atoms into bins and sorting them within each bin. Because the set of bins is independent of the number of processors, this enables a consistent assignment of new IDs to each atom.

This can be useful to do after using the “create\_atoms” command and/or “replicate” command. In general those commands do not guarantee assignment of the same atom ID to the same physical atom when LAMMPS is run on different numbers of processors. Enforcing consistent IDs can be useful for debugging or comparing output from two different runs.

Note that the spatial sort requires communication of atom IDs and coordinates between processors in an all-to-all manner. This is done efficiently in LAMMPS, but it is more expensive than how atom IDs are reset without sorting.

Note that whether sorting or not, the resetting of IDs is not a compression, where gaps in atom IDs are removed by decrementing atom IDs that are larger. Instead the IDs for all atoms are erased, and new IDs are assigned so that the atoms owned by an individual processor have consecutive IDs, as the *create\_atoms* command explains.

---

**Note:** If this command is used before a *pair style* is defined, an error about bond topology atom IDs not being found may result. This is because the cutoff distance for ghost atom communication was not sufficient to find atoms in bonds, angles, etc that are owned by other processors. The *comm\_modify cutoff* command can be used to correct this issue. Or you can define a pair style before using this command. If you do the former, you should unset the *comm\_modify cutoff* after using *reset\_atom\_ids* so that subsequent communication is not inefficient.

---

## 16.113.4 Restrictions

none

## 16.113.5 Related commands

*delete\_atoms*

### 16.113.6 Default

By default, *sort* is no.

## 16.114 reset\_mol\_ids command

### 16.114.1 Syntax

```
reset_mol_ids group-ID keyword value ...
```

- group-ID = ID of group of atoms whose molecule IDs will be reset
- zero or more keyword/value pairs may be appended
- keyword = *compress* or *offset* or *single*
  - compress* value = *yes* or *no*
  - offset* value = *Noffset* >= -1
  - single* value = *yes* or *no* to treat single atoms (no bonds) as molecules

### 16.114.2 Examples

```
reset_mol_ids all
reset_mol_ids all offset 10 single yes
reset_mol_ids solvent compress yes offset 100
reset_mol_ids solvent compress no
```

### 16.114.3 Description

Reset molecule IDs for a group of atoms based on current bond connectivity. This will typically create a new set of molecule IDs for atoms in the group. Only molecule IDs for atoms in the specified group are reset; molecule IDs for atoms not in the group are not changed.

For purposes of this operation, molecules are identified by the current bond connectivity in the system, which may or may not be consistent with the current molecule IDs. A molecule in this context is a set of atoms connected to each other with explicit bonds. The specific algorithm used is the one of *compute fragment/atom*. Once the molecules are identified and a new molecule ID computed for each, this command will update the current molecule ID for all atoms in the group with the new molecule ID. Note that if the group excludes atoms within molecules, one (physical) molecule may become two or more (logical) molecules. For example if the group excludes atoms in the middle of a linear chain, then each end of the chain is considered an independent molecule and will be assigned a different molecule ID.

This can be a useful operation to perform after running reactive molecular dynamics run with *fix bond/react*, *fix bond/create*, or *fix bond/break*, all of which can change molecule topologies. It can also be useful after molecules have been deleted with the *delete\_atoms* command or after a simulation which has lost molecules, e.g. via the *fix evaporate* command.

The *compress* keyword determines how new molecule IDs are computed. If the setting is *yes* (the default) and there are N molecules in the group, the new molecule IDs will be a set of N contiguous values. See the *offset* keyword for details on selecting the range of these values. If the setting is *no*, the molecule ID of every atom in the molecule will be set to the smallest atom ID of any atom in the molecule.

The *single* keyword determines whether single atoms (not bonded to another atom) are treated as one-atom molecules or not, based on the *yes* or *no* setting. If the setting is *no* (the default), their molecule IDs are set to 0. This setting can

be important if the new molecule IDs will be used as input to other commands such as *compute chunk/atom molecule* or *fix rigid molecule*.

The *offset* keyword is only used if the *compress* setting is *yes*. Its default value is *Noffset = -1*. In that case, if the specified group is *all*, then the new compressed molecule IDs will range from 1 to N. If the specified group is not *all* and the largest molecule ID of atoms outside that group is M, then the new compressed molecule IDs will range from M+1 to M+N, to avoid collision with existing molecule IDs. If an *Noffset*  $\geq 0$  is specified, then the new compressed molecule IDs will range from *Noffset+1* to *Noffset+N*. If the group is not *all* there may be collisions with the molecule IDs of other atoms.

---

**Note:** The same as explained for the *compute fragment/atom* command, molecules are identified using the current bond topology. This will not account for bonds broken by the *bond\_style quartic* command because it does not perform a full update of the bond topology data structures within LAMMPS.

---

#### 16.114.4 Restrictions

none

#### 16.114.5 Related commands

*reset\_atom\_ids*, *fix bond/react*, *fix bond/create*, *fix bond/break*, *fix evaporate*, *delete\_atoms*, *compute fragment/atom*

#### 16.114.6 Default

The default keyword settings are *compress = yes*, *single = no*, and *offset = -1*.

### 16.115 reset\_timestep command

#### 16.115.1 Syntax

```
reset_timestep N keyword values ...
```

- N = timestep number
- zero or more keyword/value pairs may be appended
- keyword = *time*

*time* value = *atime*

*atime* = accumulated simulation time

## 16.115.2 Examples

```
reset_timestep 0
reset_timestep 4000000
reset_timestep 1000 time 100.0
```

## 16.115.3 Description

Set the timestep counter to the specified value. This command usually comes after the timestep has been set by reading a restart file via the *read\_restart* command, or a previous simulation run or minimization advanced the timestep.

The optional *time* keyword allows to also set the accumulated simulation time. This is usually the number of timesteps times the size of the timestep, but when using variable size timesteps with *fix dt/reset* it can differ.

The *read\_data* and *create\_box* commands set the timestep to 0; the *read\_restart* command sets the timestep to the value it had when the restart file was written. The same applies to the accumulated simulation time.

## 16.115.4 Restrictions

This command cannot be used when any fixes are defined that keep track of elapsed time to perform certain kinds of time-dependent operations. Examples are the *fix deposit* and *fix dt/reset* commands. The former adds atoms on specific timesteps. The latter keeps track of accumulated time.

Various fixes use the current timestep to calculate related quantities. If the timestep is reset, this may produce unexpected behavior, but LAMMPS allows the fixes to be defined even if the timestep is reset. For example, commands which thermostat the system, e.g. *fix nvt*, allow you to specify a target temperature which ramps from Tstart to Tstop which may persist over several runs. If you change the timestep, you may induce an instantaneous change in the target temperature.

Resetting the timestep clears flags for *computes* that may have calculated some quantity from a previous run. This means these quantity cannot be accessed by a variable in between runs until a new run is performed. See the *variable* command for more details.

## 16.115.5 Related commands

*rerun*, *timestep*, *fix dt/reset*

## 16.115.6 Default

none

## 16.116 restart command

### 16.116.1 Syntax

```
restart 0
restart N root keyword value ...
restart N file1 file2 keyword value ...
```

- N = write a restart file on timesteps which are multiples of N
- N can be a variable (see below)
- root = filename to which timestep # is appended
- file1,file2 = two full filenames, toggle between them when writing file
- zero or more keyword/value pairs may be appended
- keyword = *fileper* or *nfile*

*fileper* arg = Np

Np = write one file for every this many processors

*nfile* arg = Nf

Nf = write this many files, one from each of Nf processors

## 16.116.2 Examples

```
restart 0
restart 1000 poly.restart
restart 1000 poly.restart.mpio
restart 1000 restart.*.equil
restart 10000 poly.%1 poly.%2 nfile 10
restart v_mystep poly.restart
```

## 16.116.3 Description

Write out a binary restart file with the current state of the simulation on timesteps which are a multiple of N. A value of N = 0 means do not write out any restart files, which is the default. Restart files are written in one (or both) of two modes as a run proceeds. If one filename is specified, a series of filenames will be created which include the timestep in the filename. If two filenames are specified, only 2 restart files will be created, with those names. LAMMPS will toggle between the 2 names as it writes successive restart files.

Note that you can specify the restart command twice, once with a single filename and once with two filenames. This would allow you, for example, to write out archival restart files every 100000 steps using a single filename, and more frequent temporary restart files every 1000 steps, using two filenames. Using restart 0 will turn off both modes of output.

Similar to *dump* files, the restart filename(s) can contain two wild-card characters.

If a "\*" appears in the single filename, it is replaced with the current timestep value. This is only recognized when a single filename is used (not when toggling back and forth). Thus, the third example above creates restart files as follows: restart.1000.equil, restart.2000.equil, etc. If a single filename is used with no "\*", then the timestep value is appended. E.g. the second example above creates restart files as follows: poly.restart.1000, poly.restart.2000, etc.

If a "%" character appears in the restart filename(s), then one file is written for each processor and the "%" character is replaced with the processor ID from 0 to P-1. An additional file with the "%" replaced by "base" is also written, which contains global information. For example, the files written on step 1000 for filename restart.% would be restart.base.1000, restart.0.1000, restart.1.1000, ..., restart.P-1.1000. This creates smaller files and can be a fast mode of output and subsequent input on parallel machines that support parallel I/O. The optional *fileper* and *nfile* keywords discussed below can alter the number of files written.

The restart file can also be written in parallel as one large binary file via the MPI-IO library, which is part of the MPI standard for versions 2.0 and above. Using MPI-IO requires two steps. First, build LAMMPS with its MPIIO package installed, e.g.

```
make yes-mpio    # installs the MPIIO package
make mpi        # build LAMMPS for your platform
```

Second, use a restart filename which contains “.mpio”. Note that it does not have to end in “.mpio”, just contain those characters. Unlike MPI-IO dump files, a particular restart file must be both written and read using MPI-IO.

Restart files are written on timesteps that are a multiple of N but not on the first timestep of a run or minimization. You can use the `write_restart` command to write a restart file before a run begins. A restart file is not written on the last timestep of a run unless it is a multiple of N. A restart file is written on the last timestep of a minimization if  $N > 0$  and the minimization converges.

Instead of a numeric value, N can be specified as an *equal-style variable*, which should be specified as `v_name`, where name is the variable name. In this case, the variable is evaluated at the beginning of a run to determine the next timestep at which a restart file will be written out. On that timestep, the variable will be evaluated again to determine the next timestep, etc. Thus the variable should return timestep values. See the `stagger()` and `logfreq()` and `stride()` math functions for *equal-style variables*, as examples of useful functions to use in this context. Other similar math functions could easily be added as options for *equal-style variables*.

For example, the following commands will write restart files every step from 1100 to 1200, and could be useful for debugging a simulation where something goes wrong at step 1163:

```
variable        s equal stride(1100,1200,1)
restart         v_s tmp.restart
```

---

See the `read_restart` command for information about what is stored in a restart file.

Restart files can be read by a `read_restart` command to restart a simulation from a particular state. Because the file is binary (to enable exact restarts), it may not be readable on another machine. In this case, you can use the `-r command-line switch` to convert a restart file to a data file.

---

**Note:** Although the purpose of restart files is to enable restarting a simulation from where it left off, not all information about a simulation is stored in the file. For example, the list of fixes that were specified during the initial run is not stored, which means the new input script must specify any fixes you want to use. Even when restart information is stored in the file, as it is for some fixes, commands may need to be re-specified in the new input script, in order to re-use that information. See the `read_restart` command for information about what is stored in a restart file.

---

The optional `nfile` or `fileper` keywords can be used in conjunction with the “%” wildcard character in the specified restart file name(s). As explained above, the “%” character causes the restart file to be written in pieces, one piece for each of P processors. By default  $P =$  the number of processors the simulation is running on. The `nfile` or `fileper` keyword can be used to set P to a smaller value, which can be more efficient when running on a large number of processors.

The `nfile` keyword sets P to the specified Nf value. For example, if  $N_f = 4$ , and the simulation is running on 100 processors, 4 files will be written, by processors 0,25,50,75. Each will collect information from itself and the next 24 processors and write it to a restart file.

For the `fileper` keyword, the specified value of Np means write one file for every Np processors. For example, if  $N_p = 4$ , every fourth processor (0,4,8,12,etc) will collect information from itself and the next 3 processors and write it to a restart file.

---

## 16.116.4 Restrictions

To write and read restart files in parallel with MPI-IO, the MPIIO package must be installed.

## 16.116.5 Related commands

*write\_restart, read\_restart*

## 16.116.6 Default

```
restart 0
```

## 16.117 run command

### 16.117.1 Syntax

```
run N keyword values ...
```

- N = # of timesteps
- zero or more keyword/value pairs may be appended
- keyword = *upto* or *start* or *stop* or *pre* or *post* or *every*

*upto* value = none

*start* value = N1

N1 = timestep at which 1st run started

*stop* value = N2

N2 = timestep at which last run will end

*pre* value = *no* or *yes*

*post* value = *no* or *yes*

*every* values = M c1 c2 ...

M = break the run into M-timestep segments and invoke one or more commands,  
 ↳ between each segment

c1,c2,...,cN = one or more LAMMPS commands, each enclosed in quotes

c1 = NULL means no command will be invoked

### 16.117.2 Examples

```
run 10000
run 1000000 upto
run 100 start 0 stop 1000
run 1000 pre no post yes
run 100000 start 0 stop 1000000 every 1000 "print 'Protein Rg = $r'"
run 100000 every 1000 NULL
```

### 16.117.3 Description

Run or continue dynamics for a specified number of timesteps.

When the *run style* is *respa*, N refers to outer loop (largest) timesteps.

A value of N = 0 is acceptable; only the thermodynamics of the system are computed and printed without taking a timestep.

The *upto* keyword means to perform a run starting at the current timestep up to the specified timestep. E.g. if the current timestep is 10,000 and “run 100000 upto” is used, then an additional 90,000 timesteps will be run. This can be useful for very long runs on a machine that allocates chunks of time and terminate your job when time is exceeded. If you need to restart your script multiple times (reading in the last restart file), you can keep restarting your script with the same run command until the simulation finally completes.

The *start* or *stop* keywords can be used if multiple runs are being performed and you want a *fix* command that changes some value over time (e.g. temperature) to make the change across the entire set of runs and not just a single run. See the page for individual fixes to see which ones can be used with the *start/stop* keywords.

For example, consider this fix followed by 10 run commands:

```
fix          1 all nvt 200.0 300.0 1.0
run          1000 start 0 stop 10000
run          1000 start 0 stop 10000
...
run          1000 start 0 stop 10000
```

The NVT fix ramps the target temperature from 200.0 to 300.0 during a run. If the run commands did not have the start/stop keywords (just “run 1000”), then the temperature would ramp from 200.0 to 300.0 during the 1000 steps of each run. With the start/stop keywords, the ramping takes place over the 10000 steps of all runs together.

The *pre* and *post* keywords can be used to streamline the setup, clean-up, and associated output to the screen that happens before and after a run. This can be useful if you wish to do many short runs in succession (e.g. LAMMPS is being called as a library which is doing other computations between successive short LAMMPS runs).

By default (*pre* and *post* = yes), LAMMPS creates neighbor lists, computes forces, and imposes fix constraints before every run. And after every run it gathers and prints timings statistics. If a run is just a continuation of a previous run (i.e. no settings are changed), the initial computation is not necessary; the old neighbor list is still valid as are the forces. So if *pre* is specified as “no” then the initial setup is skipped, except for printing thermodynamic info. Note that if *pre* is set to “no” for the very first run LAMMPS performs, then it is overridden, since the initial setup computations must be done.

---

**Note:** If your input script changes the system between 2 runs, then the initial setup must be performed to insure the change is recognized by all parts of the code that are affected. Examples are adding a *fix* or *dump* or *compute*, changing a *neighbor* list parameter, or writing restart file which can migrate atoms between processors. LAMMPS has no easy way to check if this has happened, but it is an error to use the *pre no* option in this case.

---

If *post* is specified as “no”, the full timing summary is skipped; only a one-line summary timing is printed.

The *every* keyword provides a means of breaking a LAMMPS run into a series of shorter runs. Optionally, one or more LAMMPS commands (c1, c2, ..., cN) will be executed in between the short runs. If used, the *every* keyword must be the last keyword, since it has a variable number of arguments. Each of the trailing arguments is a single LAMMPS command, and each command should be enclosed in quotes, so that the entire command will be treated as a single argument. This will also prevent any variables in the command from being evaluated until it is executed multiple times during the run. Note that if a command itself needs one of its arguments quoted (e.g. the *print* command), then you can use a combination of single and double quotes, as in the example above or below.



The *every* keyword is a means to avoid listing a long series of runs and interleaving commands in your input script. For example, a *print* command could be invoked or a *fix* could be redefined, e.g. to reset a thermostat temperature. Or this could be useful for invoking a command you have added to LAMMPS that wraps some other code (e.g. as a library) to perform a computation periodically during a long LAMMPS run. See the *Modify* doc page for info about how to add new commands to LAMMPS. See the *Howto couple* page for ideas about how to couple LAMMPS to other codes.

With the *every* option, N total steps are simulated, in shorter runs of M steps each. After each M-length run, the specified commands are invoked. If only a single command is specified as NULL, then no command is invoked. Thus these lines:

```
variable q equal x[100]
run 6000 every 2000 "print 'Coord = $q'"
```

are the equivalent of:

```
variable q equal x[100]
run 2000
print "Coord = $q"
run 2000
print "Coord = $q"
run 2000
print "Coord = $q"
```

which does 3 runs of 2000 steps and prints the x-coordinate of a particular atom between runs. Note that the variable “\$q” will be evaluated afresh each time the print command is executed.

Note that by using the line continuation character “&”, the run every command can be spread across many lines, though it is still a single command:

```
run 100000 every 1000 &
  "print 'Minimum value = $a'" &
  "print 'Maximum value = $b'" &
  "print 'Temp = $c'" &
  "print 'Press = $d'"
```

If the *pre* and *post* options are set to “no” when used with the *every* keyword, then the first run will do the full setup and the last run will print the full timing summary, but these operations will be skipped for intermediate runs.

---

**Note:** You might wish to specify a command that exits the run by jumping out of the loop, e.g.

---

```
variable t equal temp
run 10000 every 100 "if '$t < 300.0' then 'jump SELF afterrun'"
```

However, this will not work. The run command simply executes each command one at a time each time it pauses, then continues the run.

Instead, you should use the *fix halt* command, which has additional options for how to exit the run.

### 16.117.4 Restrictions

When not using the *upto* keyword, the number of specified timesteps *N* must fit in a signed 32-bit integer, so you are limited to slightly more than 2 billion steps ( $2^{31}$ ) in a single run. When using *upto*, *N* can be larger than a signed 32-bit integer, however the difference between *N* and the current timestep must still be no larger than  $2^{31}$  steps.

However, with or without the *upto* keyword, you can perform successive runs to run a simulation for any number of steps (ok, up to  $2^{63}$  total steps). I.e. the timestep counter within LAMMPS is a 64-bit signed integer.

### 16.117.5 Related commands

*minimize*, *run\_style*, *temper*, *fix halt*

### 16.117.6 Default

The option defaults are *start* = the current timestep, *stop* = current timestep + *N*, *pre* = yes, and *post* = yes.

## 16.118 run\_style command

### 16.118.1 Syntax

```
run_style style args
```

- *style* = *verlet* or *verlet/split* or *respa* or *respa/omp*

*verlet* args = none

*verlet/split* args = none

*respa* args = *N* *n1* *n2* ... keyword values ...

*N* = # of levels of rRESPA

*n1*, *n2*, ... = loop factors between rRESPA levels (*N*-1 values)

zero or more keyword/value pairings may be appended to the loop factors

keyword = *bond* or *angle* or *dihedral* or *improper* or

*pair* or *inner* or *middle* or *outer* or *hybrid* or *kpace*

*bond* value = *M*

*M* = which level (1-*N*) to compute bond forces in

*angle* value = *M*

*M* = which level (1-*N*) to compute angle forces in

*dihedral* value = *M*

*M* = which level (1-*N*) to compute dihedral forces in

*improper* value = *M*

*M* = which level (1-*N*) to compute improper forces in

*pair* value = *M*

*M* = which level (1-*N*) to compute pair forces in

*inner* values = *M* *cut1* *cut2*

*M* = which level (1-*N*) to compute pair inner forces in

*cut1* = inner cutoff between pair inner and  
pair middle or outer (distance units)

*cut2* = outer cutoff between pair inner and  
pair middle or outer (distance units)

*middle* values = *M* *cut1* *cut2*

*M* = which level (1-*N*) to compute pair middle forces in

```

cut1 = inner cutoff between pair middle and pair outer (distance units)
cut2 = outer cutoff between pair middle and pair outer (distance units)
outer value = M
M = which level (1-N) to compute pair outer forces in
hybrid values = M1 M2 ... (as many values as there are hybrid sub-styles
M1 = which level (1-N) to compute the first pair_style hybrid sub-style in
M2 = which level (1-N) to compute the second pair_style hybrid sub-style in
M3, etc
kspace value = M
M = which level (1-N) to compute kspace forces in

```

## 16.118.2 Examples

```

run_style verlet
run_style respa 4 2 2 2 bond 1 dihedral 2 pair 3 kspace 4
run_style respa 4 2 2 2 bond 1 dihedral 2 inner 3 5.0 6.0 outer 4 kspace 4
run_style respa 3 4 2 bond 1 hybrid 2 2 1 kspace 3

```

## 16.118.3 Description

Choose the style of time integrator used for molecular dynamics simulations performed by LAMMPS.

The *verlet* style is the velocity form of the Stoermer-Verlet time integration algorithm (velocity-Verlet)

The *verlet/split* style is also a velocity-Verlet integrator, but it splits the force calculation within each timestep over 2 partitions of processors. See the *-partition command-line switch* for info on how to run LAMMPS with multiple partitions.

Specifically, this style performs all computation except the *kspace\_style* portion of the force field on the first partition. This include the *pair style*, *bond style*, *neighbor list building*, *fixes* including time integration, and output. The *kspace\_style* portion of the calculation is performed on the second partition.

This is most useful for the PPPM *kspace\_style* when its performance on a large number of processors degrades due to the cost of communication in its 3d FFTs. In this scenario, splitting your P total processors into 2 subsets of processors, P1 in the first partition and P2 in the second partition, can enable your simulation to run faster. This is because the long-range forces in PPPM can be calculated at the same time as pairwise and bonded forces are being calculated, and the FFTs can actually speed up when running on fewer processors.

To use this style, you must define 2 partitions where P1 is a multiple of P2. Typically having P1 be 3x larger than P2 is a good choice. The 3d processor layouts in each partition must overlay in the following sense. If P1 is a Px1 by Py1 by Pz1 grid, and P2 = Px2 by Py2 by Pz2, then Px1 must be an integer multiple of Px2, and similarly for Py1 a multiple of Py2, and Pz1 a multiple of Pz2.

Typically the best way to do this is to let the first partition choose its own optimal layout, then require the second partition's layout to match the integer multiple constraint. See the *processors* command with its *part* keyword for a way to control this, e.g.

```
processors * * * part 1 2 multiple
```

You can also use the *partition* command to explicitly specify the processor layout on each partition. E.g. for 2 partitions of 60 and 15 processors each:

```
partition yes 1 processors 3 4 5
partition yes 2 processors 3 1 5
```

When you run in 2-partition mode with the *verlet/split* style, the thermodynamic data for the entire simulation will be output to the log and screen file of the first partition, which are `log.lammps.0` and `screen.0` by default; see the *-plog and -pscreen command-line switches* to change this. The log and screen file for the second partition will not contain thermodynamic output beyond the first timestep of the run.

See the *Accelerator packages* page for performance details of the speed-up offered by the *verlet/split* style. One important performance consideration is the assignment of logical processors in the 2 partitions to the physical cores of a parallel machine. The *processors* command has options to support this, and strategies are discussed in *Section 5* of the manual.

---

The *respa* style implements the rRESPA multi-timescale integrator (*Tuckerman*) with N hierarchical levels, where level 1 is the innermost loop (shortest timestep) and level N is the outermost loop (largest timestep). The loop factor arguments specify what the looping factor is between levels. N1 specifies the number of iterations of level 1 for a single iteration of level 2, N2 is the iterations of level 2 per iteration of level 3, etc. N-1 looping parameters must be specified.

Thus with a 4-level respa setting of “2 2 2” for the 3 loop factors, you could choose to have bond interactions computed 8x per large timestep, angle interactions computed 4x, pair interactions computed 2x, and long-range interactions once per large timestep.

The *timestep* command sets the large timestep for the outermost rRESPA level. Thus if the 3 loop factors are “2 2 2” for 4-level rRESPA, and the outer timestep is set to 4.0 fs, then the inner timestep would be 8x smaller or 0.5 fs. All other LAMMPS commands that specify number of timesteps (e.g. *thermo* for thermo output every N steps, *neigh\_modify delay/every* parameters, *dump* every N steps, etc) refer to the outermost timesteps.

The rRESPA keywords enable you to specify at what level of the hierarchy various forces will be computed. If not specified, the defaults are that bond forces are computed at level 1 (innermost loop), angle forces are computed where bond forces are, dihedral forces are computed where angle forces are, improper forces are computed where dihedral forces are, pair forces are computed at the outermost level, and kspace forces are computed where pair forces are. The inner, middle, outer forces have no defaults.

For fixes that support it, the rRESPA level at which a given fix is active, can be selected through the *fix\_modify* command.

The *inner* and *middle* keywords take additional arguments for cutoffs that are used by the pairwise force computations. If the 2 cutoffs for *inner* are 5.0 and 6.0, this means that all pairs up to 6.0 apart are computed by the inner force. Those between 5.0 and 6.0 have their force go ramped to 0.0 so the overlap with the next regime (middle or outer) is smooth. The next regime (middle or outer) will compute forces for all pairs from 5.0 outward, with those from 5.0 to 6.0 having their value ramped in an inverse manner.

Note that you can use *inner* and *outer* without using *middle* to split the pairwise computations into two portions instead of three. Unless you are using a very long pairwise cutoff, a 2-way split is often faster than a 3-way split, since it avoids too much duplicate computation of pairwise interactions near the intermediate cutoffs.

Also note that only a few pair potentials support the use of the *inner* and *middle* and *outer* keywords. If not, only the *pair* keyword can be used with that pair style, meaning all pairwise forces are computed at the same rRESPA level. See the doc pages for individual pair styles for details.

Another option for using pair potentials with rRESPA is with the *hybrid* keyword, which requires the use of the *pair\_style hybrid or hybrid/overlay* command. In this scenario, different sub-styles of the hybrid pair style are evaluated at different rRESPA levels. This can be useful, for example, to set different timesteps for hybrid coarse-grained/all-atom models. The *hybrid* keyword requires as many level assignments as there are hybrid sub-styles, which assigns each sub-style to a rRESPA level, following their order of definition in the *pair\_style* command. Since the *hybrid* keyword operates on pair style computations, it is mutually exclusive with either the *pair* or the *inner/middle/outer* keywords.

When using rRESPA (or for any MD simulation) care must be taken to choose a timestep size(s) that insures the Hamiltonian for the chosen ensemble is conserved. For the constant NVE ensemble, total energy must be conserved.

Unfortunately, it is difficult to know *a priori* how well energy will be conserved, and a fairly long test simulation (~10 ps) is usually necessary in order to verify that no long-term drift in energy occurs with the trial set of parameters.

With that caveat, a few rules-of-thumb may be useful in selecting *respa* settings. The following applies mostly to biomolecular simulations using the CHARMM or a similar all-atom force field, but the concepts are adaptable to other problems. Without SHAKE, bonds involving hydrogen atoms exhibit high-frequency vibrations and require a timestep on the order of 0.5 fs in order to conserve energy. The relatively inexpensive force computations for the bonds, angles, impropers, and dihedrals can be computed on this innermost 0.5 fs step. The outermost timestep cannot be greater than 4.0 fs without risking energy drift. Smooth switching of forces between the levels of the rRESPA hierarchy is also necessary to avoid drift, and a 1-2 angstrom “healing distance” (the distance between the outer and inner cutoffs) works reasonably well. We thus recommend the following settings for use of the *respa* style without SHAKE in biomolecular simulations:

```
timestep 4.0
run_style respa 4 2 2 2 inner 2 4.5 6.0 middle 3 8.0 10.0 outer 4
```

With these settings, users can expect good energy conservation and roughly a 2.5 fold speedup over the *verlet* style with a 0.5 fs timestep.

If SHAKE is used with the *respa* style, time reversibility is lost, but substantially longer time steps can be achieved. For biomolecular simulations using the CHARMM or similar all-atom force field, bonds involving hydrogen atoms exhibit high frequency vibrations and require a time step on the order of 0.5 fs in order to conserve energy. These high frequency modes also limit the outer time step sizes since the modes are coupled. It is therefore desirable to use SHAKE with *respa* in order to freeze out these high frequency motions and increase the size of the time steps in the *respa* hierarchy. The following settings can be used for biomolecular simulations with SHAKE and rRESPA:

```
fix 2 all shake 0.000001 500 0 m 1.0 a 1
timestep 4.0
run_style respa 2 2 inner 1 4.0 5.0 outer 2
```

With these settings, users can expect good energy conservation and roughly a 1.5 fold speedup over the *verlet* style with SHAKE and a 2.0 fs timestep.

For non-biomolecular simulations, the *respa* style can be advantageous if there is a clear separation of time scales - fast and slow modes in the simulation. For example, a system of slowly-moving charged polymer chains could be setup as follows:

```
timestep 4.0
run_style respa 2 8
```

This is two-level rRESPA with an 8x difference between the short and long timesteps. The bonds, angles, dihedrals will be computed every 0.5 fs (assuming real units), while the pair and kspace interactions will be computed once every 4 fs. These are the default settings for each kind of interaction, so no additional keywords are necessary.

Even a LJ system can benefit from rRESPA if the interactions are divided by the inner, middle and outer keywords. A 2-fold or more speedup can be obtained while maintaining good energy conservation. In real units, for a pure LJ fluid at liquid density, with a sigma of 3.0 angstroms, and epsilon of 0.1 Kcal/mol, the following settings seem to work well:

```
timestep 36.0
run_style respa 3 3 4 inner 1 3.0 4.0 middle 2 6.0 7.0 outer 3
```

The *respa/omp* style is a variant of *respa* adapted for use with pair, bond, angle, dihedral, improper, or kspace styles with an *omp* suffix. It is functionally equivalent to *respa* but performs additional operations required for managing *omp* styles. For more on *omp* styles see the *Speed omp* doc page. Accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

You can specify *respa/omp* explicitly in your input script, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 16.118.4 Restrictions

The *verlet/split* style can only be used if LAMMPS was built with the REPLICa package. Correspondingly the *respa/omp* style is available only if the OPENMP package was included. See the *Build package* page for more info.

Whenever using rRESPA, the user should experiment with trade-offs in speed and accuracy for their system, and verify that they are conserving energy to adequate precision.

## 16.118.5 Related commands

*timestep, run*

## 16.118.6 Default

```
run_style verlet
```

For `run_style respa`, the default assignment of interactions to rRESPA levels is as follows:

- bond forces = level 1 (innermost loop)
- angle forces = same level as bond forces
- dihedral forces = same level as angle forces
- improper forces = same level as dihedral forces
- pair forces = level N (outermost level)
- kspace forces = same level as pair forces
- inner, middle, outer forces = no default

---

(Tuckerman) Tuckerman, Berne and Martyna, J Chem Phys, 97, p 1990 (1992).

## 16.119 set command

### 16.119.1 Syntax

```
set style ID keyword values ...
```

- style = *atom* or *type* or *mol* or *group* or *region*
- ID = atom ID range or type range or mol ID range or group ID or region ID
- one or more keyword/value pairs may be appended

- keyword = *type* or *type/fraction* or *type/ratio* or *type/subset* or *mol* or *x* or *y* or *z* or *vx* or *vy* or *vz* or *charge* or *dipole* or *dipole/random* or *quat* or *spin/atom* or *spin/atom/random* or *spin/electron* or *radius/electron* or *quat/random* or *diameter* or *shape* or *length* or *tri* or *theta* or *theta/random* or *angmom* or *omega* or *mass* or *density* or *density/disc* or *volume* or *image* or *bond* or *angle* or *dihedral* or *improper* or *sph/e* or *sph/cv* or *sph/rho* or *smd/contact/radius* or *smd/mass/density* or *dpd/theta* or *edpd/temp* or *edpd/cv* or *cc* or *epsilon* or *i\_name* or *d\_name* or *i2\_name* or *d2\_name*

*type* value = atom type  
value can be an atom-style variable (see below)

*type/fraction* values = type fraction seed  
type = new atom type  
fraction = approximate fraction of selected atoms to set to new atom type  
seed = random # seed (positive integer)

*type/ratio* values = type fraction seed  
type = new atom type  
fraction = exact fraction of selected atoms to set to new atom type  
seed = random # seed (positive integer)

*type/subset* values = type Nsubset seed  
type = new atom type  
Nsubset = exact number of selected atoms to set to new atom type  
seed = random # seed (positive integer)

*mol* value = molecule ID  
value can be an atom-style variable (see below)

*x,y,z* value = atom coordinate (distance units)  
value can be an atom-style variable (see below)

*vx,vy,vz* value = atom velocity (velocity units)  
value can be an atom-style variable (see below)

*charge* value = atomic charge (charge units)  
value can be an atom-style variable (see below)

*dipole* values = x y z  
x,y,z = orientation of dipole moment vector  
any of x,y,z can be an atom-style variable (see below)

*dipole/random* value = seed Dlen  
seed = random # seed (positive integer) for dipole moment orientations  
Dlen = magnitude of dipole moment (dipole units)

*spin/atom* values = g x y z  
g = magnitude of magnetic spin vector (in Bohr magneton's unit)  
x,y,z = orientation of magnetic spin vector  
any of x,y,z can be an atom-style variable (see below)

*spin/atom/random* value = seed Dlen  
seed = random # seed (positive integer) for magnetic spin orientations  
Dlen = magnitude of magnetic spin vector (in Bohr magneton's unit)

*radius/electron* values = eradius  
eradius = electron radius (or fixed-core radius) (distance units)

*spin/electron* value = espin  
espin = electron spin (+1/-1), 0 = nuclei, 2 = fixed-core, 3 = pseudo-cores (i.e., ↪ECP)

*quat* values = a b c theta  
a,b,c = unit vector to rotate particle around via right-hand rule  
theta = rotation angle (degrees)  
any of a,b,c,theta can be an atom-style variable (see below)

*quat/random* value = seed  
seed = random # seed (positive integer) for quaternion orientations

*diameter* value = diameter of spherical particle (distance units)

value can be an atom-style variable (see below)  
*shape* value = Sx Sy Sz  
Sx,Sy,Sz = 3 diameters of ellipsoid (distance units)  
*length* value = len  
len = length of line segment (distance units)  
len can be an atom-style variable (see below)  
*tri* value = side  
side = side length of equilateral triangle (distance units)  
side can be an atom-style variable (see below)  
*theta* value = angle (degrees)  
angle = orientation of line segment with respect to x-axis  
angle can be an atom-style variable (see below)  
*theta/random* value = seed  
seed = random # seed (positive integer) for line segment orientations  
*angmom* values = Lx Ly Lz  
Lx,Ly,Lz = components of angular momentum vector (distance-mass-velocity units)  
any of Lx,Ly,Lz can be an atom-style variable (see below)  
*omega* values = Wx Wy Wz  
Wx,Wy,Wz = components of angular velocity vector (radians/time units)  
any of wx,wy,wz can be an atom-style variable (see below)  
*mass* value = per-atom mass (mass units)  
value can be an atom-style variable (see below)  
*density* value = particle density for a sphere or ellipsoid (mass/distance<sup>3</sup> units),  
↳ or for a triangle (mass/distance<sup>2</sup> units) or line (mass/distance units) particle  
value can be an atom-style variable (see below)  
*density/disc* value = particle density for a 2d disc or ellipse (mass/distance<sup>2</sup>,  
↳ units)  
value can be an atom-style variable (see below)  
*volume* value = particle volume for Peridynamic particle (distance<sup>3</sup> units)  
value can be an atom-style variable (see below)  
*image* nx ny nz  
nx,ny,nz = which periodic image of the simulation box the atom is in  
any of nx,ny,nz can be an atom-style variable (see below)  
*bond* value = bond type for all bonds between selected atoms  
*angle* value = angle type for all angles between selected atoms  
*dihedral* value = dihedral type for all dihedrals between selected atoms  
*improper* value = improper type for all improper between selected atoms  
*sph/e* value = energy of SPH particles (need units)  
value can be an atom-style variable (see below)  
*sph/cv* value = heat capacity of SPH particles (need units)  
value can be an atom-style variable (see below)  
*sph/rho* value = density of SPH particles (need units)  
value can be an atom-style variable (see below)  
*smd/contact/radius* = radius for short range interactions, i.e. contact and friction  
value can be an atom-style variable (see below)  
*smd/mass/density* = set particle mass based on volume by providing a mass density  
value can be an atom-style variable (see below)  
*dpd/theta* value = internal temperature of DPD particles (temperature units)  
value can be an atom-style variable (see below)  
value can be NULL which sets internal temp of each particle to KE temp  
*edpd/temp* value = temperature of eDPD particles (temperature units)  
value can be an atom-style variable (see below)  
*edpd/cv* value = volumetric heat capacity of eDPD particles (energy/temperature/  
↳ volume units)



value can be an atom-style variable (see below)  
 cc values = index cc  
 index = index of a chemical species (1 to Nspecies)  
 cc = chemical concentration of tDPD particles for a species (mole/volume units)  
 epsilon value = dielectric constant of the medium where the atoms reside  
 i\_name value = custom integer vector with name  
 d\_name value = custom floating-point vector with name  
 i2\_name value = column of a custom integer array with name  
                   column specified as i2\_name[N] where N is 1 to Ncol  
 d2\_name value = column of a custom floating-point array with name  
                   column specified as d2\_name[N] where N is 1 to Ncol

## 16.119.2 Examples

```
set group solvent type 2
set group solvent type/fraction 2 0.5 12393
set group edge bond 4
set region half charge 0.5
set type 3 charge 0.5
set type 1*3 charge 0.5
set atom * charge v_atomfile
set atom 100*200 x 0.5 y 1.0
set atom 100 vx 0.0 vy 0.0 vz -1.0
set atom 1492 type 3
set atom * i_myVal 5
set atom * d2_Sxyz[1] 6.4
```

## 16.119.3 Description

Set one or more properties of one or more atoms. Since atom properties are initially assigned by the *read\_data*, *read\_restart* or *create\_atoms* commands, this command changes those assignments. This can be useful for overriding the default values assigned by the *create\_atoms* command (e.g. charge = 0.0). It can be useful for altering pairwise and molecular force interactions, since force-field coefficients are defined in terms of types. It can be used to change the labeling of atoms by atom type or molecule ID when they are output in *dump* files. It can also be useful for debugging purposes; i.e. positioning an atom at a precise location to compute subsequent forces or energy.

Note that the *style* and *ID* arguments determine which atoms have their properties reset. The remaining keywords specify which properties to reset and what the new values are. Some strings like *type* or *mol* can be used as a style and/or a keyword.

---

This section describes how to select which atoms to change the properties of, via the *style* and *ID* arguments.

The style *atom* selects all the atoms in a range of atom IDs. The style *type* selects all the atoms in a range of types. The style *mol* selects all the atoms in a range of molecule IDs.

In each of the range cases, the range can be specified as a single numeric value, or a wildcard asterisk can be used to specify a range of values. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. For example, for the style *type*, if N = the number of atom types, then an asterisk with no numeric values means all types from 1 to N. A leading asterisk means all types from 1 to n (inclusive). A trailing asterisk means all types from n to N (inclusive). A middle asterisk means all types from m to n (inclusive). For all the styles except *mol*, the lowest value for the wildcard is 1; for *mol* it is 0.

The style *group* selects all the atoms in the specified group. The style *region* selects all the atoms in the specified geometric region. See the *group* and *region* commands for details of how to specify a group or region.

This section describes the keyword options for which properties to change, for the selected atoms.

Note that except where explicitly prohibited below, all of the keywords allow an *atom-style* or *atomfile-style* variable to be used as the specified value(s). If the value is a variable, it should be specified as *v\_name*, where name is the variable name. In this case, the variable will be evaluated, and its resulting per-atom value used to determine the value assigned to each selected atom. Note that the per-atom value from the variable will be ignored for atoms that are not selected via the *style* and *ID* settings explained above. A simple way to use per-atom values from the variable to reset a property for all atoms is to use style *atom* with *ID* = “\*”; this selects all atom IDs.

Atom-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. They can also include per-atom values, such as atom coordinates. Thus it is easy to specify a time-dependent or spatially-dependent set of per-atom values. As explained on the *variable* doc page, atomfile-style variables can be used in place of atom-style variables, and thus as arguments to the set command. Atomfile-style variables read their per-atoms values from a file.

---

**Note:** Atom-style and atomfile-style variables return floating point per-atom values. If the values are assigned to an integer variable, such as the molecule ID, then the floating point value is truncated to its integer portion, e.g. a value of 2.6 would become 2.

---

Keyword *type* sets the atom type for all selected atoms. The specified value must be from 1 to *ntypes*, where *ntypes* was set by the *create\_box* command or the *atom types* field in the header of the data file read by the *read\_data* command.

Keyword *type/fraction* sets the atom type for a fraction of the selected atoms. The actual number of atoms changed is not guaranteed to be exactly the specified fraction ( $0 \leq \textit{fraction} \leq 1$ ), but should be statistically close. Random numbers are used in such a way that a particular atom is changed or not changed, regardless of how many processors are being used. This keyword does not allow use of an atom-style variable.

Keywords *type/ratio* and *type/subset* also set the atom type for a fraction of the selected atoms. The actual number of atoms changed will be exactly the requested number. For *type/ratio* the specified fraction ( $0 \leq \textit{fraction} \leq 1$ ) determines the number. For *type/subset*, the specified *Nsubset* is the number. An iterative algorithm is used which insures the correct number of atoms are selected, in a perfectly random fashion. Which atoms are selected will change with the number of processors used. These keywords do not allow use of an atom-style variable.

Keyword *mol* sets the molecule ID for all selected atoms. The *atom style* being used must support the use of molecule IDs.

Keywords *x*, *y*, *z*, and *charge* set the coordinates or charge of all selected atoms. For *charge*, the *atom style* being used must support the use of atomic charge. Keywords *vx*, *vy*, and *vz* set the velocities of all selected atoms.

Keyword *dipole* uses the specified x,y,z values as components of a vector to set as the orientation of the dipole moment vectors of the selected atoms. The magnitude of the dipole moment is set by the length of this orientation vector.

Keyword *dipole/random* randomizes the orientation of the dipole moment vectors for the selected atoms and sets the magnitude of each to the specified *Dlen* value. For 2d systems, the z component of the orientation is set to 0.0. Random numbers are used in such a way that the orientation of a particular atom is the same, regardless of how many processors are being used. This keyword does not allow use of an atom-style variable.

Changed in version 15Sep2022.

Keyword *spin/atom* uses the specified *g* value to set the magnitude of the magnetic spin vectors, and the x,y,z values as components of a vector to set as the orientation of the magnetic spin vectors of the selected atoms. This keyword was previously called *spin*.

Changed in version 15Sep2022.

Keyword *spin/atom/random* randomizes the orientation of the magnetic spin vectors for the selected atoms and sets the magnitude of each to the specified *Dlen* value. This keyword was previously called *spin/random*.

New in version 15Sep2022.

Keyword *radius/electron* uses the specified value to set the radius of electrons or fixed cores.

New in version 15Sep2022.

Keyword *spin/electron* sets the spin of an electron (+/- 1) or indicates nuclei (=0), fixed-cores (=2), or pseudo-cores (=3).

Keyword *quat* uses the specified values to create a quaternion (4-vector) that represents the orientation of the selected atoms. The particles must define a quaternion for their orientation (e.g. ellipsoids, triangles, body particles) as defined by the *atom\_style* command. Note that particles defined by *atom\_style ellipsoid* have 3 shape parameters. The 3 values must be non-zero for each particle set by this command. They are used to specify the aspect ratios of an ellipsoidal particle, which is oriented by default with its x-axis along the simulation box's x-axis, and similarly for y and z. If this body is rotated (via the right-hand rule) by an angle theta around a unit rotation vector (a,b,c), then the quaternion that represents its new orientation is given by  $(\cos(\theta/2), a*\sin(\theta/2), b*\sin(\theta/2), c*\sin(\theta/2))$ . The theta and a,b,c values are the arguments to the *quat* keyword. LAMMPS normalizes the quaternion in case (a,b,c) was not specified as a unit vector. For 2d systems, the a,b,c values are ignored, since a rotation vector of (0,0,1) is the only valid choice.

Keyword *quat/random* randomizes the orientation of the quaternion for the selected atoms. The particles must define a quaternion for their orientation (e.g. ellipsoids, triangles, body particles) as defined by the *atom\_style* command. Random numbers are used in such a way that the orientation of a particular atom is the same, regardless of how many processors are being used. For 2d systems, only orientations in the xy plane are generated. As with keyword *quat*, for ellipsoidal particles, the 3 shape values must be non-zero for each particle set by this command. This keyword does not allow use of an atom-style variable.

Keyword *diameter* sets the size of the selected atoms. The particles must be finite-size spheres as defined by the *atom\_style sphere* command. The diameter of a particle can be set to 0.0, which means they will be treated as point particles. Note that this command does not adjust the particle mass, even if it was defined with a density, e.g. via the *read\_data* command.

Keyword *shape* sets the size and shape of the selected atoms. The particles must be ellipsoids as defined by the *atom\_style ellipsoid* command. The *Sx*, *Sy*, *Sz* settings are the 3 diameters of the ellipsoid in each direction. All 3 can be set to the same value, which means the ellipsoid is effectively a sphere. They can also all be set to 0.0 which means the particle will be treated as a point particle. Note that this command does not adjust the particle mass, even if it was defined with a density, e.g. via the *read\_data* command.

Keyword *length* sets the length of selected atoms. The particles must be line segments as defined by the *atom\_style line* command. If the specified value is non-zero the line segment is (re)set to a length = the specified value, centered around the particle position, with an orientation along the x-axis. If the specified value is 0.0, the particle will become a point particle. Note that this command does not adjust the particle mass, even if it was defined with a density, e.g. via the *read\_data* command.

Keyword *tri* sets the size of selected atoms. The particles must be triangles as defined by the *atom\_style tri* command. If the specified value is non-zero the triangle is (re)set to be an equilateral triangle in the xy plane with side length = the specified value, with a centroid at the particle position, with its base parallel to the x axis, and the y-axis running from the center of the base to the top point of the triangle. If the specified value is 0.0, the particle will become a point particle. Note that this command does not adjust the particle mass, even if it was defined with a density, e.g. via the *read\_data* command.

Keyword *theta* sets the orientation of selected atoms. The particles must be line segments as defined by the *atom\_style line* command. The specified value is used to set the orientation angle of the line segments with respect to the x axis.

Keyword *theta/random* randomizes the orientation of theta for the selected atoms. The particles must be line segments as defined by the *atom\_style line* command. Random numbers are used in such a way that the orientation of a particular atom is the same, regardless of how many processors are being used. This keyword does not allow use of an atom-style variable.

Keyword *angmom* sets the angular momentum of selected atoms. The particles must be ellipsoids as defined by the *atom\_style ellipsoid* command or triangles as defined by the *atom\_style tri* command. The angular momentum vector of the particles is set to the 3 specified components.

Keyword *omega* sets the angular velocity of selected atoms. The particles must be spheres as defined by the *atom\_style sphere* command. The angular velocity vector of the particles is set to the 3 specified components.

Keyword *mass* sets the mass of all selected particles. The particles must have a per-atom mass attribute, as defined by the *atom\_style* command. See the “mass” command for how to set mass values on a per-type basis.

Keyword *density* or *density/disc* also sets the mass of all selected particles, but in a different way. The particles must have a per-atom mass attribute, as defined by the *atom\_style* command. If the atom has a radius attribute (see *atom\_style sphere*) and its radius is non-zero, its mass is set from the density and particle volume for 3d systems (the input density is assumed to be in mass/distance<sup>3</sup> units). For 2d, the default is for LAMMPS to model particles with a radius attribute as spheres. However, if the *density/disc* keyword is used, then they can be modeled as 2d discs (circles). Their mass is set from the density and particle area (the input density is assumed to be in mass/distance<sup>2</sup> units).

If the atom has a shape attribute (see *atom\_style ellipsoid*) and its 3 shape parameters are non-zero, then its mass is set from the density and particle volume (the input density is assumed to be in mass/distance<sup>3</sup> units). The *density/disc* keyword has no effect; it does not (yet) treat 3d ellipsoids as 2d ellipses.

If the atom has a length attribute (see *atom\_style line*) and its length is non-zero, then its mass is set from the density and line segment length (the input density is assumed to be in mass/distance units). If the atom has an area attribute (see *atom\_style tri*) and its area is non-zero, then its mass is set from the density and triangle area (the input density is assumed to be in mass/distance<sup>2</sup> units).

If none of these cases are valid, then the mass is set to the density value directly (the input density is assumed to be in mass units).

Keyword *volume* sets the volume of all selected particles. Currently, only the *atom\_style peri* command defines particles with a volume attribute. Note that this command does not adjust the particle mass.

Keyword *image* sets which image of the simulation box the atom is considered to be in. An image of 0 means it is inside the box as defined. A value of 2 means add 2 box lengths to get the true value. A value of -1 means subtract 1 box length to get the true value. LAMMPS updates these flags as atoms cross periodic boundaries during the simulation. The flags can be output with atom snapshots via the *dump* command. If a value of NULL is specified for any of nx,ny,nz, then the current image value for that dimension is unchanged. For non-periodic dimensions only a value of 0 can be specified. This command can be useful after a system has been equilibrated and atoms have diffused one or more box lengths in various directions. This command can then reset the image values for atoms so that they are effectively inside the simulation box, e.g if a diffusion coefficient is about to be measured via the *compute msd* command. Care should be taken not to reset the image flags of two atoms in a bond to the same value if the bond straddles a periodic boundary (rather they should be different by +/- 1). This will not affect the dynamics of a simulation, but may mess up analysis of the trajectories if a LAMMPS diagnostic or your own analysis relies on the image flags to unwrap a molecule which straddles the periodic box.

Keywords *bond*, *angle*, *dihedral*, and *improper*, set the bond type (angle type, etc) of all bonds (angles, etc) of selected atoms to the specified value from 1 to nbondtypes (nangletypes, etc). All atoms in a particular bond (angle, etc) must be selected atoms in order for the change to be made. The value of nbondtype (nangletypes, etc) was set by the *bond types* (*angle types*, etc) field in the header of the data file read by the *read\_data* command. These keywords do not allow use of an atom-style variable.

Keywords *sph/e*, *sph/cv*, and *sph/rho* set the energy, heat capacity, and density of smoothed particle hydrodynamics (SPH) particles. See [this PDF guide](#) to using SPH in LAMMPS.

Keyword *smd/mass/density* sets the mass of all selected particles, but it is only applicable to the Smooth Mach Dynamics package MACHDYN. It assumes that the particle volume has already been correctly set and calculates particle mass from the provided mass density value.

Keyword *smd/contact/radius* only applies to simulations with the Smooth Mach Dynamics package MACHDYN. It sets an interaction radius for computing short-range interactions, e.g. repulsive forces to prevent different individual physical

bodies from penetrating each other. Note that the SPH smoothing kernel diameter used for computing long range, nonlocal interactions, is set using the *diameter* keyword.

Keyword *dpd/theta* sets the internal temperature of a DPD particle as defined by the DPD-REACT package. If the specified value is a number it must be  $\geq 0.0$ . If the specified value is NULL, then the kinetic temperature  $T_{kin}$  of each particle is computed as  $\frac{3}{2} k T_{kin} = KE = \frac{1}{2} m v^2 = \frac{1}{2} m (v_x^2 + v_y^2 + v_z^2)$ . Each particle's internal temperature is set to  $T_{kin}$ . If the specified value is an atom-style variable, then the variable is evaluated for each particle. If a value  $\geq 0.0$ , the internal temperature is set to that value. If it is  $< 0.0$ , the computation of  $T_{kin}$  is performed and the internal temperature is set to that value.

Keywords *edpd/temp* and *edpd/cv* set the temperature and volumetric heat capacity of an eDPD particle as defined by the DPD-MESO package. Currently, only *atom\_style edpd* defines particles with these attributes. The values for the temperature and heat capacity must be positive.

Keyword *cc* sets the chemical concentration of a tDPD particle for a specified species as defined by the DPD-MESO package. Currently, only *atom\_style tdpd* defines particles with this attribute. An integer for "index" selects a chemical species (1 to  $N_{species}$ ) where  $N_{species}$  is set by the *atom\_style* command. The value for the chemical concentration must be  $\geq 0.0$ .

Keyword *epsilon* sets the dielectric constant of a particle, precisely of the medium where the particle resides as defined by the DIELECTRIC package. Currently, only *atom\_style dielectric* defines particles with this attribute. The value for the dielectric constant must be  $\geq 0.0$ . Note that the set command with this keyword will rescale the particle charge accordingly so that the real charge (e.g., as read from a data file) stays intact. To change the real charges, one needs to use the set command with the *charge* keyword. Care must be taken to ensure that the real and scaled charges, and dielectric constants are consistent.

Keywords *i\_name*, *d\_name*, *i2\_name*, *d2\_name* refer to custom per-atom integer and floating-point vectors or arrays that have been added via the *fix property/atom* command. When that command is used specific names are given to each attribute which are the "name" portion of these keywords. For arrays *i2\_name* and *d2\_name*, the column of the array must also be included following the name in brackets: e.g. *d2\_xyz[2]*, *i2\_mySpin[3]*.

#### 16.119.4 Restrictions

You cannot set an atom attribute (e.g. *mol* or *q* or *volume*) if the *atom\_style* does not have that attribute.

This command requires inter-processor communication to coordinate the setting of bond types (angle types, etc). This means that your system must be ready to perform a simulation before using one of these keywords (force fields set, atom mass set, etc). This is not necessary for other keywords.

Using the *region* style with the bond (angle, etc) keywords can give unpredictable results if there are bonds (angles, etc) that straddle periodic boundaries. This is because the region may only extend up to the boundary and partner atoms in the bond (angle, etc) may have coordinates outside the simulation box if they are ghost atoms.

#### 16.119.5 Related commands

*create\_box*, *create\_atoms*, *read\_data*

## 16.119.6 Default

none

## 16.120 shell command

### 16.120.1 Syntax

```
shell command args
```

- `command` = `cd` or `mkdir` or `mv` or `rm` or `rmdir` or `putenv` or arbitrary command

`cd` arg = dir

dir = directory to change to

`mkdir` args = dir1 dir2 ...

dir1,dir2 = one or more directories to create

`mv` args = old new

old = old filename

new = new filename or destination folder

`rm` args = [-f] file1 file2 ...

-f = turn off warnings (optional)

file1,file2 = one or more filenames to delete

`rmdir` args = dir1 dir2 ...

dir1,dir2 = one or more directories to delete

`putenv` args = var1=value1 var2=value2

var=value = one of more definitions of environment variables

anything else is passed as a command to the shell for direct execution

### 16.120.2 Examples

```
shell cd sub1
shell cd ..
shell mkdir tmp1 tmp2/tmp3
shell rmdir tmp1 tmp2
shell mv log.lammps hold/log.1
shell rm TMP/file1 TMP/file2
shell putenv LAMMPS_POTENTIALS=../../potentials
shell my_setup file1 10 file2
shell my_post_process 100 dump.out
```

### 16.120.3 Description

Execute a shell command. A few simple file-based shell commands are supported directly, in Unix-style syntax. Any command not listed above is passed as-is to the `C-library system()` call, which invokes the command in a shell. To use the external executable instead of the built-in version one needs to use a full path, for example `/bin/rm` instead of `rm`. The built-in commands will also work on operating systems, that do not - by default - provide the corresponding external executables (like `mkdir` on Windows).

This command provides a way to invoke custom commands or executables from your input script. For example, you can move files around in preparation for the next section of the input script. Or you can run a program that pre-processes data for input into LAMMPS. Or you can run a program that post-processes LAMMPS output data.

With the exception of *cd*, all commands, including ones invoked via a `system()` call, are executed by only a single processor, so that files/directories are not being manipulated by multiple processors concurrently which may result in unexpected errors or corrupted files.

The *cd* command changes the current working directory similar to the `cd` command. All subsequent LAMMPS commands that read/write files will use the new directory. All processors execute this command.

The *mkdir* command creates directories similar to the Unix `mkdir -p` command. That is, it will attempt to create the entire path of sub-directories if they do not exist yet.

The *mv* command renames a file and/or moves it to a new directory. It cannot rename files across filesystem boundaries or between drives.

The *rm* command deletes file similar to the Unix `rm` command.

The *rmdir* command deletes directories similar to Unix `rmdir` command. If a directory is not empty, its contents are also removed recursively similar to the Unix `rm -r` command.

The *putenv* command defines or updates an environment variable directly. Since this command does not pass through the shell, no shell variable expansion or globbing is performed, only the usual substitution for LAMMPS variables defined with the *variable* command is performed. The resulting string is then used literally.

Any other command is passed as-is to the shell along with its arguments as one string, invoked by the C-library `system()` call. For example, these lines in your input script:

```
variable n equal 10
variable foo string file2
shell my_setup file1 $n ${foo}
```

would be the same as invoking

```
% my_setup file1 10 file2
```

from a command-line prompt. The executable program “my\_setup” is run with 3 arguments: file1 10 file2.

## 16.120.4 Restrictions

LAMMPS will do a best effort to detect errors and print suitable warnings, but due to the nature of delegating commands to the C-library `system()` call, this is not always reliable.

## 16.120.5 Related commands

none

## 16.120.6 Default

none

## 16.121 special\_bonds command

### 16.121.1 Syntax

```
special_bonds keyword values ...
```

- one or more keyword/value pairs may be appended
- keyword = *amber* or *charmm* or *dreiding* or *fene* or *lj/coul* or *lj* or *coul* or *angle* or *dihedral* or *one/five*

*amber* values = none

*charmm* values = none

*dreiding* values = none

*fene* values = none

*lj/coul* values = w1,w2,w3

w1,w2,w3 = weights (0.0 to 1.0) on pairwise Lennard-Jones and Coulombic interactions

*lj* values = w1,w2,w3

w1,w2,w3 = weights (0.0 to 1.0) on pairwise Lennard-Jones interactions

*coul* values = w1,w2,w3

w1,w2,w3 = weights (0.0 to 1.0) on pairwise Coulombic interactions

*angle* value = *yes* or *no*

*dihedral* value = *yes* or *no*

*one/five* value = *yes* or *no*

### 16.121.2 Examples

```
special_bonds amber
special_bonds charmm
special_bonds fene dihedral no
special_bonds lj/coul 0.0 0.0 0.5 angle yes dihedral yes
special_bonds lj 0.0 0.0 0.5 coul 0.0 0.0 0.0 dihedral yes
```

### 16.121.3 Description

Set weighting coefficients for pairwise energy and force contributions between pairs of atoms that are also permanently bonded to each other, either directly or via one or two intermediate bonds. These weighting factors are used by nearly all *pair styles* in LAMMPS that compute simple pairwise interactions. Permanent bonds between atoms are specified by defining the bond topology in the data file read by the *read\_data* command. Typically a *bond\_style* command is also used to define a bond potential. The rationale for using these weighting factors is that the interaction between a pair of bonded atoms is all (or mostly) specified by the bond, angle, dihedral potentials, and thus the non-bonded Lennard-Jones or Coulombic interaction between the pair of atoms should be excluded (or reduced by a weighting factor).

---

**Note:** These weighting factors are NOT used by *pair styles* that compute many-body interactions, since the “bonds” that result from such interactions are not permanent, but are created and broken dynamically as atom conformations change.



Examples of pair styles in this category are EAM, MEAM, Stillinger-Weber, Tersoff, COMB, AIREBO, and ReaxFF. In fact, it generally makes no sense to define permanent bonds between atoms that interact via these potentials, though such bonds may exist elsewhere in your system, e.g. when using the *pair\_style hybrid* command. Thus LAMMPS ignores *special\_bonds* settings when many-body potentials are calculated. Please note, that the existence of explicit bonds for atoms that are described by a many-body potential will alter the neighbor list and thus can render the computation of those interactions invalid, since those pairs are not only used to determine direct pairwise interactions but also neighbors of neighbors and more. The recommended course of action is to remove such bonds, or - if that is not possible - use a *special\_bonds* setting of 1.0 1.0 1.0.

---

**Note:** Unlike some commands in LAMMPS, you cannot use this command multiple times in an incremental fashion: e.g. to first set the LJ settings and then the Coulombic ones. Each time you use this command it sets all the coefficients to default values and only overrides the one you specify, so you should set all the options you need each time you use it. See more details at the bottom of this page.

---

The Coulomb factors are applied to any Coulomb (charge interaction) term that the potential calculates. The LJ factors are applied to the remaining terms that the potential calculates, whether they represent LJ interactions or not. The weighting factors are a scaling prefactor on the energy and force between the pair of atoms. A value of 1.0 means include the full interaction; a value of 0.0 means exclude it completely.

The first of the 3 coefficients (LJ or Coulombic) is the weighting factor on 1-2 atom pairs, which are pairs of atoms directly bonded to each other. The second coefficient is the weighting factor on 1-3 atom pairs which are those separated by 2 bonds (e.g. the two H atoms in a water molecule). The third coefficient is the weighting factor on 1-4 atom pairs which are those separated by 3 bonds (e.g. the first and fourth atoms in a dihedral interaction). Thus if the 1-2 coefficient is set to 0.0, then the pairwise interaction is effectively turned off for all pairs of atoms bonded to each other. If it is set to 1.0, then that interaction will be at full strength.

---

**Note:** For purposes of computing weighted pairwise interactions, 1-3 and 1-4 interactions are not defined from the list of angles or dihedrals used by the simulation. Rather, they are inferred topologically from the set of bonds specified when the simulation is defined from a data or restart file (see *read\_data* or *read\_restart* commands). Thus the set of 1-2,1-3,1-4 interactions that the weights apply to is the same whether angle and dihedral potentials are computed or not, and remains the same even if bonds are constrained, or turned off, or removed during a simulation.

---

The two exceptions to this rule are (a) if the *angle* or *dihedral* keywords are set to *yes* (see below), or (b) if the *delete\_bonds* command is used with the *special* option that re-computes the 1-2,1-3,1-4 topologies after bonds are deleted; see the *delete\_bonds* command for more details.

The *amber* keyword sets the 3 coefficients to 0.0, 0.0, 0.5 for LJ interactions and to 0.0, 0.0, 0.8333 for Coulombic interactions, which is the default for a commonly used version of the AMBER force field, where the last value is really 5/6. See (*Cornell*) for a description of the AMBER force field.

The *charmm* keyword sets the 3 coefficients to 0.0, 0.0, 0.0 for both LJ and Coulombic interactions, which is the default for a commonly used version of the CHARMM force field. Note that in pair styles *lj/charmm/coul/charmm* and *lj/charmm/coul/long* the 1-4 coefficients are defined explicitly, and these pairwise contributions are computed as part of the charmm dihedral style - see the *pair\_coeff* and *dihedral\_style* commands for more information. See (*MacKerell*) for a description of the CHARMM force field.

The *dreiding* keyword sets the 3 coefficients to 0.0, 0.0, 1.0 for both LJ and Coulombic interactions, which is the default for the Dreiding force field, as discussed in (*Mayo*).

The *fene* keyword sets the 3 coefficients to 0.0, 1.0, 1.0 for both LJ and Coulombic interactions, which is consistent with a coarse-grained polymer model with *FENE bonds*. See (*Kremer*) for a description of FENE bonds.

The *lj/coul*, *lj*, and *coul* keywords allow the 3 coefficients to be set explicitly. The *lj/coul* keyword sets both the LJ and Coulombic coefficients to the same 3 values. The *lj* and *coul* keywords only set either the LJ or Coulombic coefficients.

Use both of them if you wish to set the LJ coefficients to different values than the Coulombic coefficients.

The *angle* keyword allows the 1-3 weighting factor to be ignored for individual atom pairs if they are not listed as the first and last atoms in any angle defined in the simulation or as 1,3 or 2,4 atoms in any dihedral defined in the simulation. For example, imagine the 1-3 weighting factor is set to 0.5 and you have a linear molecule with 4 atoms and bonds as follows: 1-2-3-4. If your data file defines 1-2-3 as an angle, but does not define 2-3-4 as an angle or 1-2-3-4 as a dihedral, then the pairwise interaction between atoms 1 and 3 will always be weighted by 0.5, but different force fields use different rules for weighting the pairwise interaction between atoms 2 and 4. If the *angle* keyword is specified as *yes*, then the pairwise interaction between atoms 2 and 4 will be unaffected (full weighting of 1.0). If the *angle* keyword is specified as *no* which is the default, then the 2,4 interaction will also be weighted by 0.5.

The *dihedral* keyword allows the 1-4 weighting factor to be ignored for individual atom pairs if they are not listed as the first and last atoms in any dihedral defined in the simulation. For example, imagine the 1-4 weighting factor is set to 0.5 and you have a linear molecule with 5 atoms and bonds as follows: 1-2-3-4-5. If your data file defines 1-2-3-4 as a dihedral, but does not define 2-3-4-5 as a dihedral, then the pairwise interaction between atoms 1 and 4 will always be weighted by 0.5, but different force fields use different rules for weighting the pairwise interaction between atoms 2 and 5. If the *dihedral* keyword is specified as *yes*, then the pairwise interaction between atoms 2 and 5 will be unaffected (full weighting of 1.0). If the *dihedral* keyword is specified as *no* which is the default, then the 2,5 interaction will also be weighted by 0.5.

The *one/five* keyword enable calculation and storage of a list of 1-5 neighbors in the molecular topology for each atom. It is required by some pair styles, such as *pair\_style amoeba* and *pair\_style hippo*.

---

**Note:** LAMMPS stores and maintains a data structure with a list of the first, second, and third neighbors of each atom (within the bond topology of the system). If new bonds are created (or molecules added containing atoms with more special neighbors), the size of this list needs to grow. Note that adding a single bond always adds a new first neighbor but may also induce \*many\* new second and third neighbors, depending on the molecular topology of your system. Using the *extra/special/per/atom* keyword to either *read\_data* or *create\_box* reserves empty space in the list for this N additional first, second, or third neighbors to be added. If you do not do this, you may get an error when bonds (or molecules) are added.

---

**Note:** If you reuse this command in an input script, you should set all the options you need each time. This command cannot be used a second time incrementally. E.g. these two commands:

---

```
special_bonds lj 0.0 1.0 1.0
special_bonds coul 0.0 0.0 1.0
```

are not the same as

```
special_bonds lj 0.0 1.0 1.0 coul 0.0 0.0 1.0
```

In the first case you end up with (after the second command):

```
LJ: 0.0 0.0 0.0
Coul: 0.0 0.0 1.0
```

while only in the second case do you get the desired settings of:

```
LJ: 0.0 1.0 1.0
Coul: 0.0 0.0 1.0
```

This happens because the LJ (and Coul) settings are reset to their default values before modifying them, each time the *special\_bonds* command is issued.

### 16.121.4 Restrictions

none

### 16.121.5 Related commands

*delete\_bonds*, *fix bond/create*

### 16.121.6 Default

All 3 Lennard-Jones and 3 Coulombic weighting coefficients = 0.0, angle = no, dihedral = no.

---

**(Cornell)** Cornell, Cieplak, Bayly, Gould, Merz, Ferguson, Spellmeyer, Fox, Caldwell, Kollman, JACS 117, 5179-5197 (1995).

**(Kremer)** Kremer, Grest, J Chem Phys, 92, 5057 (1990).

**(MacKerell)** MacKerell, Bashford, Bellott, Dunbrack, Evanseck, Field, Fischer, Gao, Guo, Ha, et al, J Phys Chem, 102, 3586 (1998).

**(Mayo)** Mayo, Olfason, Goddard III, J Phys Chem, 94, 8897-8909 (1990).

## 16.122 suffix command

### 16.122.1 Syntax

```
suffix style args
```

- style = *off* or *on* or *gpu* or *intel* or *kk* or *omp* or *opt* or *hybrid*
- args = for hybrid style, default suffix to be used and alternative suffix

### 16.122.2 Examples

```
suffix off
suffix on
suffix gpu
suffix intel
suffix hybrid intel omp
suffix kk
```

### 16.122.3 Description

This command allows you to use variants of various styles if they exist. In that respect it operates the same as the *-suffix command-line switch*. It also has options to turn off or back on any suffix setting made via the command line.

The specified style can be *gpu*, *intel*, *kk*, *omp*, *opt* or *hybrid*. These refer to optional packages that LAMMPS can be built with, as described on the *Build package* doc page. The “gpu” style corresponds to the GPU package, the “intel” style to the INTEL package, the “kk” style to the KOKKOS package, the “omp” style to the OPENMP package, and the “opt” style to the OPT package.

These are the variants these packages provide:

- GPU = a handful of pair styles and the PPPM *kpace\_style*, optimized to run on one or more GPUs or multicore CPU/GPU nodes
- INTEL = a collection of pair styles and neighbor routines optimized to run in single, mixed, or double precision on CPUs and Intel(R) Xeon Phi(TM) co-processors.
- KOKKOS = a collection of atom, pair, and fix styles optimized to run using the Kokkos library on various kinds of hardware, including GPUs via CUDA and many-core chips via OpenMP or threading.
- OPENMP = a collection of pair, bond, angle, dihedral, improper, *kpace*, compute, and fix styles with support for OpenMP multi-threading
- OPT = a handful of pair styles, cache-optimized for faster CPU performance
- HYBRID = a combination of two packages can be specified (see below)

As an example, all of the packages provide a *pair\_style lj/cut* variant, with style names *lj/cut/opt*, *lj/cut/omp*, *lj/cut/gpu*, *lj/cut/intel*, or *lj/cut/kk*. A variant styles can be specified explicitly in your input script, e.g. *pair\_style lj/cut/gpu*. If the suffix command is used with the appropriate style, you do not need to modify your input script. The specified suffix (*opt,omp,gpu,intel,kk*) is automatically appended whenever your input script command creates a new *atom*, *pair*, *bond*, *angle*, *dihedral*, *improper*, *kpace*, *fix*, *compute*, or *run* style. If the variant version does not exist, the standard version is created.

For “hybrid”, two packages are specified. The first is used whenever available. If a style with the first suffix is not available, the style with the suffix for the second package will be used if available. For example, “hybrid intel omp” will use styles from the INTEL package as a first choice and styles from the OPENMP package as a second choice if no INTEL variant is available.

If the specified style is *off*, then any previously specified suffix is temporarily disabled, whether it was specified by a command-line switch or a previous suffix command. If the specified style is *on*, a disabled suffix is turned back on. The use of these 2 commands lets your input script use a standard LAMMPS style (i.e. a non-accelerated variant), which can be useful for testing or benchmarking purposes. Of course this is also possible by not using any suffix commands, and explicitly appending or not appending the suffix to the relevant commands in your input script.

---

**Note:** The default *run\_style* verlet is invoked prior to reading the input script and is therefore not affected by a suffix command in the input script. The KOKKOS package requires “*run\_style verlet/kk*”, so when using the KOKKOS package it is necessary to either use the command line “-sf kk” command or add an explicit “*run\_style verlet*” command to the input script.

---

## 16.122.4 Restrictions

none

## 16.122.5 Related commands

*-suffix command-line switch*

## 16.122.6 Default

none

## 16.123 tad command

### 16.123.1 Syntax

```
tad N t_event T_lo T_hi delta tmax compute-ID keyword value ...
```

- N = # of timesteps to run (not including dephasing/quenching)
- t\_event = timestep interval between event checks
- T\_lo = temperature at which event times are desired
- T\_hi = temperature at which MD simulation is performed
- delta = desired confidence level for stopping criterion
- tmax = reciprocal of lowest expected pre-exponential factor (time units)
- compute-ID = ID of the compute used for event detection
- zero or more keyword/value pairs may be appended
- keyword = *min* or *neb* or *min\_style* or *neb\_style* or *neb\_log*

*min* values = etol ftol maxiter maxeval

etol = stopping tolerance for energy (energy units)

ftol = stopping tolerance for force (force units)

maxiter = max iterations of minimize

maxeval = max number of force/energy evaluations

*neb* values = ftol N1 N2 Nevery

etol = stopping tolerance for energy (energy units)

ftol = stopping tolerance for force (force units)

N1 = max # of iterations (timesteps) to run initial NEB

N2 = max # of iterations (timesteps) to run barrier-climbing NEB

Nevery = print NEB statistics every this many timesteps

*neb\_style* value = *quickmin* or *fire*

*neb\_step* value = dtneb

dtneb = timestep for NEB damped dynamics minimization

*neb\_log* value = file where NEB statistics are printed

## 16.123.2 Examples

```
tad 2000 50 1800 2300 0.01 0.01 event
tad 2000 50 1800 2300 0.01 0.01 event &
  min 1e-05 1e-05 100 100 &
  neb 0.0 0.01 200 200 20 &
  min_style cg &
  neb_style fire &
  neb_log log.neb
```

## 16.123.3 Description

Run a temperature accelerated dynamics (TAD) simulation. This method requires two or more partitions to perform NEB transition state searches.

TAD is described in [this paper](#) by Art Voter. It is a method that uses accelerated dynamics at an elevated temperature to generate results at a specified lower temperature. A good overview of accelerated dynamics methods (AMD) for such systems is given in [this review paper](#) from the same group. To quote from the review paper: “The dynamical evolution is characterized by vibrational excursions within a potential basin, punctuated by occasional transitions between basins. The transition probability is characterized by  $p(t) = k \cdot \exp(-kt)$  where  $k$  is the rate constant.”

TAD is a suitable AMD method for infrequent-event systems, where in addition, the transition kinetics are well-approximated by harmonic transition state theory (hTST). In hTST, the temperature dependence of transition rates follows the Arrhenius relation. As a consequence a set of event times generated in a high-temperature simulation can be mapped to a set of much longer estimated times in the low-temperature system. However, because this mapping involves the energy barrier of the transition event, which is different for each event, the first event at the high temperature may not be the earliest event at the low temperature. TAD handles this by first generating a set of possible events from the current basin. After each event, the simulation is reflected backwards into the current basin. This is repeated until the stopping criterion is satisfied, at which point the event with the earliest low-temperature occurrence time is selected. The stopping criterion is that the confidence measure be greater than  $1-\delta$ . The confidence measure is the probability that no earlier low-temperature event will occur at some later time in the high-temperature simulation. hTST provides an lower bound for this probability, based on the user-specified minimum pre-exponential factor (reciprocal of  $t_{max}$ ).

In order to estimate the energy barrier for each event, the TAD method invokes the *NEB* method. Each NEB replica runs on a partition of processors. The current NEB implementation in LAMMPS restricts you to having exactly one processor per replica. For more information, see the documentation for the *neb* command. In the current LAMMPS implementation of TAD, all the non-NEB TAD operations are performed on the first partition, while the other partitions remain idle. See the [Howto replica](#) doc page for further discussion of multi-replica simulations.

A TAD run has several stages, which are repeated each time an event is performed. The logic for a TAD run is as follows:

```
while (time remains):
  while (time < tstop):
    until (event occurs):
      run dynamics for t_event steps
      quench
      run neb calculation using all replicas
      compute tlo from energy barrier
      update earliest event
      update tstop
      reflect back into current basin
    execute earliest event
```

Before this outer loop begins, the initial potential energy basin is identified by quenching (an energy minimization, see below) the initial state and storing the resulting coordinates for reference.

Inside the inner loop, dynamics is run continuously according to whatever integrator has been specified by the user, stopping every  $t_{event}$  steps to check if a transition event has occurred. This check is performed by quenching the system and comparing the resulting atom coordinates to the coordinates from the previous basin.

A quench is an energy minimization and is performed by whichever algorithm has been defined by the *min\_style* command; its default is the CG minimizer. The tolerances and limits for each quench can be set by the *min* keyword. Note that typically, you do not need to perform a highly-converged minimization to detect a transition event.

The event check is performed by a compute with the specified *compute-ID*. Currently there is only one compute that works with the TAD command, which is the *compute event/displace* command. Other event-checking computes may be added. *Compute event/displace* checks whether any atom in the compute group has moved further than a specified threshold distance. If so, an “event” has occurred.

The NEB calculation is similar to that invoked by the *neb* command, except that the final state is generated internally, instead of being read in from a file. The style of minimization performed by NEB is determined by the *neb\_style* keyword and must be a damped dynamics minimizer. The tolerances and limits for each NEB calculation can be set by the *neb* keyword. As discussed on the *neb*, it is often advantageous to use a larger timestep for NEB than for normal dynamics. Since the size of the timestep set by the *timestep* command is used by TAD for performing dynamics, there is a *neb\_step* keyword which can be used to set a larger timestep for each NEB calculation if desired.

A key aspect of the TAD method is setting the stopping criterion appropriately. If this criterion is too conservative, then many events must be generated before one is finally executed. Conversely, if this criterion is too aggressive, high-entropy high-barrier events will be over-sampled, while low-entropy low-barrier events will be under-sampled. If the lowest pre-exponential factor is known fairly accurately, then it can be used to estimate  $t_{max}$ , and the value of *delta* can be set to the desired confidence level e.g.  $delta = 0.05$  corresponds to 95% confidence. However, for systems where the dynamics are not well characterized (the most common case), it will be necessary to experiment with the values of *delta* and  $t_{max}$  to get a good trade-off between accuracy and performance.

A second key aspect is the choice of  $t_{hi}$ . A larger value greatly increases the rate at which new events are generated. However, too large a value introduces errors due to anharmonicity (not accounted for within hTST). Once again, for any given system, experimentation is necessary to determine the best value of  $t_{hi}$ .

Five kinds of output can be generated during a TAD run: event statistics, NEB statistics, thermodynamic output by each replica, dump files, and restart files.

Event statistics are printed to the screen and master log.lammps file each time an event is executed. The quantities are the timestep, CPU time, global event number  $N$ , local event number  $M$ , event status, energy barrier, time margin,  $t_{lo}$  and  $delt_{lo}$ . The timestep is the usual LAMMPS timestep, which corresponds to the high-temperature time at which the event was detected, in units of timestep. The CPU time is the total processor time since the start of the TAD run. The global event number  $N$  is a counter that increments with each executed event. The local event number  $M$  is a counter that resets to zero upon entering each new basin. The event status is  $E$  when an event is executed, and is  $D$  for an event that is detected, while  $DF$  is for a detected event that is also the earliest (first) event at the low temperature.

The time margin is the ratio of the high temperature time in the current basin to the stopping time. This last number can be used to judge whether the stopping time is too short or too long (see above).

$t_{lo}$  is the low-temperature event time when the current basin was entered, in units of timestep.  $delt_{lo}$  is the time of each detected event, measured relative to  $t_{lo}$ .  $delt_{lo}$  is equal to the high-temperature time since entering the current basin, scaled by an exponential factor that depends on the hi/lo temperature ratio and the energy barrier for that event.

On lines for executed events, with status  $E$ , the global event number is incremented by one, the local event number and time margin are reset to zero, while the global event number, energy barrier, and  $delt_{lo}$  match the last event with

status *DF* in the immediately preceding block of detected events. The low-temperature event time *t\_lo* is incremented by *delt\_lo*.

NEB statistics are written to the file specified by the *neb\_log* keyword. If the keyword value is “none”, then no NEB statistics are printed out. The statistics are written every *Nevery* timesteps. See the *neb* command for a full description of the NEB statistics. When invoked from TAD, NEB statistics are never printed to the screen.

Because the NEB calculation must run on multiple partitions, LAMMPS produces additional screen and log files for each partition, e.g. *log.lammps.0*, *log.lammps.1*, etc. For the TAD command, these contain the thermodynamic output of each NEB replica. In addition, the log file for the first partition, *log.lammps.0*, will contain thermodynamic output from short runs and minimizations corresponding to the dynamics and quench operations, as well as a line for each new detected event, as described above.

After the TAD command completes, timing statistics for the TAD run are printed in each replica’s log file, giving a breakdown of how much CPU time was spent in each stage (NEB, dynamics, quenching, etc).

Any *dump files* defined in the input script will be written to during a TAD run at timesteps when an event is executed. This means the requested dump frequency in the *dump* command is ignored. There will be one dump file (per dump command) created for all partitions. The atom coordinates of the dump snapshot are those of the minimum energy configuration resulting from quenching following the executed event. The timesteps written into the dump files correspond to the timestep at which the event occurred and NOT the clock. A dump snapshot corresponding to the initial minimum state used for event detection is written to the dump file at the beginning of each TAD run.

If the *restart* command is used, a single restart file for all the partitions is generated, which allows a TAD run to be continued by a new input script in the usual manner. The restart file is generated after an event is executed. The restart file contains a snapshot of the system in the new quenched state, including the event number and the low-temperature time. The restart frequency specified in the *restart* command is interpreted differently when performing a TAD run. It does not mean the timestep interval between restart files. Instead it means an event interval for executed events. Thus a frequency of 1 means write a restart file every time an event is executed. A frequency of 10 means write a restart file every 10th executed event. When an input script reads a restart file from a previous TAD run, the new script can be run on a different number of replicas or processors.

Note that within a single state, the dynamics will typically temporarily continue beyond the event that is ultimately chosen, until the stopping criterion is satisfied. When the event is eventually executed, the timestep counter is reset to the value when the event was detected. Similarly, after each quench and NEB minimization, the timestep counter is reset to the value at the start of the minimization. This means that the timesteps listed in the replica log files do not always increase monotonically. However, the timestep values printed to the master log file, dump files, and restart files are always monotonically increasing.

---

### 16.123.4 Restrictions

This command can only be used if LAMMPS was built with the REPLICA package. See the *Build package* doc page for more info.

*N* setting must be integer multiple of *t\_event*.

Runs restarted from restart files written during a TAD run will only produce identical results if the user-specified integrator supports exact restarts. So *fix nvt* will produce an exact restart, but *fix langevin* will not.

This command cannot be used when any fixes are defined that keep track of elapsed time to perform time-dependent operations. Examples include the “ave” fixes such as *fix ave/chunk*. Also *fix dt/reset* and *fix deposit*.



### 16.123.5 Related commands

*compute event/displace, min\_modify, min\_style, run\_style, minimize, temper, neb, prd*

### 16.123.6 Default

The option defaults are *min* = 0.1 0.1 40 50, *neb* = 0.01 100 100 10, *neb\_style* = *quickmin*, *neb\_step* = the same timestep set by the *timestep* command, and *neb\_log* = “none”.

(**Voter2000**) Sorensen and Voter, J Chem Phys, 112, 9599 (2000)

(**Voter2002**) Voter, Montalenti, Germann, Annual Review of Materials Research 32, 321 (2002).

## 16.124 temper command

### 16.124.1 Syntax

```
temper N M temp fix-ID seed1 seed2 index
```

- N = total # of timesteps to run
- M = attempt a tempering swap every this many steps
- temp = initial temperature for this ensemble
- fix-ID = ID of the fix that will control temperature during the run
- seed1 = random # seed used to decide on adjacent temperature to partner with
- seed2 = random # seed for Boltzmann factor in Metropolis swap
- index = which temperature (0 to N-1) I am simulating (optional)

### 16.124.2 Examples

```
temper 100000 100 $t tempfix 0 58728
temper 40000 100 $t tempfix 0 32285 $w
```

### 16.124.3 Description

Run a parallel tempering or replica exchange simulation using multiple replicas (ensembles) of a system. Two or more replicas must be used.

Each replica runs on a partition of one or more processors. Processor partitions are defined at run-time using the *-partition command-line switch*. Note that if you have MPI installed, you can run a multi-replica simulation with more replicas (partitions) than you have physical processors, e.g you can run a 10-replica simulation on one or two processors. You will simply not get the performance speed-up you would see with one or more physical processors per replica. See the *Howto replica* doc page for further discussion.

Each replica’s temperature is controlled at a different value by a fix with *fix-ID* that controls temperature. Most thermostat fix styles (with and without included time integration) are supported. The command will print an error message and abort, if the chosen fix is unsupported. The desired temperature is specified by *temp*, which is typically a variable

previously set in the input script, so that each partition is assigned a different temperature. See the *variable* command for more details. For example:

```
variable t world 300.0 310.0 320.0 330.0
fix myfix all nvt temp $t $t 100.0
temper 100000 100 $t myfix 3847 58382
```

would define 4 temperatures, and assign one of them to the thermostat used by each replica, and to the temper command.

As the tempering simulation runs for  $N$  timesteps, a temperature swap between adjacent ensembles will be attempted every  $M$  timesteps. If *seed1* is 0, then the swap attempts will alternate between odd and even pairings. If *seed1* is non-zero then it is used as a seed in a random number generator to randomly choose an odd or even pairing each time. Each attempted swap of temperatures is either accepted or rejected based on a Boltzmann-weighted Metropolis criterion which uses *seed2* in the random number generator.

As a tempering run proceeds, multiple log files and screen output files are created, one per replica. By default these files are named log.lammps.M and screen.M where M is the replica number from 0 to N-1, with  $N = \#$  of replicas. See the *-log and -screen command-line switches* for info on how to change these names.

The main screen and log file (log.lammps) will list information about which temperature is assigned to each replica at each thermodynamic output timestep. E.g. for a simulation with 16 replicas:

```
Running on 16 partitions of processors
Step T0 T1 T2 T3 T4 T5 T6 T7 T8 T9 T10 T11 T12 T13 T14 T15
0    0 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15
500  1 0 3 2 5 4 6 7 8 9 10 11 12 13 14 15
1000 2 0 4 1 5 3 6 7 8 9 10 11 12 14 13 15
1500 2 1 4 0 5 3 6 7 9 8 10 11 12 14 13 15
2000 2 1 3 0 6 4 5 7 10 8 9 11 12 14 13 15
2500 2 1 3 0 6 4 5 7 11 8 9 10 12 14 13 15
...
```

The column headings T0 to TN-1 mean which temperature is currently assigned to the replica 0 to N-1. Thus the columns represent replicas and the value in each column is its temperature (also numbered 0 to N-1). For example, a 0 in the fourth column (column T3, step 2500) means that the fourth replica is assigned temperature 0, i.e. the lowest temperature. You can verify this time sequence of temperature assignments for the Nth replica by comparing the Nth column of screen output to the thermodynamic data in the corresponding log.lammps.N or screen.N files as time proceeds.

You can have each replica create its own dump file in the following manner:

```
variable rep world 0 1 2 3 4 5 6 7
dump 1 all atom 1000 dump.temper.${rep}
```

---

**Note:** Each replica's dump file will contain a continuous trajectory for its atoms where the temperature varies over time as swaps take place involving that replica. If you want a series of dump files, each with snapshots (from all replicas) that are all at a single temperature, then you will need to post-process the dump files using the information from the log.lammps file. E.g. you could produce one dump file with snapshots at 300K (from all replicas), another with snapshots at 310K, etc. Note that these new dump files will not contain "continuous trajectories" for individual atoms, because two successive snapshots (in time) may be from different replicas. The reorder\_remd\_traj python script can do the reordering for you (and additionally also calculated configurational log-weights of trajectory snapshots in the canonical ensemble). The script can be found in the tools/replica directory while instructions on how to use it is available in doc/Tools (in brief) and as a README file in tools/replica (in detail).

---

The last argument *index* in the temper command is optional and is used when restarting a tempering run from a set of

restart files (one for each replica) which had previously swapped to new temperatures. The *index* value (from 0 to N-1, where N is the # of replicas) identifies which temperature the replica was simulating on the timestep the restart files were written. Obviously, this argument must be a variable so that each partition has the correct value. Set the variable to the *N* values listed in the log file for the previous run for the replica temperatures at that timestep. For example if the log file listed the following for a simulation with 5 replicas:

```
500000 2 4 0 1 3
```

then a setting of

```
variable w world 2 4 0 1 3
```

would be used to restart the run with a tempering command like the example above with \$w as the last argument.

### 16.124.4 Restrictions

This command can only be used if LAMMPS was built with the REPLICAs package. See the *Build package* doc page for more info.

### 16.124.5 Related commands

*variable, prd, neb*

### 16.124.6 Default

none

## 16.125 temper/grem command

### 16.125.1 Syntax

```
temper/grem N M lambda fix-ID thermostat-ID seed1 seed2 index
```

- N = total # of timesteps to run
- M = attempt a tempering swap every this many steps
- lambda = initial lambda for this ensemble
- fix-ID = ID of *fix grem*
- thermostat-ID = ID of the thermostat that controls kinetic temperature
- seed1 = random # seed used to decide on adjacent temperature to partner with
- seed2 = random # seed for Boltzmann factor in Metropolis swap
- index = which temperature (0 to N-1) I am simulating (optional)

## 16.125.2 Examples

```
temper/grem 100000 1000 ${lambda} fxgREM fxnvt 0 58728
temper/grem 40000 100 ${lambda} fxgREM fxnpt 0 32285 ${walkers}
```

## 16.125.3 Description

Run a parallel tempering or replica exchange simulation in LAMMPS partition mode using multiple generalized replicas (ensembles) of a system defined by *fix grem*, which stands for the generalized replica exchange method (gREM) originally developed by (Kim). It uses non-Boltzmann ensembles to sample over first order phase transitions. The is done by defining replicas with an enthalpy dependent effective temperature

Two or more replicas must be used. See the *temper* command for an explanation of how to run replicas on multiple partitions of one or more processors.

This command is a modification of the *temper* command and has the same dependencies, restraints, and input variables which are discussed there in greater detail.

Instead of temperature, this command performs replica exchanges in lambda as per the generalized ensemble enforced by *fix grem*. The desired lambda is specified by *lambda*, which is typically a variable previously set in the input script, so that each partition is assigned a different temperature. See the *variable* command for more details. For example:

```
variable lambda world 400 420 440 460
fix fxnvt all nvt temp 300.0 300.0 100.0
fix fxgREM all grem ${lambda} -0.05 -50000 fxnvt
temper/grem 100000 100 ${lambda} fxgREM fxnvt 3847 58382
```

would define 4 lambdas with constant kinetic temperature but unique generalized temperature, and assign one of them to *fix grem* used by each replica, and to the *grem* command.

As the gREM simulation runs for  $N$  timesteps, a swap between adjacent ensembles will be attempted every  $M$  timesteps. If *seed1* is 0, then the swap attempts will alternate between odd and even pairings. If *seed1* is non-zero then it is used as a seed in a random number generator to randomly choose an odd or even pairing each time. Each attempted swap of temperatures is either accepted or rejected based on a Metropolis criterion, derived for gREM by (Kim), which uses *seed2* in the random number generator.

File management works identical to the *temper* command. Dump files created by this fix contain continuous trajectories and require post-processing to obtain per-replica information.

The last argument *index* in the *grem* command is optional and is used when restarting a run from a set of restart files (one for each replica) which had previously swapped to new lambda. This is done using a variable. For example if the log file listed the following for a simulation with 5 replicas:

```
500000 2 4 0 1 3
```

then a setting of

```
variable walkers world 2 4 0 1 3
```

would be used to restart the run with a *grem* command like the example above with  $\{\text{walkers}\}$  as the last argument. This functionality is identical to *temper*.

---

### 16.125.4 Restrictions

This command can only be used if LAMMPS was built with the REPLICA package. See the *Build package* doc page for more info.

This command must be used with *fix grem*.

### 16.125.5 Related commands

*fix grem*, *temper*, *variable*

### 16.125.6 Default

none

(Kim) Kim, Keyes, Straub, J Chem Phys, 132, 224107 (2010).

## 16.126 temper/npt command

### 16.126.1 Syntax

```
temper/npt N M temp fix-ID seed1 seed2 pressure index
```

- N = total # of timesteps to run
- M = attempt a tempering swap every this many steps
- temp = initial temperature for this ensemble
- fix-ID = ID of the fix that will control temperature and pressure during the run
- seed1 = random # seed used to decide on adjacent temperature to partner with
- seed2 = random # seed for Boltzmann factor in Metropolis swap
- pressure = setpoint pressure for the ensemble
- index = which temperature (0 to N-1) I am simulating (optional)

### 16.126.2 Examples

```
temper/npt 100000 100 $t nptfix 0 58728 1
temper/npt 2500000 1000 300 nptfix 0 32285 $p
temper/npt 5000000 2000 $t nptfix 0 12523 1 $w
```

### 16.126.3 Description

Run a parallel tempering or replica exchange simulation using multiple replicas (ensembles) of a system in the isothermal-isobaric (NPT) ensemble. The command `temper/npt` works like `temper` but requires running replicas in the NPT ensemble instead of the canonical (NVT) ensemble and allows for pressure to be set in the ensembles. These multiple ensembles can run in parallel at different temperatures or different pressures. The acceptance criteria for `temper/npt` is specific to the NPT ensemble and can be found in references (*Okabe*) and (*Mori*).

Apart from the difference in acceptance criteria and the specification of pressure, this command works much like the `temper` command. See the documentation on `temper` for information on how the parallel tempering is handled in general.

---

### 16.126.4 Restrictions

This command can only be used if LAMMPS was built with the REPLICA package. See the *Build package* page for more info.

This command should be used with a fix that maintains the isothermal-isobaric (NPT) ensemble.

### 16.126.5 Related commands

*temper, variable, fix\_npt*

### 16.126.6 Default

none

**(Okabe)** T. Okabe, M. Kawata, Y. Okamoto, M. Masuhiro, Chem. Phys. Lett., 335, 435-439 (2001).

**(Mori)** Y. Mori, Y. Okamoto, J. Phys. Soc. Jpn., 7, 074003 (2010).

## 16.127 thermo command

### 16.127.1 Syntax

```
thermo N
```

- N = output thermodynamics every N timesteps
- N can be a variable (see below)

## 16.127.2 Examples

```
thermo 100
```

## 16.127.3 Description

Compute and print thermodynamic info (e.g. temperature, energy, pressure) on timesteps that are a multiple of N and at the beginning and end of a simulation. A value of 0 will only print thermodynamics at the beginning and end.

The content and format of what is printed is controlled by the *thermo\_style* and *thermo\_modify* commands.

Instead of a numeric value, N can be specified as an *equal-style variable*, which should be specified as v\_name, where name is the variable name. In this case, the variable is evaluated at the beginning of a run to determine the next timestep at which thermodynamic info will be written out. On that timestep, the variable will be evaluated again to determine the next timestep, etc. Thus the variable should return timestep values. See the stagger() and logfreq() and stride() math functions for *equal-style variables*, as examples of useful functions to use in this context. Other similar math functions could easily be added as options for *equal-style variables*.

For example, the following commands will output thermodynamic info at timesteps 0, 10, 20, 30, 100, 200, 300, 1000, 2000, etc:

```
variable      s equal logfreq(10,3,10)
thermo        v_s
```

## 16.127.4 Restrictions

none

## 16.127.5 Related commands

*thermo\_style*, *thermo\_modify*

## 16.127.6 Default

```
thermo 0
```

## 16.128 thermo\_modify command

### 16.128.1 Syntax

```
thermo_modify keyword value ...
```

- one or more keyword/value pairs may be listed
- keyword = *lost* or *lost/bond* or *warn* or *norm* or *flush* or *line* or *colname* or *format* or *temp* or *press*

*lost* value = *error* or *warn* or *ignore*  
*lost/bond* value = *error* or *warn* or *ignore*  
*warn* value = *ignore* or *reset* or *default* or a number  
*norm* value = *yes* or *no*  
*flush* value = *yes* or *no*  
*line* value = *one* or *multi* or *yaml*  
*colname* values = ID string, or *default*  
    string = new column header name  
    ID = integer from 1 to N, or integer from -1 to -N, where N = # of quantities.  
    ↪being output  
        or a thermo keyword or reference to compute, fix, property or variable.  
*format* values = *line* string, *int* string, *float* string, ID string, or *none*  
    string = C-style format string  
    ID = integer from 1 to N, or integer from -1 to -N, where N = # of quantities.  
    ↪being output  
        or a thermo keyword or reference to compute, fix, property or variable.  
*temp* value = compute ID that calculates a temperature  
*press* value = compute ID that calculates a pressure

## 16.128.2 Examples

```
thermo_modify lost ignore flush yes
thermo_modify temp myTemp format 3 %15.8g
thermo_modify temp myTemp format line "%ld %g %g %15.8g"
thermo_modify line multi format float %g
thermo_modify line yaml format none
thermo_modify colname 1 Timestep colname -2 Pressure colname f_1[1] AvgDensity
```

## 16.128.3 Description

Set options for how thermodynamic information is computed and printed by LAMMPS.

---

**Note:** These options apply to the *currently defined* thermo style. When you specify a *thermo\_style* command, all thermodynamic settings are restored to their default values, including those previously reset by a *thermo\_modify* command. Thus if your input script specifies a *thermo\_style* command, you should use the *thermo\_modify* command **after** it.

---

The *lost* keyword determines whether LAMMPS checks for lost atoms each time it computes thermodynamics and what it does if atoms are lost. An atom can be “lost” if it moves across a non-periodic simulation box *boundary* or if it moves more than a box length outside the simulation domain (or more than a processor sub-domain length) before reneighboring occurs. The latter case is typically due to bad dynamics, e.g. too large a timestep or huge forces and velocities. If the value is *ignore*, LAMMPS does not check for lost atoms. If the value is *error* or *warn*, LAMMPS checks and either issues an error or warning. The code will exit with an error and continue with a warning. A warning will only be issued once, the first time an atom is lost. This can be a useful debugging option.

The *lost/bond* keyword determines whether LAMMPS throws an error or not if an atom in a bonded interaction (bond, angle, etc) cannot be found when it creates bonded neighbor lists. By default this is a fatal error. However in some scenarios it may be desirable to only issue a warning or ignore it and skip the computation of the missing bond, angle, etc. An example would be when gas molecules in a vapor are drifting out of the box through a fixed boundary condition (see the *boundary* command). In this case one atom may be deleted before the rest of the molecule is, on a later timestep.



The *warn* keyword allows you to control whether LAMMPS will print warning messages and how many of them. Most warning messages are only printed by MPI rank 0. They are usually pointing out important issues that should be investigated, but LAMMPS cannot determine for certain whether they are an indication of an error.

Some warning messages are printed during a run (or immediately before) each time a specific MPI rank encounters the issue, e.g. bonds that are stretched too far or dihedrals in extreme configurations. These number of these can quickly blow up the size of the log file and screen output. Thus a limit of 100 warning messages is applied by default. The warning count is applied to the entire input unless reset with a `thermo_modify warn reset` command. If there are more warnings than the limit, LAMMPS will print one final warning that it will not print any additional warning messages.

---

**Note:** The warning limit is enforced on either the per-processor count or the total count across all processors. For efficiency reasons, however, the total count is only updated at steps with thermodynamic output. Thus when running on a large number of processors in parallel, the total number of warnings printed can be significantly larger than the given limit.

---

Any number after the keyword *warn* will change the warning limit accordingly. With the value *ignore* all warnings will be suppressed, with the value *always* no limit will be applied and warnings will always be printed, with the value *reset* the internal warning counter will be reset to zero, and with the value *default*, the counter is reset and the limit set to 100. An example usage of either *reset* or *default* would be to re-enable warnings that were disabled or have reached the limit during equilibration, where the warnings would be acceptable while the system is still adjusting, but then change to all warnings for the production run, where they would indicate problems that would require a closer look at what is causing them.

The *norm* keyword determines whether various thermodynamic output values are normalized by the number of atoms or not, depending on whether it is set to *yes* or *no*. Different unit styles have different defaults for this setting (see below). Even if *norm* is set to *yes*, a value is only normalized if it is an “extensive” quantity, meaning that it scales with the number of atoms in the system. For the thermo keywords described by the page for the *thermo\_style* command, all energy-related keywords are extensive, such as *pe* or *ebond* or *enthalpy*. Other keywords such as *temp* or *press* are “intensive” meaning their value is independent (in a statistical sense) of the number of atoms in the system and thus are never normalized. For thermodynamic output values extracted from *fixes* and *computes* in a *thermo\_style custom* command, the page for the individual *fix* or *compute* lists whether the value is “extensive” or “intensive” and thus whether it is normalized. Thermodynamic output values calculated by a variable formula are assumed to be “intensive” and thus are never normalized. You can always include a divide by the number of atoms in the variable formula if this is not the case.

The *flush* keyword invokes a flush operation after thermodynamic info is written to the screen and log file. This insures the output is updated and not buffered (by the application) even if LAMMPS halts before the simulation completes. Please note that this does not affect buffering by the OS or devices, so you may still lose data in case the simulation stops due to a hardware failure.

The *line* keyword determines whether thermodynamics will be output as a series of numeric values on one line (“one”), in a multi-line format with 3 quantities with text strings per line and a dashed-line header containing the timestep and CPU time (“multi”), or in a YAML format block (“yaml”). This modify option overrides the *one*, *multi*, or *yaml* *thermo\_style* settings.

New in version 4May2022.

The *colname* keyword can be used to change the default header keyword for a column or field of thermodynamic output. The setting for *ID string* replaces the default text with the provided string. *ID* can be a positive integer when it represents the column number counting from the left, a negative integer when it represents the column number from the right (i.e. -1 is the last column/keyword), or a thermo keyword (or compute, fix, property, or variable reference) and then it replaces the string for that specific thermo keyword.

The *colname* keyword can be used multiple times. If multiple *colname* settings refer to the same keyword, the last setting has precedence. A setting of *default* clears all previous settings, reverting all values to their default values.

The *format* keyword can be used to change the default numeric format of any of quantities the *thermo\_style* command outputs. All the specified format strings are C-style formats, e.g. as used by the C/C++ `printf()` command. The *line* keyword takes a single argument which is the format string for the entire line of thermo output, with N fields, which you must enclose in quotes if it is more than one field. The *int* and *float* keywords take a single format argument and are applied to all integer or floating-point quantities output. The setting for *ID string* also takes a single format argument which is used for the indexed value in each line. The interpretation is the same as for *colname*, i.e. a positive integer is the n-th value corresponding to the n-th thermo keyword, a negative integer is counting backwards, and a string matches the entry with the thermo keyword., e.g. the fifth column is output in high precision for “format 5 %20.15g” and the pair energy for “format epair %20.15g”.

The *format* keyword can be used multiple times. The precedence is that for each value in a line of output, the *ID* format (if specified) is used, else the *int* or *float* setting (if specified) is used, else the *line* setting (if specified) for that value is used, else the default setting is used. A setting of *none* clears all previous settings, reverting all values to their default format.

---

**Note:** The thermo output values *step* and *atoms* are stored internally as 8-byte signed integers, rather than the usual 4-byte signed integers. When specifying the *format int* option you can use a “%d”-style format identifier in the format string and LAMMPS will convert this to the corresponding 8-byte form when it is applied to those keywords. However, when specifying the *line* option or *format ID string* option for *step* and *atoms*, you should specify a format string appropriate for an 8-byte signed integer, e.g. one with “%ld” or “%lld” depending on the platform.

---

The *temp* keyword is used to determine how thermodynamic temperature is calculated, which is used by all thermo quantities that require a temperature (“temp”, “press”, “ke”, “etotal”, “enthalpy”, “pxx”, etc). The specified compute ID must have been previously defined by the user via the *compute* command and it must be a style of compute that calculates a temperature. As described in the *thermo\_style* command, thermo output uses a default compute for temperature with ID = *thermo\_temp*. This option allows the user to override the default.

The *press* keyword is used to determine how thermodynamic pressure is calculated, which is used by all thermo quantities that require a pressure (“press”, “enthalpy”, “pxx”, etc). The specified compute ID must have been previously defined by the user via the *compute* command and it must be a style of compute that calculates a pressure. As described in the *thermo\_style* command, thermo output uses a default compute for pressure with ID = *thermo\_press*. This option allows the user to override the default.

---

**Note:** If both the *temp* and *press* keywords are used in a single *thermo\_modify* command (or in two separate commands), then the order in which the keywords are specified is important. Note that a *pressure compute* defines its own temperature compute as an argument when it is specified. The *temp* keyword will override this (for the pressure compute being used by thermodynamics), but only if the *temp* keyword comes after the *press* keyword. If the *temp* keyword comes before the *press* keyword, then the new pressure compute specified by the *press* keyword will be unaffected by the *temp* setting.

---

## 16.128.4 Restrictions

none

## 16.128.5 Related commands

*thermo*, *thermo\_style*

## 16.128.6 Default

The option defaults are `lost = error`, `warn = 100`, `norm = yes` for unit style of *lj*, `norm = no` for unit style of *real* and *metal*, `flush = no`, and `temp/press = compute IDs defined by thermo_style`.

The defaults for the line and format options depend on the thermo style. For styles “one” and “custom”, the line and format defaults are “one”, “%10d”, and “%14.8g”. For style “multi”, the line and format defaults are “multi”, “%14d”, and “%14.4f”. For style “yaml”, the line and format defaults are “%d” and “%.15g”.

## 16.129 thermo\_style command

### 16.129.1 Syntax

```
thermo_style style args
```

- `style = one` or `multi` or `yaml` or `custom`
- `args = list of arguments for a particular style`

`one args = none`

`multi args = none`

`yaml args = none`

`custom args = list of keywords`

```
possible keywords = step, elapsed, elaplong, dt, time,
                    cpu, tpcpu, spcpu, cpuremain, part, timeremain,
                    atoms, temp, press, pe, ke, etotal,
                    evdwl, ecoul, epair, ebond, eangle, edihed, eimp,
                    emol, elong, etail,
                    enthalpy, ecouple, econserve,
                    vol, density, lx, ly, lz, xlo, xhi, ylo, yhi, zlo, zhi,
                    xy, xz, yz, xlat, ylat, zlat,
                    bonds, angles, dihedrals, impropers,
                    pxx, pyy, pzz, pxy, pxz, pyz,
                    fmax, fnorm, nbuild, ndanger,
                    cella, cellb, cellc, cellalpha, cellbeta, cellgamma,
                    c_ID, c_ID[I], c_ID[I][J],
                    f_ID, f_ID[I], f_ID[I][J],
                    v_name, v_name[I]
```

`step = timestep`

`elapsed = timesteps since start of this run`

`elaplong = timesteps since start of initial run in a series of runs`

`dt = timestep size`

`time = simulation time`

```
cpu = elapsed CPU time in seconds since start of this run
tpcpu = time per CPU second
spcpu = timesteps per CPU second
cpuremain = estimated CPU time remaining in run
part = which partition (0 to Npartition-1) this is
timeremain = remaining time in seconds on timer timeout.
atoms = # of atoms
temp = temperature
press = pressure
pe = total potential energy
ke = kinetic energy
etotal = total energy (pe + ke)
evdwl = van der Waals pairwise energy (includes etail)
ecoul = Coulombic pairwise energy
epair = pairwise energy (evdwl + ecoul + elong)
ebond = bond energy
eangle = angle energy
ediهد = dihedral energy
eimp = improper energy
emol = molecular energy (ebond + eangle + ediهد + eimp)
elong = long-range kspace energy
etail = van der Waals energy long-range tail correction
enthalpy = enthalpy (etotal + press*vol)
ecouple = cumulative energy change due to thermo/baro statting fixes
econserve = pe + ke + ecouple = etotal + ecouple
vol = volume
density = mass density of system
lx,ly,lz = box lengths in x,y,z
xlo,xhi,ylo,yhi,zlo,zhi = box boundaries
xy,xz,yz = box tilt for triclinic (non-orthogonal) simulation boxes
xlat,ylat,zlat = lattice spacings as calculated by lattice command
bonds,angles,dihedrals,impropers = # of these interactions defined
pxx,pyy,pzz,pxy,pxz,pyz = 6 components of pressure tensor
fmax = max component of force on any atom in any dimension
fnorm = length of force vector for all atoms
nbuild = # of neighbor list builds
ndanger = # of dangerous neighbor list builds
cella,cellb,cellc = periodic cell lattice constants a,b,c
cellalpha, cellbeta, cellgamma = periodic cell angles alpha,beta,gamma
c_ID = global scalar value calculated by a compute with ID
c_ID[I] = Ith component of global vector calculated by a compute with ID, I can_
↪include wildcard (see below)
c_ID[I][J] = I,J component of global array calculated by a compute with ID
f_ID = global scalar value calculated by a fix with ID
f_ID[I] = Ith component of global vector calculated by a fix with ID, I can_
↪include wildcard (see below)
f_ID[I][J] = I,J component of global array calculated by a fix with ID
v_name = value calculated by an equal-style variable with name
v_name[I] = value calculated by a vector-style variable with name, I can_
↪include wildcard (see below)
```

## 16.129.2 Examples

```
thermo_style multi
thermo_style yaml
thermo_style one
thermo_style custom step temp pe etotal press vol
thermo_style custom step temp etotal c_myTemp v_abc
thermo_style custom step temp etotal c_myTemp[*] v_abc
```

## 16.129.3 Description

Set the style and content for printing thermodynamic data to the screen and log files.

Style *one* prints a single line of thermodynamic info that is the equivalent of “thermo\_style custom step temp epair emol etotal press”. The line contains only numeric values.

Style *multi* prints a multiple-line listing of thermodynamic info that is the equivalent of “thermo\_style custom etotal ke temp pe ebond eangle edihed eimp evdwl ecoul elong press”. The listing contains numeric values and a string ID for each quantity.

New in version 24Mar2022.

Style *yaml* is similar to style *one* but prints the output in [YAML](#) format which can be easily read by a variety of script languages and data handling packages. Since LAMMPS may print other output before, after, or in between thermodynamic output, the YAML format content needs to be separated from the rest. All YAML format thermodynamic output can be matched with a regular expression and can thus be extracted with commands like `egrep` as follows:

```
egrep '^(\bkeywords:|data:$|---$|\.\.\.\$| - \D)' log.lammps > log.yaml
```

Information about processing such YAML files is in the [structured data output howto](#).

Style *custom* is the most general setting and allows you to specify which of the keywords listed above you want printed on each thermodynamic timestep. Note that the keywords `c_ID`, `f_ID`, `v_name` are references to [computes](#), [fixes](#), and equal-style [variables](#) that have been defined elsewhere in the input script or can even be new styles which users have added to LAMMPS. See the [Modify](#) page for details on the latter. Thus the *custom* style provides a flexible means of outputting essentially any desired quantity as a simulation proceeds.

All styles except *custom* have *vol* appended to their list of outputs if the simulation box volume changes during the simulation.

The values printed by the various keywords are instantaneous values, calculated on the current timestep. Time-averaged quantities, which include values from previous timesteps, can be output by using the `f_ID` keyword and accessing a `fix` that does time-averaging such as the [fix ave/time](#) command.

Options invoked by the [thermo\\_modify](#) command can be used to set the one- or multi-line format of the print-out, the normalization of thermodynamic output (total values versus per-atom values for extensive quantities (ones which scale with the number of atoms in the system)), and the numeric precision of each printed value.

---

**Note:** When you use a “thermo\_style” command, all thermodynamic settings are restored to their default values, including those previously set by a [thermo\\_modify](#) command. Thus if your input script specifies a thermo\_style command, you should use the thermo\_modify command after it.

---

Several of the thermodynamic quantities require a temperature to be computed: “temp”, “press”, “ke”, “etotal”, “enthalpy”, “pxx”, etc. By default this is done by using a [temperature](#) compute which is created when LAMMPS starts up, as if this command had been issued:

```
compute thermo_temp all temp
```

See the `compute temp` command for details. Note that the ID of this compute is `thermo_temp` and the group is `all`. You can change the attributes of this temperature (e.g. its degrees-of-freedom) via the `compute_modify` command. Alternatively, you can directly assign a new compute (that calculates temperature) which you have defined, to be used for calculating any thermodynamic quantity that requires a temperature. This is done via the `thermo_modify` command.

Several of the thermodynamic quantities require a pressure to be computed: “press”, “enthalpy”, “pxx”, etc. By default this is done by using a `pressure` compute which is created when LAMMPS starts up, as if this command had been issued:

```
compute thermo_press all pressure thermo_temp
```

See the `compute pressure` command for details. Note that the ID of this compute is `thermo_press` and the group is `all`. You can change the attributes of this pressure via the `compute_modify` command. Alternatively, you can directly assign a new compute (that calculates pressure) which you have defined, to be used for calculating any thermodynamic quantity that requires a pressure. This is done via the `thermo_modify` command.

Several of the thermodynamic quantities require a potential energy to be computed: “pe”, “etotal”, “ebond”, etc. This is done by using a `pe` compute which is created when LAMMPS starts up, as if this command had been issued:

```
compute thermo_pe all pe
```

See the `compute pe` command for details. Note that the ID of this compute is `thermo_pe` and the group is `all`. You can change the attributes of this potential energy via the `compute_modify` command.

---

The kinetic energy of the system `ke` is inferred from the temperature of the system with  $\frac{1}{2}k_B T$  of energy for each degree of freedom. Thus, using different `compute commands` for calculating temperature, via the `thermo_modify temp` command, may yield different kinetic energies, since different computes that calculate temperature can subtract out different non-thermal components of velocity and/or include different degrees of freedom (translational, rotational, etc).

The potential energy of the system `pe` will include contributions from fixes if the `fix_modify energy yes` option is set for a fix that calculates such a contribution. For example, the `fix wall/lj93` fix calculates the energy of atoms interacting with the wall. See the doc pages for “individual fixes” to see which ones contribute and whether their default `fix_modify energy` setting is `yes` or `no`.

A long-range tail correction `etail` for the van der Waals pairwise energy will be non-zero only if the `pair_modify tail` option is turned on. The `etail` contribution is included in `evdwl`, `epair`, `pe`, and `etotal`, and the corresponding tail correction to the pressure is included in `press` and `pxx`, `pyy`, etc.

---

Here is more information on other keywords whose meaning may not be clear:

The `step`, `elapsed`, and `elaplong` keywords refer to timestep count. `Step` is the current timestep, or iteration count when a `minimization` is being performed. `Elapsed` is the number of timesteps elapsed since the beginning of this run. `Elaplong` is the number of timesteps elapsed since the beginning of an initial run in a series of runs. See the `start` and `stop` keywords for the `run` for info on how to invoke a series of runs that keep track of an initial starting time. If these keywords are not used, then `elapsed` and `elaplong` are the same value.

The `dt` keyword is the current timestep size in time `units`. The `time` keyword is the current elapsed simulation time, also in time `units`, which is simply (step\*dt) if the timestep size has not changed and the timestep has not been reset. If the timestep has changed (e.g. via `fix dt/reset`) or the timestep has been reset (e.g. via the “reset\_timestep” command), then the simulation time is effectively a cumulative value up to the current point.

The `cpu` keyword is elapsed CPU seconds since the beginning of this run. The `tpcpu` and `spcpu` keywords are measures of how fast your simulation is currently running. The `tpcpu` keyword is simulation time per CPU second, where simulation

time is in time *units*. E.g. for metal units, the *tpcpu* value would be picoseconds per CPU second. The *spcpu* keyword is the number of timesteps per CPU second. Both quantities are on-the-fly metrics, measured relative to the last time they were invoked. Thus if you are printing out thermodynamic output every 100 timesteps, the two keywords will continually output the time and timestep rate for the last 100 steps. The *tpcpu* keyword does not attempt to track any changes in timestep size, e.g. due to using the *fix dt/reset* command.

The *cpuremain* keyword estimates the CPU time remaining in the current run, based on the time elapsed thus far. It will only be a good estimate if the CPU time/timestep for the rest of the run is similar to the preceding timesteps. On the initial timestep the value will be 0.0 since there is no history to estimate from. For a minimization run performed by the “minimize” command, the estimate is based on the *maxiter* parameter, assuming the minimization will proceed for the maximum number of allowed iterations.

The *part* keyword is useful for multi-replica or multi-partition simulations to indicate which partition this output and this file corresponds to, or for use in a *variable* to append to a filename for output specific to this partition. See discussion of the *-partition command-line switch* for details on running in multi-partition mode.

The *timeremain* keyword is the seconds remaining when a timeout has been configured via the *timer timeout* command. If the timeout timer is inactive, the value of this keyword is 0.0 and if the timer is expired, it is negative. This allows for example to exit loops cleanly, if the timeout is expired with:

```
if "${timeremain} < 0.0" then "quit 0"
```

The *ecouple* keyword is cumulative energy change in the system due to any thermostating or barostating fixes that are being used. A positive value means that energy has been subtracted from the system (added to the coupling reservoir). See the *econserve* keyword for an explanation of why this sign choice makes sense.

The *econserve* keyword is the sum of the potential and kinetic energy of the system as well as the energy that has been transferred by thermostating or barostating to their coupling reservoirs. I.e. it is  $pe + ke + econserve$ . Ideally, for a simulation in the NVE, NPH, or NPT ensembles, the *econserve* quantity should remain constant over time.

The *fmax* and *fnorm* keywords are useful for monitoring the progress of an *energy minimization*. The *fmax* keyword calculates the maximum force in any dimension on any atom in the system, or the infinity-norm of the force vector for the system. The *fnorm* keyword calculates the 2-norm or length of the force vector.

The *nbuild* and *ndanger* keywords are useful for monitoring neighbor list builds during a run. Note that both these values are also printed with the end-of-run statistics. The *nbuild* keyword is the number of re-builds during the current run. The *ndanger* keyword is the number of re-builds that LAMMPS considered potentially “dangerous”. If atom movement triggered neighbor list rebuilding (see the *neigh\_modify* command), then dangerous reneighborings are those that were triggered on the first timestep atom movement was checked for. If this count is non-zero you may wish to reduce the delay factor to insure no force interactions are missed by atoms moving beyond the neighbor skin distance before a rebuild takes place.

The keywords *cella*, *cellb*, *cellc*, *cellalpha*, *cellbeta*, *cellgamma*, correspond to the usual crystallographic quantities that define the periodic unit cell of a crystal. See the *Howto triclinic* page for a geometric description of triclinic periodic cells, including a precise definition of these quantities in terms of the internal LAMMPS cell dimensions  $lx$ ,  $ly$ ,  $lz$ ,  $yz$ ,  $xz$ ,  $xy$ .

For output values from a compute or fix or variable, the bracketed index I used to index a vector, as in *c\_ID[I]* or *f\_ID[I]* or *v\_name[I]*, can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If N = the size of the vector, then an asterisk with no numeric values means all indices from 1 to N. A leading asterisk means all indices from 1 to n (inclusive). A trailing asterisk means all indices from n to N (inclusive). A middle asterisk means all indices from m to n (inclusive).

Using a wildcard is the same as if the individual elements of the vector had been listed one by one. E.g. these 2 thermo\_style commands are equivalent, since the *compute temp* command creates a global vector with 6 values.

```
compute myTemp all temp
thermo_style custom step temp etotal c_myTemp[*]
thermo_style custom step temp etotal &
                c_myTemp[1] c_myTemp[2] c_myTemp[3] &
                c_myTemp[4] c_myTemp[5] c_myTemp[6]
```

---

**Note:** For a vector-style variable, only the wildcard forms “\*n” or “m\*n” are allowed. You must specify the upper bound, because vector-style variable lengths are not determined until the variable is evaluated. If n is specified larger than the vector length turns out to be, zeroes are output for missing vector values.

---

The *c\_ID* and *c\_ID[I]* and *c\_ID[I][J]* keywords allow global values calculated by a compute to be output. As discussed on the *compute* doc page, computes can calculate global, per-atom, or local values. Only global values can be referenced by this command. However, per-atom compute values for an individual atom can be referenced in a *variable* and the variable referenced by *thermo\_style* custom, as discussed below. See the discussion above for how the I in *c\_ID[I]* can be specified with a wildcard asterisk to effectively specify multiple values from a global compute vector.

The ID in the keyword should be replaced by the actual ID of a compute that has been defined elsewhere in the input script. See the *compute* command for details. If the compute calculates a global scalar, vector, or array, then the keyword formats with 0, 1, or 2 brackets will reference a scalar value from the compute.

Note that some computes calculate “intensive” global quantities like temperature; others calculate “extensive” global quantities like kinetic energy that are summed over all atoms in the compute group. Intensive quantities are printed directly without normalization by *thermo\_style* custom. Extensive quantities may be normalized by the total number of atoms in the simulation (NOT the number of atoms in the compute group) when output, depending on the *thermo\_modify norm* option being used.

The *f\_ID* and *f\_ID[I]* and *f\_ID[I][J]* keywords allow global values calculated by a fix to be output. As discussed on the *fix* doc page, fixes can calculate global, per-atom, or local values. Only global values can be referenced by this command. However, per-atom fix values can be referenced for an individual atom in a *variable* and the variable referenced by *thermo\_style* custom, as discussed below. See the discussion above for how the I in *f\_ID[I]* can be specified with a wildcard asterisk to effectively specify multiple values from a global fix vector.

The ID in the keyword should be replaced by the actual ID of a fix that has been defined elsewhere in the input script. See the *fix* command for details. If the fix calculates a global scalar, vector, or array, then the keyword formats with 0, 1, or 2 brackets will reference a scalar value from the fix.

Note that some fixes calculate “intensive” global quantities like timestep size; others calculate “extensive” global quantities like energy that are summed over all atoms in the fix group. Intensive quantities are printed directly without normalization by *thermo\_style* custom. Extensive quantities may be normalized by the total number of atoms in the simulation (NOT the number of atoms in the fix group) when output, depending on the *thermo\_modify norm* option being used.

The *v\_name* keyword allow the current value of a variable to be output. The name in the keyword should be replaced by the variable name that has been defined elsewhere in the input script. Only equal-style and vector-style variables can be referenced; the latter requires a bracketed term to specify the Ith element of the vector calculated by the variable. However, an atom-style variable can be referenced for an individual atom by an equal-style variable and that variable referenced. See the *variable* command for details. Variables of style *equal* and *vector* and *atom* define a formula which can reference per-atom properties or thermodynamic keywords, or they can invoke other computes, fixes, or variables when evaluated, so this is a very general means of creating thermodynamic output.

Note that equal-style and vector-style variables are assumed to produce “intensive” global quantities, which are thus printed as-is, without normalization by *thermo\_style* custom. You can include a division by “natoms” in the variable formula if this is not the case.

---



## 16.129.4 Restrictions

This command must come after the simulation box is defined by a *read\_data*, *read\_restart*, or *create\_box* command.

## 16.129.5 Related commands

*thermo*, *thermo\_modify*, *fix\_modify*, *compute temp*, *compute pressure*

## 16.129.6 Default

```
thermo_style one
```

## 16.130 third\_order command

Accelerator Variants: *third\_order/kk*

### 16.130.1 Syntax

```
third_order group-ID style delta args keyword value ...
```

- group-ID = ID of group of atoms to displace
- style = *regular* or *eskm*
- delta = finite different displacement length (distance units)
- one or more keyword/arg pairs may be appended

keyword = *file* or *binary*

*file* name = name of output file for the third order tensor

*binary* arg = *yes* or *no* or *gzip*

### 16.130.2 Examples

```
third_order 1 regular 0.000001
third_order 1 eskm 0.000001
third_order 3 regular 0.00004 file third_order.dat
third_order 5 eskm 0.00000001 file third_order.dat binary yes
```

### 16.130.3 Description

Calculate the third order force constant tensor by finite difference of the selected group,

$$\Phi_{ijk}^{\alpha\beta\gamma} = \frac{\partial^3 U}{\partial x_{i,\alpha} \partial x_{j,\beta} \partial x_{k,\gamma}}$$

where Phi is the third order force constant tensor.

The output of the command is the tensor, three elements at a time. The three elements correspond to the three gamma elements for a specific *i/alpha/j/beta/k*. The initial five numbers are *i*, *alpha*, *j*, *beta*, and *k* respectively.

If the style *eskm* is selected, the tensor will be using energy units of 10 J/mol. These units conform to *eskm* style from the *dynamical\_matrix* command, which will simplify operations using dynamical matrices with third order tensors.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 16.130.4 Restrictions

The command collects a 9 times the number of atoms in the group on every single MPI rank, so the memory requirements can be very significant for large systems.

This command is part of the PHONON package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 16.130.5 Related commands

*fix phonon dynamical\_matrix*

### 16.130.6 Default

The default settings are *file* = "third\_order.dat", *binary* = no

## 16.131 timer command

### 16.131.1 Syntax

```
timer args
```

- *args* = one or more of *off* or *loop* or *normal* or *full* or *sync* or *nosync* or *timeout* or *every*

*off* = do not collect or print any timing information

*loop* = collect only the total time for the simulation loop

*normal* = collect timer information broken down by sections (default)

*full* = like *normal* but also include CPU and thread utilization

*sync* = explicitly synchronize MPI tasks between sections

*nosync* = do not synchronize MPI tasks between sections (default)

*timeout* *elapse* = set wall time limit to *elapse*

*every* *Ncheck* = perform timeout check every *Ncheck* steps

### 16.131.2 Examples

```
timer full sync
timer timeout 2:00:00 every 100
timer loop
```

### 16.131.3 Description

Select the level of detail at which LAMMPS performs its CPU timings. Multiple keywords can be specified with the *timer* command. For keywords that are mutually exclusive, the last one specified takes precedence.

During a simulation run LAMMPS collects information about how much time is spent in different sections of the code and thus can provide information for determining performance and load imbalance problems. This can be done at different levels of detail and accuracy. For more information about the timing output, see the [Run output](#) doc page.

The *off* setting will turn all time measurements off. The *loop* setting will only measure the total time for a run and not collect any detailed per section information. With the *normal* setting, timing information for portions of the timestep (pairwise calculations, neighbor list construction, output, etc) are collected as well as information about load imbalances for those sections across processors. The *full* setting adds information about CPU utilization and thread utilization, when multi-threading is enabled.

With the *sync* setting, all MPI tasks are synchronized at each timer call which measures load imbalance for each section more accurately, though it can also slow down the simulation by prohibiting overlapping independent computations on different MPI ranks. Using the *nosync* setting (which is the default) turns this synchronization off.

With the *timeout* keyword a wall time limit can be imposed, that affects the *run* and *minimize* commands. This can be convenient when calculations have to comply with execution time limits, e.g. when running under a batch system when you want to maximize the utilization of the batch time slot, especially for runs where the time per timestep varies much and thus it becomes difficult to predict how many steps a simulation can perform for a given wall time limit. This also applies for difficult to converge minimizations. The *timeout* *elapse* value should be somewhat smaller than the maximum wall time requested from the batch system, as there is usually some overhead to launch jobs, and it is advisable to write out a restart after terminating a run due to a timeout.

The *timeout* timer starts when the command is issued. When the time limit is reached, the run or energy minimization will exit on the next step or iteration that is a multiple of the *Ncheck* value which can be set with the *every* keyword. Default is checking every 10 steps. After the timer timeout has expired all subsequent run or minimize commands in the

input script will be skipped. The remaining time or timer status can be accessed with the *thermo* variable *timeremain*, which will be zero, if the timeout is inactive (default setting), it will be negative, if the timeout time is expired and positive if there is time remaining and in this case the value of the variable are the number of seconds remaining.

When the *timeout* key word is used a second time, the timer is restarted with a new time limit. The timeout *elapse* value can be specified as *off* or *unlimited* to impose a no timeout condition (which is the default). The *elapse* setting can be specified as a single number for seconds, two numbers separated by a colon (MM:SS) for minutes and seconds, or as three numbers separated by colons for hours, minutes, and seconds (H:MM:SS).

The *every* keyword sets how frequently during a run or energy minimization the wall clock will be checked. This check count applies to the outer iterations or time steps during minimizations or *r-RESPA runs*, respectively. Checking for timeout too often, can slow a calculation down. Checking too infrequently can make the timeout measurement less accurate, with the run being stopped later than desired.

---

**Note:** Using the *full* and *sync* options provides the most detailed and accurate timing information, but can also have a negative performance impact due to the overhead of the many required system calls. It is thus recommended to use these settings only when testing tests to identify performance bottlenecks. For calculations with few atoms or a very large number of processors, even the *normal* setting can have a measurable negative performance impact. In those cases you can just use the *loop* or *off* setting.

---

### 16.131.4 Restrictions

none

### 16.131.5 Related commands

*run post no, kspace\_modify fftbench*

### 16.131.6 Default

```
timer normal nosync
timer timeout off
timer every 10
```

## 16.132 timestep command

### 16.132.1 Syntax

```
timestep dt
```

- dt = timestep size (time units)

## 16.132.2 Examples

```
timestep 2.0
timestep 0.003
```

## 16.132.3 Description

Set the timestep size for subsequent molecular dynamics simulations. See the *units* command for the time units associated with each choice of units that LAMMPS supports.

The default value for the timestep size also depends on the choice of units for the simulation; see the default values below.

When the *run style* is *respa*, dt is the timestep for the outer loop (largest) timestep.

## 16.132.4 Restrictions

none

## 16.132.5 Related commands

*fix dt/reset*, *run*, *run\_style respa*, *units*

## 16.132.6 Default

choice of <i>units</i>	time units	default timestep size
lj	$\tau$	0.005 $\tau$
real	fs	1.0 fs
metal	ps	0.001 ps
si	s	1.0e-8 s (10 ns)
cgs	s	1.0e-8 s (10 ns)
electron	fs	0.001 fs
micro	$\mu$ s	2.0 $\mu$ s
nano	ns	0.00045 ns

## 16.133 uncompute command

### 16.133.1 Syntax

```
uncompute compute-ID
```

- compute-ID = ID of a previously defined compute

## 16.133.2 Examples

```
uncompute 2
uncompute lower-boundary
```

## 16.133.3 Description

Delete a compute that was previously defined with a *compute* command. This also wipes out any additional changes made to the compute via the *compute\_modify* command.

## 16.133.4 Restrictions

none

## 16.133.5 Related commands

*compute*

## 16.133.6 Default

none

# 16.134 undump command

## 16.134.1 Syntax

```
undump dump-ID
```

- dump-ID = ID of previously defined dump

## 16.134.2 Examples

```
undump mine
undump 2
```

## 16.134.3 Description

Turn off a previously defined dump so that it is no longer active. This closes the file associated with the dump.

## 16.134.4 Restrictions

none

## 16.134.5 Related commands

*dump*

## 16.134.6 Default

none

# 16.135 unfix command

## 16.135.1 Syntax

```
unfix fix-ID
```

- fix-ID = ID of a previously defined fix

## 16.135.2 Examples

```
unfix 2  
unfix lower-boundary
```

## 16.135.3 Description

Delete a fix that was previously defined with a *fix* command. This also wipes out any additional changes made to the fix via the *fix\_modify* command.

## 16.135.4 Restrictions

none

## 16.135.5 Related commands

*fix*

## 16.135.6 Default

none

## 16.136 units command

### 16.136.1 Syntax

```
units style
```

- style = *lj* or *real* or *metal* or *si* or *cgs* or *electron* or *micro* or *nano*

### 16.136.2 Examples

```
units metal  
units lj
```

### 16.136.3 Description

This command sets the style of units used for a simulation. It determines the units of all quantities specified in the input script and data file, as well as quantities output to the screen, log file, and dump files. Typically, this command is used at the very beginning of an input script.

For all units except *lj*, LAMMPS uses physical constants from [www.physics.nist.gov](http://www.physics.nist.gov). For the definition of Kcal in real units, LAMMPS uses the thermochemical calorie = 4.184 J.

The choice you make for units simply sets some internal conversion factors within LAMMPS. This means that any simulation you perform for one choice of units can be duplicated with any other unit setting LAMMPS supports. In this context “duplicate” means the particles will have identical trajectories and all output generated by the simulation will be identical. This will be the case for some number of timesteps until round-off effects accumulate, since the conversion factors for two different unit systems are not identical to infinite precision.

To perform the same simulation in a different set of units you must change all the unit-based input parameters in your input script and other input files (data file, potential files, etc) correctly to the new units. And you must correctly convert all output from the new units to the old units when comparing to the original results. That is often not simple to do.

Potential or table files may have a `UNITS:` tag included in the first line indicating the unit style those files were created for. If the tag exists, its value will be compared to the chosen unit style and LAMMPS will stop with an error message if there is a mismatch. In some select cases and for specific combinations of unit styles, LAMMPS is capable of automatically converting potential parameters from a file. In those cases, a warning message signaling that an automatic conversion has happened is printed to the screen.

---

For style *lj*, all quantities are unitless. Without loss of generality, LAMMPS sets the fundamental quantities mass,  $\sigma$ ,  $\epsilon$ , and the Boltzmann constant  $k_B = 1$ . The masses, distances, energies you specify are multiples of these fundamental values. The formulas relating the reduced or unitless quantity (with an asterisk) to the same quantity with units is also given. Thus you can use the mass,  $\sigma$ , and  $\epsilon$  values for a specific material and convert the results from a unitless LJ simulation into physical quantities. Please note that using these three properties as base, your unit of time has to conform to the relation  $\epsilon = \frac{m\sigma^2}{\tau^2}$  since energy is a derived unit (in SI units you equivalently have the relation  $1\text{J} = 1 \frac{\text{kg}\cdot\text{m}^2}{\text{s}^2}$ ).

- mass = mass or  $m$ , where  $M^* = \frac{M}{m}$



- distance =  $\sigma$ , where  $x^* = \frac{x}{\sigma}$
- time =  $\tau$ , where  $\tau^* = \tau \sqrt{\frac{\epsilon}{m\sigma^2}}$
- energy =  $\epsilon$ , where  $E^* = \frac{E}{\epsilon}$
- velocity =  $\frac{\sigma}{\tau}$ , where  $v^* = v \frac{\tau}{\sigma}$
- force =  $\frac{\epsilon}{\sigma}$ , where  $f^* = f \frac{\sigma}{\epsilon}$
- torque =  $\epsilon$ , where  $t^* = \frac{t}{\epsilon}$
- temperature = reduced LJ temperature, where  $T^* = \frac{Tk_B}{\epsilon}$
- pressure = reduced LJ pressure, where  $p^* = p \frac{\sigma^3}{\epsilon}$
- dynamic viscosity = reduced LJ viscosity, where  $\eta^* = \eta \frac{\sigma^3}{\epsilon\tau}$
- charge = reduced LJ charge, where  $q^* = q \frac{1}{\sqrt{4\pi\epsilon_0\sigma\epsilon}}$
- dipole = reduced LJ dipole, moment where  $\mu^* = \mu \frac{1}{\sqrt{4\pi\epsilon_0\sigma^3\epsilon}}$
- electric field = force/charge, where  $E^* = E \frac{\sqrt{4\pi\epsilon_0\sigma\epsilon}}{\epsilon}$
- density = mass/volume, where  $\rho^* = \rho \frac{\sigma^{dim}}{m}$

Note that for LJ units, the default mode of thermodynamic output via the *thermo\_style* command is to normalize all extensive quantities by the number of atoms. E.g. potential energy is extensive because it is summed over atoms, so it is output as energy/atom. Temperature is intensive since it is already normalized by the number of atoms, so it is output as-is. This behavior can be changed via the *thermo\_modify norm* command.

For style *real*, these are the units:

- mass = grams/mole
- distance = Angstroms
- time = femtoseconds
- energy = Kcal/mole
- velocity = Angstroms/femtosecond
- force = Kcal/mole-Angstrom
- torque = Kcal/mole
- temperature = Kelvin
- pressure = atmospheres
- dynamic viscosity = Poise
- charge = multiple of electron charge (1.0 is a proton)
- dipole = charge\*Angstroms
- electric field = volts/Angstrom
- density = gram/cm<sup>dim</sup>

For style *metal*, these are the units:

- mass = grams/mole
- distance = Angstroms
- time = picoseconds

- energy = eV
- velocity = Angstroms/picosecond
- force = eV/Angstrom
- torque = eV
- temperature = Kelvin
- pressure = bars
- dynamic viscosity = Poise
- charge = multiple of electron charge (1.0 is a proton)
- dipole = charge\*Angstroms
- electric field = volts/Angstrom
- density = gram/cm<sup>dim</sup>

For style *si*, these are the units:

- mass = kilograms
- distance = meters
- time = seconds
- energy = Joules
- velocity = meters/second
- force = Newtons
- torque = Newton-meters
- temperature = Kelvin
- pressure = Pascals
- dynamic viscosity = Pascal\*second
- charge = Coulombs (1.6021765e-19 is a proton)
- dipole = Coulombs\*meters
- electric field = volts/meter
- density = kilograms/meter<sup>dim</sup>

For style *cgs*, these are the units:

- mass = grams
- distance = centimeters
- time = seconds
- energy = ergs
- velocity = centimeters/second
- force = dynes
- torque = dyne-centimeters
- temperature = Kelvin
- pressure = dyne/cm<sup>2</sup> or barye = 1.0e-6 bars

- dynamic viscosity = Poise
- charge = statcoulombs or esu (4.8032044e-10 is a proton)
- dipole = statcoul-cm =  $10^{18}$  debye
- electric field = statvolt/cm or dyne/esu
- density = grams/cm<sup>dim</sup>

For style *electron*, these are the units:

- mass = atomic mass units
- distance = Bohr
- time = femtoseconds
- energy = Hartrees
- velocity = Bohr/atomic time units [1.03275e-15 seconds]
- force = Hartrees/Bohr
- temperature = Kelvin
- pressure = Pascals
- charge = multiple of electron charge (1.0 is a proton)
- dipole moment = Debye
- electric field = volts/cm

For style *micro*, these are the units:

- mass = picograms
- distance = micrometers
- time = microseconds
- energy = picogram-micrometer<sup>2</sup>/microsecond<sup>2</sup>
- velocity = micrometers/microsecond
- force = picogram-micrometer/microsecond<sup>2</sup>
- torque = picogram-micrometer<sup>2</sup>/microsecond<sup>2</sup>
- temperature = Kelvin
- pressure = picogram/(micrometer-microsecond<sup>2</sup>)
- dynamic viscosity = picogram/(micrometer-microsecond)
- charge = picocoulombs (1.6021765e-7 is a proton)
- dipole = picocoulomb-micrometer
- electric field = volt/micrometer
- density = picograms/micrometer<sup>dim</sup>

For style *nano*, these are the units:

- mass = attograms
- distance = nanometers
- time = nanoseconds

- energy = attogram-nanometer<sup>2</sup>/nanosecond<sup>2</sup>
- velocity = nanometers/nanosecond
- force = attogram-nanometer/nanosecond<sup>2</sup>
- torque = attogram-nanometer<sup>2</sup>/nanosecond<sup>2</sup>
- temperature = Kelvin
- pressure = attogram/(nanometer-nanosecond<sup>2</sup>)
- dynamic viscosity = attogram/(nanometer-nanosecond)
- charge = multiple of electron charge (1.0 is a proton)
- dipole = charge-nanometer
- electric field = volt/nanometer
- density = attograms/nanometer<sup>dim</sup>

The units command also sets the timestep size and neighbor skin distance to default values for each style:

- For style *lj* these are  $dt = 0.005 \tau$  and  $skin = 0.3 \sigma$ .
- For style *real* these are  $dt = 1.0$  femtoseconds and  $skin = 2.0$  Angstroms.
- For style *metal* these are  $dt = 0.001$  picoseconds and  $skin = 2.0$  Angstroms.
- For style *si* these are  $dt = 1.0e-8$  seconds and  $skin = 0.001$  meters.
- For style *cgs* these are  $dt = 1.0e-8$  seconds and  $skin = 0.1$  centimeters.
- For style *electron* these are  $dt = 0.001$  femtoseconds and  $skin = 2.0$  Bohr.
- For style *micro* these are  $dt = 2.0$  microseconds and  $skin = 0.1$  micrometers.
- For style *nano* these are  $dt = 0.00045$  nanoseconds and  $skin = 0.1$  nanometers.

#### 16.136.4 Restrictions

This command cannot be used after the simulation box is defined by a *read\_data* or *create\_box* command.

#### 16.136.5 Related commands

none

#### 16.136.6 Default

```
units lj
```

## 16.137 variable command

### 16.137.1 Syntax

```
variable name style args ...
```

- name = name of variable to define
- style = *delete* or *index* or *loop* or *world* or *universe* or *uloop* or *string* or *format* or *getenv* or *file* or *atomfile* or *python* or *timer* or *internal* or *equal* or *vector* or *atom*

*delete* = no args

*index* args = one or more strings

*loop* args = N

N = integer size of loop, loop from 1 to N inclusive

*loop* args = N pad

N = integer size of loop, loop from 1 to N inclusive

pad = all values will be same length, e.g. 001, 002, ..., 100

*loop* args = N1 N2

N1,N2 = loop from N1 to N2 inclusive

*loop* args = N1 N2 pad

N1,N2 = loop from N1 to N2 inclusive

pad = all values will be same length, e.g. 050, 051, ..., 100

*world* args = one string for each partition of processors

*universe* args = one or more strings

*uloop* args = N

N = integer size of loop

*uloop* args = N pad

N = integer size of loop

pad = all values will be same length, e.g. 001, 002, ..., 100

*string* arg = one string

*format* args = vname fstr

vname = name of equal-style variable to evaluate

fstr = C-style format string

*getenv* arg = one string

*file* arg = filename

*atomfile* arg = filename

*python* arg = function

*timer* arg = no arguments

*internal* arg = numeric value

*equal* or *vector* or *atom* args = one formula containing numbers, thermo keywords, ↵

↵math operations, group functions, atom values and vectors, compute/fix/variable ↵

↵references

numbers = 0.0, 100, -5.4, 2.8e-4, etc

constants = PI, version, on, off, true, false, yes, no

thermo keywords = vol, ke, press, etc from *thermo\_style*

math operators = (), -x, x+y, x-y, x\*y, x/y, x^y, x%y,

x == y, x != y, x < y, x <= y, x > y, x >= y, x && y, x || y, x ↵

↵|^ y, !x

math functions = sqrt(x), exp(x), ln(x), log(x), abs(x),  
sin(x), cos(x), tan(x), asin(x), acos(x), atan(x), atan2(y,x),  
random(x,y,z), normal(x,y,z), ceil(x), floor(x), round(x)  
ramp(x,y), stagger(x,y), logfreq(x,y,z), logfreq2(x,y,z),  
logfreq3(x,y,z), stride(x,y,z), stride2(x,y,z,a,b,c),

```

        vdisplace(x,y), swiggle(x,y,z), cwiggle(x,y,z)
group functions = count(group), mass(group), charge(group),
                 xcm(group,dim), vcm(group,dim), fcm(group,dim),
                 bound(group,dir), gyration(group), ke(group),
                 angmom(group,dim), torque(group,dim),
                 inertia(group,dimdim), omega(group,dim)
region functions = count(group,region), mass(group,region), charge(group,region),
                 xcm(group,dim,region), vcm(group,dim,region), fcm(group,dim,
↪region),
                 bound(group,dir,region), gyration(group,region), ke(group,
↪reigon),
                 angmom(group,dim,region), torque(group,dim,region),
                 inertia(group,dimdim,region), omega(group,dim,region)
special functions = sum(x), min(x), max(x), ave(x), trap(x), slope(x), gmask(x), ↪
↪rmask(x), grmask(x,y), next(x), is_file(name), is_os(name), extract_setting(name),
↪ label2type(kind,label)
feature functions = is_active(category,feature), is_available(category,feature), ↪
↪is_defined(category,id)
atom value = id[i], mass[i], type[i], mol[i], x[i], y[i], z[i], vx[i], vy[i], ↪
↪vz[i], fx[i], fy[i], fz[i], q[i]
atom vector = id, mass, type, mol, x, y, z, vx, vy, vz, fx, fy, fz, q
compute references = c_ID, c_ID[i], c_ID[i][j], C_ID, C_ID[i]
fix references = f_ID, f_ID[i], f_ID[i][j], F_ID, F_ID[i]
variable references = v_name, v_name[i]

```

## 16.137.2 Examples

```

variable x index run1 run2 run3 run4 run5 run6 run7 run8
variable LoopVar loop $n
variable beta equal temp/3.0
variable b1 equal x[234]+0.5*vol
variable b1 equal "x[234] + 0.5*vol"
variable b equal xcm(mol1,x)/2.0
variable b equal c_myTemp
variable b atom x*y/vol
variable foo string myfile
variable foo internal 3.5
variable myPy python increase
variable f file values.txt
variable temp world 300.0 310.0 320.0 ${Tfinal}
variable x universe 1 2 3 4 5 6 7 8 9 10 11 12 13 14 15
variable x uloop 15 pad
variable str format x %.6g
variable x delete

```

```

variable start timer
other commands
variable stop timer
print "Elapsed time: $(v_stop-v_start:%.6f)"

```

### 16.137.3 Description

This command assigns one or more strings to a variable name for evaluation later in the input script or during a simulation.

Variables can thus be useful in several contexts. A variable can be defined and then referenced elsewhere in an input script to become part of a new input command. For variable styles that store multiple strings, the *next* command can be used to increment which string is assigned to the variable. Variables of style *equal* store a formula which when evaluated produces a single numeric value which can be output either directly (see the *print*, *fix print*, and *run every* commands) or as part of thermodynamic output (see the *thermo\_style* command), or used as input to an averaging fix (see the *fix ave/time* command). Variables of style *vector* store a formula which produces a vector of such values which can be used as input to various averaging fixes, or elements of which can be part of thermodynamic output. Variables of style *atom* store a formula which when evaluated produces one numeric value per atom which can be output to a dump file (see the *dump custom* command) or used as input to an averaging fix (see the *fix ave/chunk* and *fix ave/atom* commands). Variables of style *atomfile* can be used anywhere in an input script that atom-style variables are used; they get their per-atom values from a file rather than from a formula. Variables of style *python* can be hooked to Python functions using code you provide, so that the variable gets its value from the evaluation of the Python code. Variables of style *internal* are used by a few commands which set their value directly.

---

**Note:** As discussed on the *Commands parse* doc page, an input script can use “immediate” variables, specified as \$(formula) with parenthesis, where the numeric formula has the same syntax as equal-style variables described on this page. This is a convenient way to evaluate a formula immediately without using the variable command to define a named variable and then evaluate that variable. The formula can include a trailing colon and format string which determines the precision with which the numeric value is generated. This is also explained on the *Commands parse* doc page.

---

In the discussion that follows, the “name” of the variable is the arbitrary string that is the first argument in the variable command. This name can only contain alphanumeric characters and underscores. The “string” is one or more of the subsequent arguments. The “string” can be simple text as in the first example above, it can contain other variables as in the second example, or it can be a formula as in the third example. The “value” is the numeric quantity resulting from evaluation of the string. Note that the same string can generate different values when it is evaluated at different times during a simulation.

---

**Note:** When an input script line is encountered that defines a variable of style *equal* or *vector* or *atom* or *python* that contains a formula or Python code, the formula is NOT immediately evaluated. It will be evaluated every time when the variable is **used** instead. If you simply want to evaluate a formula in place you can use as so-called. See the section below about “Immediate Evaluation of Variables” for more details on the topic. This is also true of a *format* style variable since it evaluates another variable when it is invoked.

---

Variables of style *equal* and *vector* and *atom* can be used as inputs to various other commands which evaluate their formulas as needed, e.g. at different timesteps during a *run*. In this context, variables of style *timer* or *internal* or *python* can be used in place of an equal-style variable, with the following two caveats.

First, internal-style variables can be used except by commands that set the value stored by the internal variable. When the LAMMPS command evaluates the internal-style variable, it will use the value set (internally) by another command. Second, python-style variables can be used so long as the associated Python function, as defined by the *python* command, returns a numeric value. When the LAMMPS command evaluates the python-style variable, the Python function will be executed.

---

**Note:** When a variable command is encountered in the input script and the variable name has already been specified, the command is ignored. This means variables can NOT be re-defined in an input script (with two exceptions, read further). This is to allow an input script to be processed multiple times without resetting the variables; see the *jump* or

*include* commands. It also means that using the *command-line switch* `-var` will override a corresponding index variable setting in the input script.

---

There are two exceptions to this rule. First, variables of style *string*, *getenv*, *internal*, *equal*, *vector*, *atom*, and *python* ARE redefined each time the command is encountered. This allows these style of variables to be redefined multiple times in an input script. In a loop, this means the formula associated with an *equal* or *atom* style variable can change if it contains a substitution for another variable, e.g. `$x` or `v_x`.

Second, as described below, if a variable is iterated on to the end of its list of strings via the *next* command, it is removed from the list of active variables, and is thus available to be re-defined in a subsequent variable command. The *delete* style does the same thing.

---

The *Commands parse* page explains how occurrences of a variable name in an input script line are replaced by the variable's string. The variable name can be referenced as `$x` if the name "x" is a single character, or as `${LoopVar}` if the name "LoopVar" is one or more characters.

As described below, for variable styles *index*, *loop*, *file*, *universe*, and *uloop*, which string is assigned to a variable can be incremented via the *next* command. When there are no more strings to assign, the variable is exhausted and a flag is set that causes the next *jump* command encountered in the input script to be skipped. This enables the construction of simple loops in the input script that are iterated over and then exited from.

As explained above, an exhausted variable can be re-used in an input script. The *delete* style also removes the variable, the same as if it were exhausted, allowing it to be redefined later in the input script or when the input script is looped over. This can be useful when breaking out of a loop via the *if* and *jump* commands before the variable would become exhausted. For example,

```
label      loop
variable   a loop 5
print      "A = $a"
if         "$a > 2" then "jump in.script break"
next       a
jump       in.script loop
label      break
variable   a delete
```

---

This section describes how all the various variable styles are defined and what they store. Except for the *equal* and *vector* and *atom* styles, which are explained in the next section.

Many of the styles store one or more strings. Note that a single string can contain spaces (multiple words), if it is enclosed in quotes in the variable command. When the variable is substituted for in another input script command, its returned string will then be interpreted as multiple arguments in the expanded command.

For the *index* style, one or more strings are specified. Initially, the first string is assigned to the variable. Each time a *next* command is used with the variable name, the next string is assigned. All processors assign the same string to the variable.

*Index* style variables with a single string value can also be set by using the *command-line switch* `-var`.

The *loop* style is identical to the *index* style except that the strings are the integers from 1 to N inclusive, if only one argument N is specified. This allows generation of a long list of runs (e.g. 1000) without having to list N strings in the input script. Initially, the string "1" is assigned to the variable. Each time a *next* command is used with the variable name, the next string ("2", "3", etc) is assigned. All processors assign the same string to the variable. The *loop* style can also be specified with two arguments N1 and N2. In this case the loop runs from N1 to N2 inclusive, and the string N1 is initially assigned to the variable.  $N1 \leq N2$  and  $N2 \geq 0$  is required.



For the *world* style, one or more strings are specified. There must be one string for each processor partition or “world”. LAMMPS can be run with multiple partitions via the *-partition command-line switch*. This variable command assigns one string to each world. All processors in the world are assigned the same string. The next command cannot be used with *equal* style variables, since there is only one value per world. This style of variable is useful when you wish to run different simulations on different partitions, or when performing a parallel tempering simulation (see the *temper* command), to assign different temperatures to different partitions.

For the *universe* style, one or more strings are specified. There must be at least as many strings as there are processor partitions or “worlds”. LAMMPS can be run with multiple partitions via the *-partition command-line switch*. This variable command initially assigns one string to each world. When a *next* command is encountered using this variable, the first processor partition to encounter it, is assigned the next available string. This continues until all the variable strings are consumed. Thus, this command can be used to run 50 simulations on 8 processor partitions. The simulations will be run one after the other on whatever partition becomes available, until they are all finished. *Universe* style variables are incremented using the files “tmp.lammps.variable” and “tmp.lammps.variable.lock” which you will see in your directory during such a LAMMPS run.

The *uloop* style is identical to the *universe* style except that the strings are the integers from 1 to N. This allows generation of long list of runs (e.g. 1000) without having to list N strings in the input script.

For the *string* style, a single string is assigned to the variable. Two differences between this style and using the *index* style exist: a variable with *string* style can be redefined, e.g. by another command later in the input script, or if the script is read again in a loop. The other difference is that *string* performs variable substitution even if the string parameter is quoted.

For the *format* style, an equal-style variable is specified along with a C-style format string, e.g. “%f” or “%.10g”, which must be appropriate for formatting a double-precision floating-point value. The default format is “%.15g”. This variable style allows an equal-style variable to be formatted precisely when it is evaluated.

Note that if you simply wish to print a variable value with desired precision to the screen or logfile via the *print* or *fix print* commands, you can also do this by specifying an “immediate” variable with a trailing colon and format string, as part of the string argument of those commands. This is explained on the *Commands parse* doc page.

For the *getenv* style, a single string is assigned to the variable which should be the name of an environment variable. When the variable is evaluated, it returns the value of the environment variable, or an empty string if it not defined. This style of variable can be used to adapt the behavior of LAMMPS input scripts via environment variable settings, or to retrieve information that has been previously stored with the *shell putenv* command. Note that because environment variable settings are stored by the operating systems, they persist beyond a *clear* command.

For the *file* style, a filename is provided which contains a list of strings to assign to the variable, one per line. The strings can be numeric values if desired. See the discussion of the *next()* function below for equal-style variables, which will convert the string of a file-style variable into a numeric value in a formula.

When a file-style variable is defined, the file is opened and the string on the first line is read and stored with the variable. This means the variable can then be evaluated as many times as desired and will return that string. There are two ways to cause the next string from the file to be read: use the *next* command or the *next()* function in an equal- or atom-style variable, as discussed below.

The rules for formatting the file are as follows. A comment character “#” can be used anywhere on a line; text starting with the comment character is stripped. Blank lines are skipped. The first “word” of a non-blank line, delimited by white-space, is the “string” assigned to the variable.

For the *atomfile* style, a filename is provided which contains one or more sets of values, to assign on a per-atom basis to the variable. The format of the file is described below.

When an atomfile-style variable is defined, the file is opened and the first set of per-atom values are read and stored with the variable. This means the variable can then be evaluated as many times as desired and will return those values. There are two ways to cause the next set of per-atom values from the file to be read: use the *next* command or the *next()* function in an atom-style variable, as discussed below.

The rules for formatting the file are as follows. Each time a set of per-atom values is read, a non-blank line is searched for in the file. The file is read line by line but only up to 254 characters are used. The rest are ignored. A comment character “#” can be used anywhere on a line and all text following and the “#” character are ignored; text starting with the comment character is stripped. Blank lines are skipped. The first “word” of a non-blank line, delimited by white-space, is read as the count N of per-atom lines to immediately follow. N can be the total number of atoms in the system, or only a subset. The next N lines have the following format

```
ID value
```

where ID is an atom ID and value is the per-atom numeric value that will be assigned to that atom. IDs can be listed in any order.

---

**Note:** Every time a set of per-atom lines is read, the value for all atoms is first set to 0.0. Thus values for atoms whose ID does not appear in the set, will remain 0.0.

---

For the *python* style a Python function name is provided. This needs to match a function name specified in a *python* command which returns a value to this variable as defined by its *return* keyword. For example these two commands would be self-consistent:

```
variable foo python myMultiply
python myMultiply return v_foo format f file funcs.py
```

The two commands can appear in either order so long as both are specified before the Python function is invoked for the first time.

Each time the variable is evaluated, the associated Python function is invoked, and the value it returns is also returned by the variable. Since the Python function can use other LAMMPS variables as input, or query internal LAMMPS quantities to perform its computation, this means the variable can return a different value each time it is evaluated.

The type of value stored in the variable is determined by the *format* keyword of the *python* command. It can be an integer (i), floating point (f), or string (s) value. As mentioned above, if it is a numeric value (integer or floating point), then the python-style variable can be used in place of an equal-style variable anywhere in an input script, e.g. as an argument to another command that allows for equal-style variables.

For the *timer* style no additional argument is specified. The value of the variable is set by querying the current elapsed wall time of the simulation. This is done at the point in time when the variable is defined in the input script. If a second timer-style variable is also defined, then a simple formula can be used to calculate the elapsed time between the two timers, as in the example at the top of this manual entry. As mentioned above, timer-style variables can be redefined elsewhere in the input script, so the same pair of variables can be used in a loop or to time a series of operations.

For the *internal* style a numeric value is provided. This value will be assigned to the variable until a LAMMPS command sets it to a new value. There are currently only two LAMMPS commands that require *internal* variables as inputs, because they reset them: *create\_atoms* and *fix controller*. As mentioned above, an internal-style variable can be used in place of an equal-style variable anywhere else in an input script, e.g. as an argument to another command that allows for equal-style variables.

---

For the *equal* and *vector* and *atom* styles, a single string is specified which represents a formula that will be evaluated afresh each time the variable is used. If you want spaces in the string, enclose it in double quotes so the parser will treat it as a single argument. For *equal*-style variables the formula computes a scalar quantity, which becomes the value of the variable whenever it is evaluated. For *vector*-style variables the formula must compute a vector of quantities, which becomes the value of the variable whenever it is evaluated. The calculated vector can be of length one, but it cannot be a simple scalar value like that produced by an equal-style compute. I.e. the formula for a vector-style variable must have at least one quantity in it that refers to a global vector produced by a compute, fix, or other vector-style variable. For *atom*-style variables the formula computes one quantity for each atom whenever it is evaluated.

Note that *equal*, *vector*, and *atom* variables can produce different values at different stages of the input script or at different times during a run. For example, if an *equal* variable is used in a *fix print* command, different values could be printed each timestep it was invoked. If you want a variable to be evaluated immediately, so that the result is stored by the variable instead of the string, see the section below on “Immediate Evaluation of Variables”.

The next command cannot be used with *equal* or *vector* or *atom* style variables, since there is only one string.

The formula for an *equal*, *vector*, or *atom* variable can contain a variety of quantities. The syntax for each kind of quantity is simple, but multiple quantities can be nested and combined in various ways to build up formulas of arbitrary complexity. For example, this is a valid (though strange) variable formula:

```
variable x equal "pe + c_MyTemp / vol^(1/3)"
```

Specifically, a formula can contain numbers, constants, thermo keywords, math operators, math functions, group functions, region functions, atom values, atom vectors, compute references, fix references, and references to other variables.

Number	0.2, 100, 1.0e20, -15.4, etc
Constant	PI, version, on, off, true, false, yes, no
Thermodynamic keywords	vol, pe, ebond, etc
Math operators	( ), -x, x+y, x-y, x*y, x/y, x^y, x%y, x == y, x != y, x < y, x <= y, x > y, x >= y, x && y, x    y, x  ^ y, !x
Math functions	sqrt(x), exp(x), ln(x), log(x), abs(x), sin(x), cos(x), tan(x), asin(x), acos(x), atan(x), atan2(y,x), random(x,y,z), normal(x,y,z), ceil(x), floor(x), round(x), ramp(x,y), stagger(x,y), logfreq(x,y,z), logfreq2(x,y,z), logfreq3(x,y,z), stride(x,y,z), stride2(x,y,z,a,b,c), vdisplace(x,y), swiggle(x,y,z), cwiggle(x,y,z)
Group functions	count(ID), mass(ID), charge(ID), xcm(ID,dim), vcm(ID,dim), fcm(ID,dim), bound(ID,dir), gyration(ID), ke(ID), angmom(ID,dim), torque(ID,dim), inertia(ID,dimdim), omega(ID,dim)
Region functions	count(ID,IDR), mass(ID,IDR), charge(ID,IDR), xcm(ID,dim,IDR), vcm(ID,dim,IDR), fcm(ID,dim,IDR), bound(ID,dir,IDR), gyration(ID,IDR), ke(ID,IDR), angmom(ID,dim,IDR), torque(ID,dim,IDR), inertia(ID,dimdim,IDR), omega(ID,dim,IDR)
Special functions	sum(x), min(x), max(x), ave(x), trap(x), slope(x), gmask(x), rmask(x), grmask(x,y), next(x), label2type(kind,label)
Atom values	id[i], mass[i], type[i], mol[i], x[i], y[i], z[i], vx[i], vy[i], vz[i], fx[i], fy[i], fz[i], q[i]
Atom vectors	id, mass, type, mol, x, y, z, vx, vy, vz, fx, fy, fz, q
Compute references	c_ID, c_ID[i], c_ID[i][j], C_ID, C_ID[i]
Fix references	f_ID, f_ID[i], f_ID[i][j], F_ID, F_ID[i]
Other variables	v_name, v_name[i]

Most of the formula elements produce a scalar value. Some produce a global or per-atom vector of values. Global vectors can be produced by computes or fixes or by other vector-style variables. Per-atom vectors are produced by atom vectors, compute references that represent a per-atom vector, fix references that represent a per-atom vector, and variables that are atom-style variables. Math functions that operate on scalar values produce a scalar value; math function that operate on global or per-atom vectors do so element-by-element and produce a global or per-atom vector.

A formula for equal-style variables cannot use any formula element that produces a global or per-atom vector. A formula for a vector-style variable can use formula elements that produce either a scalar value or a global vector value, but cannot use a formula element that produces a per-atom vector. A formula for an atom-style variable can use formula elements that produce either a scalar value or a per-atom vector, but not one that produces a global vector. Atom-style variables are evaluated by other commands that define a *group* on which they operate, e.g. a *dump* or *compute* or *fix* command. When they invoke the atom-style variable, only atoms in the group are included in the formula evaluation.

The variable evaluates to 0.0 for atoms not in the group.

## Numbers, constants, and thermo keywords

Numbers can contain digits, scientific notation (3.0e20,3.0e-20,3.0E20,3.0E-20), and leading minus signs.

Constants are set at compile time and cannot be changed. *PI* will return the number 3.14159265358979323846; *on*, *true* or *yes* will return 1.0; *off*, *false* or *no* will return 0.0; *version* will return a numeric version code of the current LAMMPS version (e.g. version 2 Sep 2015 will return the number 20150902). The corresponding value for newer versions of LAMMPS will be larger, for older versions of LAMMPS will be smaller. This can be used to have input scripts adapt automatically to LAMMPS versions, when non-backwards compatible syntax changes are introduced. Here is an illustrative example (which will not work, since the *version* has been introduced more recently):

```
if $(version<20140513) then "communicate vel yes" else "comm_modify vel yes"
```

The thermo keywords allowed in a formula are those defined by the *thermo\_style custom* command. Thermo keywords that require a *compute* to calculate their values such as “temp” or “press”, use computes stored and invoked by the *thermo\_style* command. This means that you can only use those keywords in a variable if the style you are using with the *thermo\_style* command (and the thermo keywords associated with that style) also define and use the needed compute. Note that some thermo keywords use a compute indirectly to calculate their value (e.g. the enthalpy keyword uses temp, pe, and pressure). If a variable is evaluated directly in an input script (not during a run), then the values accessed by the thermo keyword must be current. See the discussion below about “Variable Accuracy”.

## Math Operators

Math operators are written in the usual way, where the “x” and “y” in the examples can themselves be arbitrarily complex formulas, as in the examples above. In this syntax, “x” and “y” can be scalar values or per-atom vectors. For example, “ke/natoms” is the division of two scalars, where “vy+vz” is the element-by-element sum of two per-atom vectors of y and z velocities.

Operators are evaluated left to right and have the usual C-style precedence: unary minus and unary logical NOT operator “!” have the highest precedence, exponentiation “^” is next; multiplication and division and the modulo operator “%” are next; addition and subtraction are next; the 4 relational operators “<”, “<=”, “>”, and “>=” are next; the two remaining relational operators “==” and “!=” are next; then the logical AND operator “&&”; and finally the logical OR operator “||” and logical XOR (exclusive or) operator “|^” have the lowest precedence. Parenthesis can be used to group one or more portions of a formula and/or enforce a different order of evaluation than what would occur with the default precedence.

**Note:** Because a unary minus is higher precedence than exponentiation, the formula “-2^2” will evaluate to 4, not -4. This convention is compatible with some programming languages, but not others. As mentioned, this behavior can be easily overridden with parenthesis; the formula “-(2^2)” will evaluate to -4.

The 6 relational operators return either a 1.0 or 0.0 depending on whether the relationship between x and y is TRUE or FALSE. For example the expression  $x < 10.0$  in an atom-style variable formula will return 1.0 for all atoms whose x-coordinate is less than 10.0, and 0.0 for the others. The logical AND operator will return 1.0 if both its arguments are non-zero, else it returns 0.0. The logical OR operator will return 1.0 if either of its arguments is non-zero, else it returns 0.0. The logical XOR operator will return 1.0 if one of its arguments is zero and the other non-zero, else it returns 0.0. The logical NOT operator returns 1.0 if its argument is 0.0, else it returns 0.0.

These relational and logical operators can be used as a masking or selection operation in a formula. For example, the number of atoms whose properties satisfy one or more criteria could be calculated by taking the returned per-atom vector of ones and zeroes and passing it to the *compute reduce* command.

---

## Math Functions

Math functions are specified as keywords followed by one or more parenthesized arguments “x”, “y”, “z”, each of which can themselves be arbitrarily complex formulas. In this syntax, the arguments can represent scalar values or global vectors or per-atom vectors. In the latter case, the math operation is performed on each element of the vector. For example, “sqrt(natoms)” is the sqrt() of a scalar, where “sqrt(y\*z)” yields a per-atom vector with each element being the sqrt() of the product of one atom’s y and z coordinates.

Most of the math functions perform obvious operations. The ln() is the natural log; log() is the base 10 log.

The random(x,y,z) function takes 3 arguments: x = lo, y = hi, and z = seed. It generates a uniform random number between lo and hi. The normal(x,y,z) function also takes 3 arguments: x = mu, y = sigma, and z = seed. It generates a Gaussian variate centered on mu with variance sigma<sup>2</sup>. In both cases the seed is used the first time the internal random number generator is invoked, to initialize it. For equal-style and vector-style variables, every processor uses the same seed so that they each generate the same sequence of random numbers. For atom-style variables, a unique seed is created for each processor, based on the specified seed. This effectively generates a different random number for each atom being looped over in the atom-style variable.

---

**Note:** Internally, there is just one random number generator for all equal-style and vector-style variables and another one for all atom-style variables. If you define multiple variables (of each style) which use the random() or normal() math functions, then the internal random number generators will only be initialized once, which means only one of the specified seeds will determine the sequence of generated random numbers.

---

The ceil(), floor(), and round() functions are those in the C math library. Ceil() is the smallest integer not less than its argument. Floor() is the largest integer not greater than its argument. Round() is the nearest integer to its argument.

The ramp(x,y) function uses the current timestep to generate a value linearly interpolated between the specified x,y values over the course of a run, according to this formula:

$$\text{value} = x + (y-x) * (\text{timestep}-\text{startstep}) / (\text{stopstep}-\text{startstep})$$

The run begins on startstep and ends on stopstep. Startstep and stopstep can span multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this. If called in between runs or during a *run 0* command, the ramp(x,y) function will return the value of x.

The stagger(x,y) function uses the current timestep to generate a new timestep. X,y > 0 and x > y are required. The generated timesteps increase in a staggered fashion, as the sequence x,x+y,2x,2x+y,3x,3x+y,etc. For any current timestep, the next timestep in the sequence is returned. Thus if stagger(1000,100) is used in a variable by the *dump\_modify every* command, it will generate the sequence of output timesteps:

100, 1000, 1100, 2000, 2100, 3000, etc
--

The logfreq(x,y,z) function uses the current timestep to generate a new timestep. X,y,z > 0 and y < z are required. The generated timesteps are on a base-z logarithmic scale, starting with x, and the y value is how many of the z-1 possible timesteps within one logarithmic interval are generated. I.e. the timesteps follow the sequence x,2x,3x,...y\*x,x\*z,2x\*z,3x\*z,...y\*x\*z,x\*z<sup>2</sup>,2x\*z<sup>2</sup>,etc. For any current timestep, the next timestep in the sequence is returned. Thus if logfreq(100,4,10) is used in a variable by the *dump\_modify every* command, it will generate this sequence of output timesteps:

```
100, 200, 300, 400, 1000, 2000, 3000, 4000, 10000, 20000, etc
```

The `logfreq2(x,y,z)` function is similar to `logfreq`, except a single logarithmic interval is divided into  $y$  equally-spaced timesteps and all of them are output.  $Y < z$  is not required. Thus, if `logfreq2(100,18,10)` is used in a variable by the `dump_modify every` command, then the interval between 100 and 1000 is divided as  $900/18 = 50$  steps, and it will generate the sequence of output timesteps:

```
100, 150, 200, . . . 950, 1000, 1500, 2000, . . . 9500, 10000, 15000, etc
```

The `logfreq3(x,y,z)` function generates  $y$  points between  $x$  and  $z$  (inclusive), that are separated by a multiplicative ratio:  $(z/x)^{1/(y-1)}$ . Constraints are:  $x, z > 0$ ,  $y > 1$ ,  $z-x \geq y-1$ . For eg., if `logfreq3(10,25,1000)` is used in a variable by the `fix print` command, then the interval between 10 and 1000 is divided into 24 parts with a multiplicative separation of  $\sim 1.21$ , and it will generate the following sequence of output timesteps:

```
10, 13, 15, 18, 22, 27, 32, . . . 384, 465, 563, 682, 826, 1000
```

The `stride(x,y,z)` function uses the current timestep to generate a new timestep.  $X, y \geq 0$  and  $z > 0$  and  $x \leq y$  are required. The generated timesteps increase in increments of  $z$ , from  $x$  to  $y$ , i.e. it generates the sequence  $x, x+z, x+2z, \dots, y$ . If  $y-x$  is not a multiple of  $z$ , then similar to the way a for loop operates, the last value will be one that does not exceed  $y$ . For any current timestep, the next timestep in the sequence is returned. Thus if `stride(1000,2000,100)` is used in a variable by the `dump_modify every` command, it will generate the sequence of output timesteps:

```
1000, 1100, 1200, . . . , 1900, 2000
```

The `stride2(x,y,z,a,b,c)` function is similar to the `stride()` function except it generates two sets of strided timesteps, one at a coarser level and one at a finer level. Thus it is useful for debugging, e.g. to produce output every timestep at the point in simulation when a problem occurs.  $X, y \geq 0$  and  $z > 0$  and  $x \leq y$  are required, as are  $a, b \geq 0$  and  $c > 0$  and  $a < b$ . Also,  $a \geq x$  and  $b \leq y$  are required so that the second stride is inside the first. The generated timesteps increase in increments of  $z$ , starting at  $x$ , until  $a$  is reached. At that point the timestep increases in increments of  $c$ , from  $a$  to  $b$ , then after  $b$ , increments by  $z$  are resumed until  $y$  is reached. For any current timestep, the next timestep in the sequence is returned. Thus if `stride2(1000,2000,100,1350,1360,1)` is used in a variable by the `dump_modify every` command, it will generate the sequence of output timesteps:

```
1000, 1100, 1200, 1300, 1350, 1351, 1352, . . . 1359, 1360, 1400, 1500, . . . , 2000
```

The `vdisplace(x,y)` function takes 2 arguments:  $x = \text{value0}$  and  $y = \text{velocity}$ , and uses the elapsed time to change the value by a linear displacement due to the applied velocity over the course of a run, according to this formula:

$$\text{value} = \text{value0} + \text{velocity} * (\text{timestep} - \text{startstep}) * \text{dt}$$

where  $\text{dt} =$  the timestep size.

The run begins on `startstep`. `Startstep` can span multiple runs, using the `start` keyword of the `run` command. See the `run` command for details of how to do this. Note that the `thermo_style` keyword `elaplong = timestep-startstep`. If used between runs this function will return the value according to the end of the last run or the value of  $x$  if used before *any* runs. This function assumes the length of the time step does not change and thus may not be used in combination with `fix dt/reset`.

The `swiggle(x,y,z)` and `cwiggle(x,y,z)` functions each take 3 arguments:  $x = \text{value0}$ ,  $y = \text{amplitude}$ ,  $z = \text{period}$ . They use the elapsed time to oscillate the value by a `sin()` or `cos()` function over the course of a run, according to one of these formulas, where  $\omega = 2 \text{PI} / \text{period}$ :

$$\text{value} = \text{value0} + \text{Amplitude} * \sin(\omega * (\text{timestep} - \text{startstep}) * \text{dt})$$

$$\text{value} = \text{value0} + \text{Amplitude} * (1 - \cos(\omega * (\text{timestep} - \text{startstep}) * \text{dt}))$$

where  $\text{dt} =$  the timestep size.

The run begins on `startstep`. `Startstep` can span multiple runs, using the `start` keyword of the `run` command. See the `run` command for details of how to do this. Note that the `thermo_style` keyword `elaplong = timestep-startstep`. If used between runs these functions will return the value according to the end of the last run or the value of `x` if used before *any* runs. These functions assume the length of the time step does not change and thus may not be used in combination with `fix dt/reset`.

---

## Group and Region Functions

Group functions are specified as keywords followed by one or two parenthesized arguments. The first argument `ID` is the group-ID. The `dim` argument, if it exists, is `x` or `y` or `z`. The `dir` argument, if it exists, is `xmin`, `xmax`, `ymin`, `ymax`, `zmin`, or `zmax`. The `dimdim` argument, if it exists, is `xx` or `yy` or `zz` or `xy` or `yz` or `xz`.

The group function `count()` is the number of atoms in the group. The group functions `mass()` and `charge()` are the total mass and charge of the group. `Xcm()` and `vcm()` return components of the position and velocity of the center of mass of the group. `Fcm()` returns a component of the total force on the group of atoms. `Bound()` returns the min/max of a particular coordinate for all atoms in the group. `Gyration()` computes the radius-of-gyration of the group of atoms. See the `compute gyration` command for a definition of the formula. `Angmom()` returns components of the angular momentum of the group of atoms around its center of mass. `Torque()` returns components of the torque on the group of atoms around its center of mass, based on current forces on the atoms. `Inertia()` returns one of 6 components of the symmetric inertia tensor of the group of atoms around its center of mass, ordered as `Ixx`, `Iyy`, `Izz`, `Ixy`, `Iyz`, `Ixz`. `Omega()` returns components of the angular velocity of the group of atoms around its center of mass.

Region functions are specified exactly the same way as group functions except they take an extra final argument `IDR` which is the region ID. The function is computed for all atoms that are in both the group and the region. If the group is “all”, then the only criteria for atom inclusion is that it be in the region.

---

## Special Functions

Special functions take specific kinds of arguments, meaning their arguments cannot be formulas themselves.

The `sum(x)`, `min(x)`, `max(x)`, `ave(x)`, `trap(x)`, and `slope(x)` functions each take 1 argument which is of the form “`c_ID`” or “`c_ID[N]`” or “`f_ID`” or “`f_ID[N]`” or “`v_name`”. The first two are computes and the second two are fixes; the ID in the reference should be replaced by the ID of a compute or fix defined elsewhere in the input script. The compute or fix must produce either a global vector or array. If it produces a global vector, then the notation without “[N]” should be used. If it produces a global array, then the notation with “[N]” should be used, when N is an integer, to specify which column of the global array is being referenced. The last form of argument “`v_name`” is for a vector-style variable where “`name`” is replaced by the name of the variable.

These functions operate on a global vector of inputs and reduce it to a single scalar value. This is analogous to the operation of the `compute reduce` command, which performs similar operations on per-atom and local vectors.

The `sum()` function calculates the sum of all the vector elements. The `min()` and `max()` functions find the minimum and maximum element respectively. The `ave()` function is the same as `sum()` except that it divides the result by the length of the vector.

The `trap()` function is the same as `sum()` except the first and last elements are multiplied by a weighting factor of 1/2 when performing the sum. This effectively implements an integration via the trapezoidal rule on the global vector of data. I.e. consider a set of points, equally spaced by 1 in their x coordinate: (1,V1), (2,V2), ..., (N,VN), where the `Vi` are the values in the global vector of length N. The integral from 1 to N of these points is `trap()`. When appropriately normalized by the timestep size, this function is useful for calculating integrals of time-series data, like that generated by the `fix ave/correlate` command.



The `slope()` function uses linear regression to fit a line to the set of points, equally spaced by 1 in their x coordinate: (1,V1), (2,V2), ..., (N,VN), where the  $V_i$  are the values in the global vector of length N. The returned value is the slope of the line. If the line has a single point or is vertical, it returns 1.0e20.

The `gmask(x)` function takes 1 argument which is a group ID. It can only be used in atom-style variables. It returns a 1 for atoms that are in the group, and a 0 for atoms that are not.

The `rmask(x)` function takes 1 argument which is a region ID. It can only be used in atom-style variables. It returns a 1 for atoms that are in the geometric region, and a 0 for atoms that are not.

The `grmask(x,y)` function takes 2 arguments. The first is a group ID, and the second is a region ID. It can only be used in atom-style variables. It returns a 1 for atoms that are in both the group and region, and a 0 for atoms that are not in both.

The `next(x)` function takes 1 argument which is a variable ID (not “v\_foo”, just “foo”). It must be for a file-style or atomfile-style variable. Each time the `next()` function is invoked (i.e. each time the equal-style or atom-style variable is evaluated), the following steps occur.

For file-style variables, the current string value stored by the file-style variable is converted to a numeric value and returned by the function. And the next string value in the file is read and stored. Note that if the line previously read from the file was not a numeric string, then it will typically evaluate to 0.0, which is likely not what you want.

For atomfile-style variables, the current per-atom values stored by the atomfile-style variable are returned by the function. And the next set of per-atom values in the file is read and stored.

Since file-style and atomfile-style variables read and store the first line of the file or first set of per-atoms values when they are defined in the input script, these are the value(s) that will be returned the first time the `next()` function is invoked. If `next()` is invoked more times than there are lines or sets of lines in the file, the variable is deleted, similar to how the `next` command operates.

The `is_file(name)` function is a test whether *name* is a (readable) file and returns 1 in this case, otherwise it returns 0. For that *name* is taken as a literal string and must not have any blanks in it.

The `is_os(name)` function is a test whether *name* is part of the OS information that LAMMPS collects and provides in the `platform::os_info()` function. The argument *name* is interpreted as a regular expression as documented for the `utils::strmatch()` function. This allows to adapt LAMMPS inputs to the OS it runs on:

```
if $(is_os(^Windows)) then &
  "shell copy ${input_dir}\some_file.txt ." &
else &
  "shell cp ${input_dir}/some_file.txt ."
```

The `extract_setting(name)` function enables access to basic settings for the LAMMPS executable and the running simulation via calling the `lammps_extract_setting()` library function. For example, the number of processors (MPI ranks) being used by the simulation or the MPI process ID (for this processor) can be queried, or the number of atom types, bond types and so on. For the full list of available keywords *name* and their meaning, see the documentation for `extract_setting()` via the link in this paragraph.

The `label2type()` function converts type labels into numeric types, using label maps created by the `labelmap` or `read_data` commands. The first argument is the label map kind (atom, bond, angle, dihedral, or improper) and the second argument is the label. The function returns the corresponding numeric type.

## Feature Functions

Feature functions allow to probe the running LAMMPS executable for whether specific features are either active, defined, or available. The functions take two arguments, a *category* and a corresponding *argument*. The arguments are strings and thus cannot be formulas themselves; only  $\$$ -style immediate variable expansion is possible. Return value is either 1.0 or 0.0 depending on whether the function evaluates to true or false, respectively.

The *is\_active(category,feature)* function allows to query for active settings which are grouped by categories. Currently supported categories and arguments are:

- *package*: argument = *gpu* or *intel* or *kokkos* or *omp*
- *newton*: argument = *pair* or *bond* or *any*
- *pair*: argument = *single* or *respa* or *manybody* or *tail* or *shift*
- *comm\_style*: argument = *brick* or *tiled*
- *min\_style*: argument = any of the compiled in minimizer styles
- *run\_style*: argument = any of the compiled in run styles
- *atom\_style*: argument = any of the compiled in atom style
- *pair\_style*: argument = any of the compiled in pair styles
- *bond\_style*: argument = any of the compiled in bond styles
- *angle\_style*: argument = any of the compiled in angle styles
- *dihedral\_style*: argument = any of the compiled in dihedral styles
- *improper\_style*: argument = any of the compiled in improper styles
- *kspace\_style*: argument = any of the compiled in kspace styles

Most of the settings are self-explanatory, the *single* argument in the *pair* category allows to check whether a pair style supports a `Pair::single()` function as needed by compute group/group and others features or LAMMPS, *respa* allows to check whether the inner/middle/outer mode of r-RESPA is supported. In the various style categories, the checking is also done using suffix flags, if available and enabled.

Example 1: disable use of suffix for pppm when using GPU package (i.e. run it on the CPU concurrently to running the pair style on the GPU), but do use the suffix otherwise (e.g. with OPENMP).

```
pair_style lj/cut/coul/long 14.0
if $(is_active(package,gpu)) then "suffix off"
kspace_style pppm
```

Example 2: use r-RESPA with inner/outer cutoff, if supported by pair style, otherwise fall back to using pair and reducing the outer time step

```
timestep $(2.0*(1.0+2.0*is_active(pair,respa)))
if $(is_active(pair,respa)) then "run_style respa 4 3 2 2 improper 1 inner 2 5.5 7.0_
↳outer 3 kspace 4" else "run_style respa 3 3 2 improper 1 pair 2 kspace 3"
```

The *is\_available(category,name)* function allows to query whether a specific optional feature is available, i.e. compiled in. This currently works for the following categories: *command*, *compute*, *fix*, *pair\_style* and *feature*. For all categories except *command* and *feature* also appending active suffixes is tried before reporting failure.

The *feature* category is used to check the availability of compiled in features such as GZIP support, PNG support, JPEG support, FFMPEG support, and C++ exceptions for error handling. Corresponding values for name are *gzip*, *png*, *jpeg*, *ffmpeg* and *exceptions*.

This enables writing input scripts which only dump using a given format if the compiled binary supports it.

```
if "${is_available(feature,png)}" then "print 'PNG supported'" else "print 'PNG not_
↳supported'"

if "${is_available(feature,ffmpeg)}" then "dump 3 all movie 25 movie.mp4 type type zoom 1.
↳6 adiam 1.0"
```

The `is_defined(category,id)` function allows to query categories like `compute`, `dump`, `fix`, `group`, `region`, and `variable` whether an entry with the provided name or id is defined.

## Atom Values and Vectors

Atom values take an integer argument  $I$  from 1 to  $N$ , where  $I$  is the atom-ID, e.g. `x[243]`, which means use the  $x$  coordinate of the atom with ID = 243. Or they can take a variable name, specified as `v_name`, where name is the name of the variable, like `x[v_myIndex]`. The variable can be of any style except *vector* or *atom* or *atomfile* variables. The variable is evaluated and the result is expected to be numeric and is cast to an integer (i.e. 3.4 becomes 3), to use an index, which must be a value from 1 to  $N$ . Note that a “formula” cannot be used as the argument between the brackets, e.g. `x[243+10]` or `x[v_myIndex+1]` are not allowed. To do this a single variable can be defined that contains the needed formula.

Note that the  $0 < \text{atom-ID} \leq N$ , where  $N$  is the largest atom ID in the system. If an ID is specified for an atom that does not currently exist, then the generated value is 0.0.

Atom vectors generate one value per atom, so that a reference like “vx” means the  $x$ -component of each atom’s velocity will be used when evaluating the variable.

The meaning of the different atom values and vectors is mostly self-explanatory. *Mol* refers to the molecule ID of an atom, and is only defined if an *atom\_style* is being used that defines molecule IDs.

Note that many other atom attributes can be used as inputs to a variable by using the *compute property/atom* command and then specifying a quantity from that compute.

## Compute References

Compute references access quantities calculated by a *compute*. The ID in the reference should be replaced by the ID of a compute defined elsewhere in the input script. As discussed in the page for the *compute* command, computes can produce global, per-atom, or local values. Only global and per-atom values can be used in a variable. Computes can also produce a scalar, vector, or array.

An equal-style variable can only use scalar values, which means a global scalar, or an element of a global or per-atom vector or array. A vector-style variable can use scalar values or a global vector of values, or a column of a global array of values. Atom-style variables can use global scalar values. They can also use per-atom vector values, or a column of a per-atom array. See the doc pages for individual computes to see what kind of values they produce.

Examples of different kinds of compute references are as follows. There is typically no ambiguity (see exception below) as to what a reference means, since computes only produce either global or per-atom quantities, never both.

<code>c_ID</code>   global scalar, or per-atom vector
<code>c_ID[I]</code>   $I$ th element of global vector, or atom $I$ ’s value in per-atom vector, or $I$ th column from per-atom array
<code>c_ID[I][J]</code>   $I,J$ element of global array, or atom $I$ ’s $J$ th value in per-atom array

For I and J indices, integers can be specified or a variable name, specified as `v_name`, where name is the name of the variable. The rules for this syntax are the same as for the “Atom Values and Vectors” discussion above.

One source of ambiguity for compute references is when a vector-style variable refers to a compute that produces both a global scalar and a global vector. Consider a compute with ID “foo” that does this, referenced as follows by variable “a”, where “myVec” is another vector-style variable:

```
variable a vector c_foo*v_myVec
```

The reference “c\_foo” could refer to either the global scalar or global vector produced by compute “foo”. In this case, “c\_foo” will always refer to the global scalar, and “C\_foo” can be used to reference the global vector. Similarly if the compute produces both a global vector and global array, then “c\_foo[I]” will always refer to an element of the global vector, and “C\_foo[I]” can be used to reference the Ith column of the global array.

Note that if a variable containing a compute is evaluated directly in an input script (not during a run), then the values accessed by the compute must be current. See the discussion below about “Variable Accuracy”.

---

## Fix References

Fix references access quantities calculated by a *fix*. The ID in the reference should be replaced by the ID of a fix defined elsewhere in the input script. As discussed in the page for the *fix* command, fixes can produce global, per-atom, or local values. Only global and per-atom values can be used in a variable. Fixes can also produce a scalar, vector, or array. An equal-style variable can only use scalar values, which means a global scalar, or an element of a global or per-atom vector or array. Atom-style variables can use the same scalar values. They can also use per-atom vector values. A vector value can be a per-atom vector itself, or a column of an per-atom array. See the doc pages for individual fixes to see what kind of values they produce.

The different kinds of fix references are exactly the same as the compute references listed in the above table, where “c\_” is replaced by “f\_”. Again, there is typically no ambiguity (see exception below) as to what a reference means, since fixes only produce either global or per-atom quantities, never both.

f_ID   global scalar, or per-atom vector
f_ID[I]   Ith element of global vector, or atom I’s value in per-atom vector, or Ith column from per-atom array
f_ID[I][J]   I,J element of global array, or atom I’s Jth value in per-atom array

For I and J indices, integers can be specified or a variable name, specified as `v_name`, where name is the name of the variable. The rules for this syntax are the same as for the “Atom Values and Vectors” discussion above.

One source of ambiguity for fix references is the same ambiguity discussed for compute references above. Namely when a vector-style variable refers to a fix that produces both a global scalar and a global vector. The solution is the same as for compute references. For a fix with ID “foo”, “f\_foo” will always refer to the global scalar, and “F\_foo” can be used to reference the global vector. And similarly for distinguishing between a fix’s global vector versus global array with “f\_foo[I]” versus “F\_foo[I]”.

Note that if a variable containing a fix is evaluated directly in an input script (not during a run), then the values accessed by the fix should be current. See the discussion below about “Variable Accuracy”.

Note that some fixes only generate quantities on certain timesteps. If a variable attempts to access the fix on non-allowed timesteps, an error is generated. For example, the *fix ave/time* command may only generate averaged quantities every 100 steps. See the doc pages for individual fix commands for details.

---

## Variable References

Variable references access quantities stored or calculated by other variables, which will cause those variables to be evaluated. The name in the reference should be replaced by the name of a variable defined elsewhere in the input script.

As discussed on this doc page, equal-style variables generate a single global numeric value, vector-style variables generate a vector of global numeric values, and atom-style and atomfile-style variables generate a per-atom vector of numeric values. All other variables store one or more strings.

The formula for an equal-style variable can use any style of variable including a vector\_style or atom-style or atomfile-style. For these 3 styles, a subscript must be used to access a single value from the vector-, atom-, or atomfile-style variable. If a string-storing variable is used, the string is converted to a numeric value. Note that this will typically produce a 0.0 if the string is not a numeric string, which is likely not what you want.

The formula for a vector-style variable can use any style of variable, including atom-style or atomfile-style variables. For these 2 styles, a subscript must be used to access a single value from the atom-, or atomfile-style variable.

The formula for an atom-style variable can use any style of variable, including other atom-style or atomfile-style variables. If it uses a vector-style variable, a subscript must be used to access a single value from the vector-style variable.

Examples of different kinds of variable references are as follows. There is no ambiguity as to what a reference means, since variables produce only a global scalar or global vector or per-atom vector.

v_name	global scalar from equal-style variable
v_name	global vector from vector-style variable
v_name	per-atom vector from atom-style or atomfile-style variable
v_name[I]	Ith element of a global vector from vector-style variable
v_name[I]	value of atom with ID = I from atom-style or atomfile-style variable

For the I index, an integer can be specified or a variable name, specified as v\_name, where name is the name of the variable. The rules for this syntax are the same as for the “Atom Values and Vectors” discussion above.

### 16.137.4 Immediate Evaluation of Variables

If you want an equal-style variable to be evaluated immediately, it may be the case that you do not need to define a variable at all. See the [Commands parse](#) page for info on how to use “immediate” variables in an input script, specified as \$(formula) with parenthesis, where the formula has the same syntax as equal-style variables described on this page. This effectively evaluates a formula immediately without using the variable command to define a named variable.

More generally, there is a difference between referencing a variable with a leading \$ sign (e.g. \$x or \${abc}) versus with a leading “v\_” (e.g. v\_x or v\_abc). The former can be used in any input script command, including a variable command. The input script parser evaluates the reference variable immediately and substitutes its value into the command. As explained on the [Commands parse](#) doc page, you can also use un-named “immediate” variables for this purpose. For example, a string like this  $\$((x_l + x_h)/2 + \sqrt{v\_area})$  in an input script command evaluates the string between the parenthesis as an equal-style variable formula.

Referencing a variable with a leading “v\_” is an optional or required kind of argument for some commands (e.g. the [fix ave/chunk](#) or [dump custom](#) or [thermo\\_style](#) commands) if you wish it to evaluate a variable periodically during a run. It can also be used in a variable formula if you wish to reference a second variable. The second variable will be evaluated whenever the first variable is evaluated.

As an example, suppose you use this command in your input script to define the variable “v” as

```
variable v equal vol
```

before a run where the simulation box size changes. You might think this will assign the initial volume to the variable “v”. That is not the case. Rather it assigns a formula which evaluates the volume (using the `thermo_style` keyword “vol”) to the variable “v”. If you use the variable “v” in some other command like `fix ave/time` then the current volume of the box will be evaluated continuously during the run.

If you want to store the initial volume of the system, you can do it this way:

```
variable v equal vol
variable v0 equal $v
```

The second command will force “v” to be evaluated (yielding the initial volume) and assign that value to the variable “v0”. Thus the command

```
thermo_style custom step v_v v_v0
```

would print out both the current and initial volume periodically during the run.

Note that it is a mistake to enclose a variable formula in double quotes if it contains variables preceded by \$ signs. For example,

```
variable vratio equal "${vfinal}/${v0}"
```

This is because the quotes prevent variable substitution (explained on the [Commands parse](#) doc page), and thus an error will occur when the formula for “vratio” is evaluated later.

---

### 16.137.5 Variable Accuracy

Obviously, LAMMPS attempts to evaluate variables containing formulas (*equal* and *atom* style variables) accurately whenever the evaluation is performed. Depending on what is included in the formula, this may require invoking a *compute*, either directly or indirectly via a *thermo* keyword, or accessing a value previously calculated by a *compute*, or accessing a value calculated and stored by a *fix*. If the *compute* is one that calculates the pressure or energy of the system, then these quantities need to be tallied during the evaluation of the interatomic potentials (pair, bond, etc) on timesteps that the variable will need the values.

LAMMPS keeps track of all of this during a *run* or *energy minimization*. An error will be generated if you attempt to evaluate a variable on timesteps when it cannot produce accurate values. For example, if a *thermo\_style custom* command prints a variable which accesses values stored by a *fix ave/time* command and the timesteps on which thermo output is generated are not multiples of the averaging frequency used in the *fix* command, then an error will occur.

An input script can also request variables be evaluated before or after or in between runs, e.g. by including them in a *print* command. In this case, if a *compute* is needed to evaluate a variable (either directly or indirectly), LAMMPS will not invoke the *compute*, but it will use a value previously calculated by the *compute*, and can do this only if it was invoked on the current timestep. Fixes will always provide a quantity needed by a variable, but the quantity may or may not be current. This leads to one of three kinds of behavior:

- (1) The variable may be evaluated accurately. If it contains references to a *compute* or *fix*, and these values were calculated on the last timestep of a preceding run, then they will be accessed and used by the variable and the result will be accurate.
- (2) LAMMPS may not be able to evaluate the variable and will generate an error message stating so. For example, if the variable requires a quantity from a *compute* that has not been invoked on the current timestep, LAMMPS will generate an error. This means, for example, that such a variable cannot be evaluated before the first run has occurred.

Likewise, in between runs, a variable containing a compute cannot be evaluated unless the compute was invoked on the last timestep of the preceding run, e.g. by thermodynamic output.

One way to get around this problem is to perform a 0-timestep run before using the variable. For example, these commands

```
variable t equal temp
print "Initial temperature = $t"
run 1000
```

will generate an error if the run is the first run specified in the input script, because generating a value for the “t” variable requires a compute for calculating the temperature to be invoked.

However, this sequence of commands would be fine:

```
run 0
variable t equal temp
print "Initial temperature = $t"
run 1000
```

The 0-timestep run initializes and invokes various computes, including the one for temperature, so that the value it stores is current and can be accessed by the variable “t” after the run has completed. Note that a 0-timestep run does not alter the state of the system, so it does not change the input state for the 1000-timestep run that follows. Also note that the 0-timestep run must actually use and invoke the compute in question (e.g. via *thermo* or *dump* output) in order for it to enable the compute to be used in a variable after the run. Thus if you are trying to print a variable that uses a compute you have defined, you can insure it is invoked on the last timestep of the preceding run by including it in thermodynamic output.

Unlike computes, *fixes* will never generate an error if their values are accessed by a variable in between runs. They always return some value to the variable. However, the value may not be what you expect if the fix has not yet calculated the quantity of interest or it is not current. For example, the *fix indent* command stores the force on the indenter. But this is not computed until a run is performed. Thus if a variable attempts to print this value before the first run, zeroes will be output. Again, performing a 0-timestep run before printing the variable has the desired effect.

(3) The variable may be evaluated incorrectly and LAMMPS may have no way to detect this has occurred. Consider the following sequence of commands:

```
pair_coeff 1 1 1.0 1.0
run 1000
pair_coeff 1 1 1.5 1.0
variable e equal pe
print "Final potential energy = $e"
```

The first run is performed using one setting for the pairwise potential defined by the *pair\_style* and *pair\_coeff* commands. The potential energy is evaluated on the final timestep and stored by the *compute pe* compute (this is done by the *thermo\_style* command). Then a pair coefficient is changed, altering the potential energy of the system. When the potential energy is printed via the “e” variable, LAMMPS will use the potential energy value stored by the *compute pe* compute, thinking it is current. There are many other commands which could alter the state of the system between runs, causing a variable to evaluate incorrectly.

The solution to this issue is the same as for case (2) above, namely perform a 0-timestep run before the variable is evaluated to insure the system is up-to-date. For example, this sequence of commands would print a potential energy that reflected the changed pairwise coefficient:

```
pair_coeff 1 1 1.0 1.0
run 0
```

(continues on next page)

(continued from previous page)

```
pair_coeff 1 1 1.5 1.0
run 0
variable e equal pe
print "Final potential energy = $e"
```

### 16.137.6 Restrictions

Indexing any formula element by global atom ID, such as an atom value, requires the *atom style* to use a global mapping in order to look up the vector indices. By default, only atom styles with molecular information create global maps. The *atom\_modify map* command can override the default, e.g. for atomic-style atom styles.

All *universe*- and *uloop*-style variables defined in an input script must have the same number of values.

### 16.137.7 Related commands

*next*, *jump*, *include*, *temper*, *fix print*, *print*

### 16.137.8 Default

none

## 16.138 velocity command

### 16.138.1 Syntax

```
velocity group-ID style args keyword value ...
```

- group-ID = ID of group of atoms whose velocity will be changed
- style = *create* or *set* or *scale* or *ramp* or *zero*
  - create* args = temp seed
    - temp = temperature value (temperature units)
    - seed = random # seed (positive integer)
  - set* args = vx vy vz
    - vx,vy,vz = velocity value or NULL (velocity units)
    - any of vx,vy,vz can be a variable (see below)
  - scale* arg = temp
    - temp = temperature value (temperature units)
  - ramp* args = vdim vlo vhi dim clo chi
    - vdim = vx or vy or vz
    - vlo,vhi = lower and upper velocity value (velocity units)
    - dim = x or y or z
    - clo,chi = lower and upper coordinate bound (distance units)
  - zero* arg = *linear* or *angular*
    - linear* = zero the linear momentum
    - angular* = zero the angular momentum



- zero or more keyword/value pairs may be appended
- keyword = *dist* or *sum* or *mom* or *rot* or *temp* or *bias* or *loop* or *units*

```

dist value = uniform or gaussian
sum value = no or yes
mom value = no or yes
rot value = no or yes
temp value = temperature compute ID
bias value = no or yes
loop value = all or local or geom
rigid value = fix-ID
    fix-ID = ID of rigid body fix
units value = box or lattice

```

## 16.138.2 Examples

```

velocity all create 300.0 4928459 rot yes dist gaussian
velocity border set NULL 4.0 v_vz sum yes units box
velocity flow scale 300.0
velocity flow ramp vx 0.0 5.0 y 5 25 temp mytemp
velocity all zero linear

```

## 16.138.3 Description

Set or change the velocities of a group of atoms in one of several styles. For each style, there are required arguments and optional keyword/value parameters. Not all options are used by each style. Each option has a default as listed below.

The *create* style generates an ensemble of velocities using a random number generator with the specified seed at the specified temperature.

The *set* style sets the velocities of all atoms in the group to the specified values. If any component is specified as NULL, then it is not set. Any of the vx,vy,vz velocity components can be specified as an equal-style or atom-style *variable*. If the value is a variable, it should be specified as v\_name, where name is the variable name. In this case, the variable will be evaluated, and its value used to determine the velocity component. Note that if a variable is used, the velocity it calculates must be in box units, not lattice units; see the discussion of the *units* keyword below.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters or other parameters.

Atom-style variables can specify the same formulas as equal-style variables but can also include per-atom values, such as atom coordinates. Thus it is easy to specify a spatially-dependent velocity field.

The *scale* style computes the current temperature of the group of atoms and then rescales the velocities to the specified temperature.

The *ramp* style is similar to that used by the *compute temp/ramp* command. Velocities ramped uniformly from v<sub>lo</sub> to v<sub>hi</sub> are applied to dimension vx, or vy, or vz. The value assigned to a particular atom depends on its relative coordinate value (in dim) from c<sub>lo</sub> to c<sub>hi</sub>. For the example above, an atom with y-coordinate of 10 (1/4 of the way from 5 to 25), would be assigned a x-velocity of 1.25 (1/4 of the way from 0.0 to 5.0). Atoms outside the coordinate bounds (less than 5 or greater than 25 in this case), are assigned velocities equal to v<sub>lo</sub> or v<sub>hi</sub> (0.0 or 5.0 in this case).

The *zero* style adjusts the velocities of the group of atoms so that the aggregate linear or angular momentum is zero. No other changes are made to the velocities of the atoms. If the *rigid* option is specified (see below), then the zeroing is performed on individual rigid bodies, as defined by the *fix rigid* or *fix rigid/small* commands. In other words, zero

linear will set the linear momentum of each rigid body to zero, and zero angular will set the angular momentum of each rigid body to zero. This is done by adjusting the velocities of the atoms in each rigid body.

All temperatures specified in the velocity command are in temperature units; see the *units* command. The units of velocities and coordinates depend on whether the *units* keyword is set to *box* or *lattice*, as discussed below.

For all styles, no atoms are assigned z-component velocities if the simulation is 2d; see the *dimension* command.

---

The keyword/value options are used in the following ways by the various styles.

The *dist* keyword is used by *create*. The ensemble of generated velocities can be a *uniform* distribution from some minimum to maximum value, scaled to produce the requested temperature. Or it can be a *gaussian* distribution with a mean of 0.0 and a sigma scaled to produce the requested temperature.

The *sum* keyword is used by all styles, except *zero*. The new velocities will be added to the existing ones if *sum* = yes, or will replace them if *sum* = no.

The *mom* and *rot* keywords are used by *create*. If *mom* = yes, the linear momentum of the newly created ensemble of velocities is zeroed; if *rot* = yes, the angular momentum is zeroed.

---

If specified, the *temp* keyword is used by *create* and *scale* to specify a *compute* that calculates temperature in a desired way, e.g. by first subtracting out a velocity bias, as discussed on the *Howto thermostat* doc page. If this keyword is not specified, *create* and *scale* calculate temperature using a compute that is defined internally as follows:

```
compute velocity_temp group-ID temp
```

where group-ID is the same ID used in the velocity command. i.e. the group of atoms whose velocity is being altered. This compute is deleted when the velocity command is finished. See the *compute temp* command for details. If the calculated temperature should have degrees-of-freedom removed due to fix constraints (e.g. SHAKE or rigid-body constraints), then the appropriate fix command must be specified before the velocity command is issued.

The *bias* keyword with a *yes* setting is used by *create* and *scale*, but only if the *temp* keyword is also used to specify a *compute* that calculates temperature in a desired way. If the temperature compute also calculates a velocity bias, the bias is subtracted from atom velocities before the *create* and *scale* operations are performed. After the operations, the bias is added back to the atom velocities. See the *Howto thermostat* page for more discussion of temperature computes with biases. Note that the velocity bias is only applied to atoms in the temperature compute specified with the *temp* keyword.

As an example, assume atoms are currently streaming in a flow direction (which could be separately initialized with the *ramp* style), and you wish to initialize their thermal velocity to a desired temperature. In this context thermal velocity means the per-particle velocity that remains when the streaming velocity is subtracted. This can be done using the *create* style with the *temp* keyword specifying the ID of a *compute temp/ramp* or *compute temp/profile* command, and the *bias* keyword set to a *yes* value.

---

The *loop* keyword is used by *create* in the following ways.

If *loop* = all, then each processor loops over all atoms in the simulation to create velocities, but only stores velocities for atoms it owns. This can be a slow loop for a large simulation. If atoms were read from a data file, the velocity assigned to a particular atom will be the same, independent of how many processors are being used. This will not be the case if atoms were created using the *create\_atoms* command, since atom IDs will likely be assigned to atoms differently.

If *loop* = local, then each processor loops over only its atoms to produce velocities. The random number seed is adjusted to give a different set of velocities on each processor. This is a fast loop, but the velocity assigned to a particular atom will depend on which processor owns it. Thus the results will always be different when a simulation is run on a different number of processors.

---

If `loop = geom`, then each processor loops over only its atoms. For each atom a unique random number seed is created, based on the atom's xyz coordinates. A velocity is generated using that seed. This is a fast loop and the velocity assigned to a particular atom will be the same, independent of how many processors are used. However, the set of generated velocities may be more correlated than if the *all* or *local* keywords are used.

Note that the *loop geom* keyword will not necessarily assign identical velocities for two simulations run on different machines. This is because the computations based on xyz coordinates are sensitive to tiny differences in the double-precision value for a coordinate as stored on a particular machine.

---

The *rigid* keyword only has meaning when used with the *zero* style. It allows specification of a fix-ID for one of the *rigid-body fix* variants which defines a set of rigid bodies. The zeroing of linear or angular momentum is then performed for each rigid body defined by the fix, as described above.

The *units* keyword is used by *set* and *ramp*. If `units = box`, the velocities and coordinates specified in the velocity command are in the standard units described by the *units* command (e.g. Angstroms/fs for real units). If `units = lattice`, velocities are in units of lattice spacings per time (e.g. spacings/fs) and coordinates are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing.

---

## 16.138.4 Restrictions

Assigning a temperature via the *create* style to a system with *rigid bodies* or *SHAKE constraints* may not have the desired outcome for two reasons. First, the velocity command can be invoked before all of the relevant fixes are created and initialized and the number of adjusted degrees of freedom (DOFs) is known. Thus it is not possible to compute the target temperature correctly. Second, the assigned velocities may be partially canceled when constraints are first enforced, leading to a different temperature than desired. A workaround for this is to perform a *run 0* command, which insures all DOFs are accounted for properly, and then rescale the temperature to the desired value before performing a simulation. For example:

```
velocity all create 300.0 12345
run 0 # temperature may not be 300K
velocity all scale 300.0 # now it should be
```

## 16.138.5 Related commands

*fix rigid, fix shake, lattice*

## 16.138.6 Default

The keyword defaults are `dist = uniform`, `sum = no`, `mom = yes`, `rot = no`, `bias = no`, `loop = all`, and `units = lattice`. The `temp` and `rigid` keywords are not defined by default.

## 16.139 write\_coeff command

### 16.139.1 Syntax

```
write_coeff file
```

- file = name of data file to write out

### 16.139.2 Examples

```
write_coeff polymer.coeff
```

### 16.139.3 Description

Write a text format file with the currently defined force field coefficients in a way, that it can be read by LAMMPS with the *include* command. In combination with the *nocoeff* option of *write\_data* this can be used to move the Coeff's sections from a data file into a separate file.

---

**Note:** The *write\_coeff* command is not yet fully implemented as some pair styles do not output their coefficient information. This means you will need to add/copy this information manually.

---

### 16.139.4 Restrictions

none

### 16.139.5 Related commands

*read\_data*, *write\_restart*, *write\_data*

## 16.140 write\_data command

### 16.140.1 Syntax

```
write_data file keyword value ...
```

- file = name of data file to write out
- zero or more keyword/value pairs may be appended
- keyword = *pair* or *nocoeff* or *nofix* or *nolabelmap*

```

nocoeff = do not write out force field info
nofix = do not write out extra sections read by fixes
nolabelmap = do not write out type labels
types value = numeric or labels
pair value = ii or ij
  ii = write one line of pair coefficient info per atom type
  ij = write one line of pair coefficient info per IJ atom type pair

```

## 16.140.2 Examples

```

write_data data.polymer
write_data data.*

```

## 16.140.3 Description

Write a data file in text format of the current state of the simulation. Data files can be read by the *read data* command to begin a simulation. The *read\_data* command also describes their format.

Similar to *dump* files, the data filename can contain a “\*” wild-card character. The “\*” is replaced with the current timestep value.

### Data in Coeff sections

The *write\_data* command may not always write all coefficient settings to the corresponding Coeff sections of the data file. This can have one of multiple reasons. 1) A few styles may be missing the code that would write those sections (if you come across one, please notify the LAMMPS developers). 2) Some pair styles require a single *pair\_coeff* statement and those are not compatible with data files. 3) The default for *write\_data* is to write a *PairCoeff* section, which has only entries for atom types  $i == j$ . The remaining coefficients would be inferred through the currently selected mixing rule. If there has been a *pair\_coeff* command with  $i != j$ , this setting would be lost. LAMMPS will detect this and print a warning message unless *pair ij* is appended to the *write\_data* command. This will request writing a *PairIJCoeff* section which has information for all pairs of atom types. In cases where the coefficient data in the data file is incomplete, you will need to re-specify that information in your input script that reads the data file.

Because a data file is in text format, if you use a data file written out by this command to restart a simulation, the initial state of the new run will be slightly different than the final state of the old run (when the file was written) which was represented internally by LAMMPS in binary format. A new simulation which reads the data file will thus typically diverge from a simulation that continued in the original input script.

If you want to do more exact restarts, using binary files, see the *restart*, *write\_restart*, and *read\_restart* commands. You can also convert binary restart files to text data files, after a simulation has run, using the *-r command-line switch*.

**Note:** Only limited information about a simulation is stored in a data file. For example, no information about atom *groups* and *fixes* are stored. *Binary restart files* store more information.

Bond interactions (angle, etc) that have been turned off by the *fix shake* or *delete\_bonds* command will be written to a data file as if they are turned on. This means they will need to be turned off again in a new run after the data file is read.

Bonds that are broken (e.g. by a bond-breaking potential) are not written to the data file. Thus these bonds will not exist when the data file is read.

Use of the *nocoeff* keyword means no force field parameters are written to the data file. This can be helpful, for example, if you want to make significant changes to the force field or if the force field parameters are read in separately, e.g. from an include file.

Use of the *nofix* keyword means no extra sections read by fixes are written to the data file (see the *fix* option of the *read\_data* command for details). For example, this option excludes sections for user-created per-atom properties from *fix property/atom*.

The *nolabelmap* and *types* keywords refer to type labels that may be defined for numeric atom types, bond types, angle types, etc. The label map can be defined in two ways, either by the *labelmap* command or in data files read by the *read\_data* command which have sections for Atom Type Labels, Bond Type Labels, Angle Type Labels, etc. See the *Howto type labels* doc page for the allowed syntax of type labels and a general discussion of how type labels can be used.

Use of the *nolabelmap* keyword means that even if type labels exist for a given type-kind (Atoms, Bonds, Angles, etc.), type labels are not written to the data file. By default, they are written if they exist. A type label must be defined for every numeric type (within a given type-kind) to be written to the data file.

The *types* keyword determines how atom types, bond types, angle types, etc are written into these data file sections: Atoms, Bonds, Angles, etc. The default is the *numeric* setting, even if type label maps exist. If the *labels* setting is used, type labels will be written to the data file, if the corresponding label map exists. Note that when using *types labels*, the *nolabelmap* keyword cannot be used.

The *pair* keyword lets you specify in what format the pair coefficient information is written into the data file. If the value is specified as *ii*, then one line per atom type is written, to specify the coefficients for each of the I=J interactions. This means that no cross-interactions for I != J will be specified in the data file and the pair style will apply its mixing rule, as documented on individual *pair\_style* doc pages. Of course this behavior can be overridden in the input script after reading the data file, by specifying additional *pair\_coeff* commands for any desired I,J pairs.

If the value is specified as *ij*, then one line of coefficients is written for all I,J pairs where I <= J. These coefficients will include any specific settings made in the input script up to that point. The presence of these I != J coefficients in the data file will effectively turn off the default mixing rule for the pair style. Again, the coefficient values in the data file can be overridden in the input script after reading the data file, by specifying additional *pair\_coeff* commands for any desired I,J pairs.

---

#### 16.140.4 Restrictions

This command requires inter-processor communication to migrate atoms before the data file is written. This means that your system must be ready to perform a simulation before using this command (force fields setup, atom masses initialized, etc).

#### 16.140.5 Related commands

*read\_data*, *write\_restart*

## 16.140.6 Default

The option defaults are `pair = ii` and `types_style = numeric`.

## 16.141 write\_dump command

### 16.141.1 Syntax

```
write_dump group-ID style file dump-args modify dump_modify-args
```

- `group-ID` = ID of the group of atoms to be dumped
- `style` = any of the supported *dump styles*
- `file` = name of file to write dump info to
- `dump-args` = any additional args needed for a particular *dump style*
- `modify` = all args after this keyword are passed to *dump\_modify* (optional)
- `dump-modify-args` = args for *dump\_modify* (optional)

### 16.141.2 Examples

```
write_dump all atom dump.atom
write_dump subgroup atom dump.run.bin
write_dump all custom dump.myforce.* id type x y vx fx
write_dump flow custom dump.%.myforce id type c_myF[3] v_ke modify sort id
write_dump all xyz system.xyz modify sort id element O H
write_dump all image snap*.jpg type type size 960 960 modify bgcolor white
write_dump all image snap*.jpg element element &
    bond atom 0.3 shiny 0.1 ssao yes 6345 0.2 size 1600 1600 &
    modify bgcolor white element C C O H N C C C O H H S O H
write_dump all atom/gz dump.atom.gz modify compression_level 9
write_dump flow custom/zstd dump.%.myforce.zst &
    id type c_myF[3] v_ke &
    modify sort id &
    compression_level 15
```

### 16.141.3 Description

Dump a single snapshot of atom quantities to one or more files for the current state of the system. This is a one-time immediate operation, in contrast to the *dump* command which will set up a dump style to write out snapshots periodically during a running simulation.

The syntax for this command is mostly identical to that of the *dump* and *dump\_modify* commands as if they were concatenated together, with the following exceptions: There is no need for a dump ID or dump frequency and the keyword *modify* is added. The latter is so that the full range of *dump\_modify* options can be specified for the single snapshot, just as they can be for multiple snapshots. The *modify* keyword separates the arguments that would normally be passed to the *dump* command from those that would be given the *dump\_modify*. Both support optional arguments and thus LAMMPS needs to be able to cleanly separate the two sets of args.

Note that if the specified filename uses wildcard characters “\*” or “%”, as supported by the *dump* command, they will operate in the same fashion to create the new filename(s). Normally, *dump image* files require a filename with a “\*” character for the timestep. That is not the case for the *write\_dump* command; no wildcard “\*” character is necessary.

---

#### 16.141.4 Restrictions

All restrictions for the *dump* and *dump\_modify* commands apply to this command as well, with the exception of the *dump image* filename not requiring a wildcard “\*” character, as noted above.

Since dumps are normally written during a *run* or *energy minimization*, the simulation has to be ready to run before this command can be used. Similarly, if the dump requires information from a compute, fix, or variable, the information needs to have been calculated for the current timestep (e.g. by a prior run), else LAMMPS will generate an error message.

For example, it is not possible to dump per-atom energy with this command before a run has been performed, since no energies and forces have yet been calculated. See the *variable* doc page section on Variable Accuracy for more information on this topic.

#### 16.141.5 Related commands

*dump*, *dump image*, *dump\_modify*

#### 16.141.6 Default

The defaults are listed on the doc pages for the *dump* and *dump image* and *dump\_modify* commands.

### 16.142 write\_restart command

#### 16.142.1 Syntax

```
write_restart file keyword value ...
```

- file = name of file to write restart information to
- zero or more keyword/value pairs may be appended
- keyword = *fileper* or *nfile*

*fileper* arg = Np

Np = write one file for every this many processors

*nfile* arg = Nf

Nf = write this many files, one from each of Nf processors



## 16.142.2 Examples

```
write_restart restart.equil
write_restart restart.equil.mpio
write_restart poly.%.* nfile 10
```

## 16.142.3 Description

Write a binary restart file of the current state of the simulation.

During a long simulation, the *restart* command is typically used to output restart files periodically. The `write_restart` command is useful after a minimization or whenever you wish to write out a single current restart file.

Similar to *dump* files, the restart filename can contain two wild-card characters. If a “\*” appears in the filename, it is replaced with the current timestep value. If a “%” character appears in the filename, then one file is written by each processor and the “%” character is replaced with the processor ID from 0 to P-1. An additional file with the “%” replaced by “base” is also written, which contains global information. For example, the files written for filename `restart.%` would be `restart.base`, `restart.0`, `restart.1`, ... `restart.P-1`. This creates smaller files and can be a fast mode of output and subsequent input on parallel machines that support parallel I/O. The optional *fileper* and *nfile* keywords discussed below can alter the number of files written.

The restart file can also be written in parallel as one large binary file via the MPI-IO library, which is part of the MPI standard for versions 2.0 and above. Using MPI-IO requires two steps. First, build LAMMPS with its MPIIO package installed, e.g.

### CMake build

```
cmake . -DPKG_MPIIO=on # enables the MPIIO package in the build folder
cmake --build . # recompiles LAMMPS with the package code included
```

### Traditional make

```
make yes-mpiio # installs the MPIIO package
make mpi # build LAMMPS for your platform
```

Second, use a restart filename which contains “.mpio”. Note that it does not have to end in “.mpio”, just contain those characters. Unlike MPI-IO dump files, a particular restart file must be both written and read using MPI-IO.

Restart files can be read by a *read\_restart* command to restart a simulation from a particular state. Because the file is binary (to enable exact restarts), it may not be readable on another machine. In this case, you can use the *-r command-line switch* to convert a restart file to a data file.

**Note:** Although the purpose of restart files is to enable restarting a simulation from where it left off, not all information about a simulation is stored in the file. For example, the list of fixes that were specified during the initial run is not stored, which means the new input script must specify any fixes you want to use. Even when restart information is stored in the file, as it is for some fixes, commands may need to be re-specified in the new input script, in order to re-use that information. Details are usually given in the documentation of the respective command. Also, see the *read\_restart* command for general information about what is stored in a restart file.

The optional *nfile* or *fileper* keywords can be used in conjunction with the “%” wildcard character in the specified restart file name. As explained above, the “%” character causes the restart file to be written in pieces, one piece for each of P processors. By default P = the number of processors the simulation is running on. The *nfile* or *fileper* keyword can be used to set P to a smaller value, which can be more efficient when running on a large number of processors.

The *nfile* keyword sets P to the specified Nf value. For example, if Nf = 4, and the simulation is running on 100 processors, 4 files will be written, by processors 0,25,50,75. Each will collect information from itself and the next 24 processors and write it to a restart file.

For the *fileper* keyword, the specified value of Np means write one file for every Np processors. For example, if Np = 4, every fourth processor (0,4,8,12,etc) will collect information from itself and the next 3 processors and write it to a restart file.

---

#### 16.142.4 Restrictions

This command requires inter-processor communication to migrate atoms before the restart file is written. This means that your system must be ready to perform a simulation before using this command (force fields setup, atom masses initialized, etc).

To write and read restart files in parallel with MPI-IO, the MPIIO package must be installed.

#### 16.142.5 Related commands

*restart, read\_restart, write\_data*

#### 16.142.6 Default

none

## 17.1 fix accelerate/cos command

### 17.1.1 Syntax

```
fix ID group-ID accelerate value
```

- ID, group-ID are documented in *fix* command
- accelerate/cos = style name of this fix command
- value = amplitude of acceleration (in unit of velocity/time)

### 17.1.2 Examples

```
fix 1 all accelerate/cos 0.02e-5
```

### 17.1.3 Description

Give each atom a acceleration in x-direction based on its z coordinate. The acceleration is a periodic function along the z-direction:

$$a_x(z) = A \cos\left(\frac{2\pi z}{l_z}\right)$$

where  $A$  is the acceleration amplitude,  $l_z$  is the z-length of the simulation box. At steady state, the acceleration generates a velocity profile:

$$v_x(z) = V \cos\left(\frac{2\pi z}{l_z}\right)$$

The generated velocity amplitude  $V$  is related to the shear viscosity  $\eta$  by:

$$V = \frac{A\rho}{\eta} \left(\frac{l_z}{2\pi}\right)^2$$

and it can be obtained from ensemble average of the velocity profile:

$$V = \frac{\sum_i 2m_i v_{i,x} \cos\left(\frac{2\pi z_i}{l_z}\right)}{\sum_i m_i},$$

where  $m_i$ ,  $v_{i,x}$ , and  $z_i$  are the mass,  $x$ -component velocity, and  $z$ -coordinate of a particle, respectively.

The velocity amplitude  $V$  can be calculated with `compute viscosity/cos`, which enables viscosity calculation with periodic perturbation method, as described by [Hess](#). Because the applied acceleration drives the system away from equilibration, the calculated shear viscosity is lower than the intrinsic viscosity due to the shear-thinning effect. Extrapolation to zero acceleration should generally be performed to predict the zero-shear viscosity. As the shear stress decreases, the signal-noise ratio decreases rapidly, the simulation time must be extended accordingly to get converged result.

In order to get meaningful result, the group ID of this fix should be all.

---

### 17.1.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to binary restart files. None of the `fix_modify` options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various output commands. No parameter of this fix can be used with the start/stop keywords of the run command. This fix is not invoked during energy minimization.

### 17.1.5 Restrictions

This command is only available when LAMMPS was built with the MISC package. Since this fix depends on the  $z$ -coordinate of atoms, it cannot be used in 2d simulations.

### 17.1.6 Related commands

`compute viscosity/cos`

### 17.1.7 Default

none

---

(Hess) Hess, B. Journal of Chemical Physics 2002, 116 (1), 209–217.

## 17.2 fix acks2/reaxff command

Accelerator Variants: `acks2/reaxff/kk`

### 17.2.1 Syntax

```
fix ID group-ID acks2/reaxff Nevery cutlo cuthi tolerance params args
```

- ID, group-ID are documented in `fix` command
- `acks2/reaxff` = style name of this fix command
- `Nevery` = perform ACKS2 every this many steps
- `cutlo,cuthi` = lo and hi cutoff for Taper radius

- tolerance = precision to which charges will be equilibrated
- params = reaxff or a filename
- one or more keywords or keyword/value pairs may be appended

```
keyword = maxiter
maxiter N = limit the number of iterations to N
```

## 17.2.2 Examples

```
fix 1 all acks2/reaxff 1 0.0 10.0 1.0e-6 reaxff
fix 1 all acks2/reaxff 1 0.0 10.0 1.0e-6 param.acks2 maxiter 500
```

## 17.2.3 Description

Perform the atom-condensed Kohn–Sham DFT to second order (ACKS2) charge equilibration method as described in (*Verstraelen*). ACKS2 impedes unphysical long-range charge transfer sometimes seen with QEq (e.g., for dissociation of molecules), at increased computational cost. It is typically used in conjunction with the ReaxFF force field model as implemented in the *pair\_style reaxff* command, but it can be used with any potential in LAMMPS, so long as it defines and uses charges on each atom. For more technical details about the charge equilibration performed by `fix acks2/reaxff`, see the (*O’Hearn*) paper.

The ACKS2 method minimizes the electrostatic energy of the system by adjusting the partial charge on individual atoms based on interactions with their neighbors. It requires some parameters for each atom type. If the *params* setting above is the word “reaxff”, then these are extracted from the *pair\_style reaxff* command and the ReaxFF force field file it reads in. If a file name is specified for *params*, then the parameters are taken from the specified file and the file must contain one line for each atom type. The latter form must be used when performing QeQ with a non-ReaxFF potential. The lines should be formatted as follows:

```
bond_softness
itype chi eta gamma bcut
```

where the first line is the global parameter *bond\_softness*. The remaining 1 to Ntypes lines include *itype*, the atom type from 1 to Ntypes, *chi*, the electronegativity in eV, *eta*, the self-Coulomb potential in eV, *gamma*, the valence orbital exponent, and *bcut*, the bond cutoff distance. Note that these 4 quantities are also in the ReaxFF potential file, except that *eta* is defined here as twice the *eta* value in the ReaxFF file. Note that unlike the rest of LAMMPS, the units of this `fix` are hard-coded to be Å, eV, and electronic charge.

The optional *maxiter* keyword allows changing the max number of iterations in the linear solver. The default value is 200.

---

**Note:** In order to solve the self-consistent equations for electronegativity equalization, LAMMPS imposes the additional constraint that all the charges in the `fix` group must add up to zero. The initial charge assignments should also satisfy this constraint. LAMMPS will print a warning if that is not the case.

---

## 17.2.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. This fix computes a global scalar (the number of iterations) for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

This fix is invoked during *energy minimization*.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 17.2.5 Restrictions

This fix is part of the REAXFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix does not correctly handle interactions involving multiple periodic images of the same atom. Hence, it should not be used for periodic cell dimensions less than 10 Å.

This fix may be used in combination with *fix efield* and will apply the external electric field during charge equilibration, but there may be only one *fix efield* instance used, it may only use a constant electric field, and the electric field vector may only have components in non-periodic directions.

## 17.2.6 Related commands

*pair\_style reaxff*, *fix qeq/reaxff*

## 17.2.7 Default

maxiter 200

---

(O’Hearn) O’Hearn, Alperen, Aktulga, SIAM J. Comput., 42(1), C1–C22 (2020).

(Verstraelen) Verstraelen, Ayers, Speybroeck, Waroquier, J. Chem. Phys. 138, 074108 (2013).

## 17.3 fix adapt command

### 17.3.1 Syntax

```
fix ID group-ID adapt N attribute args ... keyword value ...
```

- ID, group-ID are documented in *fix* command
- adapt = style name of this fix command
- N = adapt simulation settings every this many timesteps
- one or more attribute/arg pairs may be appended
- attribute = *pair* or *bond* or *angle* or *kspace* or *atom*

```
pair args = pstyle pparam I J v_name
  pstyle = pair style name (e.g., lj/cut)
  pparam = parameter to adapt over time
  I,J = type pair(s) to set parameter for
  v_name = variable with name that calculates value of pparam
```

```
bond args = bstyle bparam I v_name
  bstyle = bond style name (e.g., harmonic)
  bparam = parameter to adapt over time
  I = type bond to set parameter for
  v_name = variable with name that calculates value of bparam
```

```
angle args = astyle aparam I v_name
  astyle = angle style name (e.g., harmonic)
  aparam = parameter to adapt over time
  I = type angle to set parameter for
  v_name = variable with name that calculates value of aparam
```

```
kspace arg = v_name
  v_name = variable with name that calculates scale factor on k-space terms
```

```
atom args = atomparam v_name
  atomparam = charge or diameter or diameter/disc = parameter to adapt over time
  v_name = variable with name that calculates value of atomparam
```

- zero or more keyword/value pairs may be appended
- keyword = *scale* or *reset* or *mass*

```
scale value = no or yes
  no = the variable value is the new setting
  yes = the variable value multiplies the original setting
```

```
reset value = no or yes
  no = values will remain altered at the end of a run
  yes = reset altered values to their original values at the end of a run
```

```
mass value = no or yes
  no = mass is not altered by changes in diameter
  yes = mass is altered by changes in diameter
```

## 17.3.2 Examples

```

fix 1 all adapt 1 pair soft a 1 1 v_prefactor
fix 1 all adapt 1 pair soft a 2* 3 v_prefactor
fix 1 all adapt 1 pair lj/cut epsilon * * v_scale1 pair coul/cut scale 3 3 v_scale2_
→scale yes reset yes
fix 1 all adapt 10 atom diameter v_size

variable ramp_up equal "ramp(0.01,0.5)"
fix stretch all adapt 1 bond harmonic r0 1 v_ramp_up

```

## 17.3.3 Description

Change or adapt one or more specific simulation attributes or settings over time as a simulation runs. Pair potential and  $k$ -space and atom attributes which can be varied by this fix are discussed below. Many other fixes can also be used to time-vary simulation parameters (e.g., the *fix deform* command will change the simulation box size/shape and the *fix move* command will change atom positions and velocities in a prescribed manner). Also note that many commands allow variables as arguments for specific parameters, if described in that manner on their doc pages. An equal-style variable can calculate a time-dependent quantity, so this is another way to vary a simulation parameter over time.

If  $N$  is specified as 0, the specified attributes are only changed once, before the simulation begins. This is all that is needed if the associated variables are not time-dependent. If  $N > 0$ , then changes are made every  $N$  steps during the simulation, presumably with a variable that is time-dependent.

Depending on the value of the *reset* keyword, attributes changed by this fix will or will not be reset back to their original values at the end of a simulation. Even if *reset* is specified as *yes*, a restart file written during a simulation will contain the modified settings.

If the *scale* keyword is set to *no*, which is the default, then the value of the altered parameter will be whatever the variable generates. If the *scale* keyword is set to *yes*, then the value of the altered parameter will be the initial value of that parameter multiplied by whatever the variable generates (i.e., the variable is now a “scale factor” applied in (presumably) a time-varying fashion to the parameter).

Note that whether *scale* is *no* or *yes*, internally, the parameters themselves are actually altered by this fix. Make sure you use the *reset yes* option if you want the parameters to be restored to their initial values after the run.

The *pair* keyword enables various parameters of potentials defined by the *pair\_style* command to be changed, if the pair style supports it. Note that the *pair\_style* and *pair\_coeff* commands must be used in the usual manner to specify these parameters initially; the fix *adapt* command simply overrides the parameters.

The *pstyle* argument is the name of the pair style. If *pair\_style hybrid* or *hybrid/overlay* is used, *pstyle* should be a sub-style name. If there are multiple sub-styles using the same pair style, then *pstyle* should be specified as “style: $N$ ”, where  $N$  is which instance of the pair style you wish to adapt (e.g., the first or second). For example, *pstyle* could be specified as “soft” or “lubricate” or “lj/cut:1” or “lj/cut:2”. The *pparam* argument is the name of the parameter to change. This is the current list of pair styles and parameters that can be varied by this fix. See the doc pages for individual pair styles and their energy formulas for the meaning of these parameters:

<i>born</i>	a,b,c	type pairs
<i>born/coul/long</i> , <i>born/coul/msm</i>	coulombic_cutoff	type global
<i>buck</i> , <i>buck/coul/cut</i>	a,c	type pairs
<i>buck/coul/long</i> , <i>buck/coul/msm</i>	a,c,coulombic_cutoff	type pairs
<i>buck/mdf</i>	a,c	type pairs

continues on next page



Table 1 – continued from previous page

<i>coul/cut</i>	scale	type pairs
<i>coul/cut/soft</i>	lambda	type pairs
<i>coul/debye</i>	scale	type pairs
<i>coul/dsf</i>	coulombic_cutoff	type global
<i>coul/long, coul/msm</i>	coulombic_cutoff, scale	type pairs
<i>coul/long/soft</i>	scale, lambda, coulombic_cutoff	type pairs
<i>eam, eam/alloy, eam/fs</i>	scale	type pairs
<i>gauss</i>	a	type pairs
<i>lennard/mdf</i>	A,B	type pairs
<i>lj/class2</i>	epsilon,sigma	type pairs
<i>lj/class2/coul/cut, lj/class2/coul/long</i>	epsilon,sigma,coulombic_cutoff	type pairs
<i>lj/cut</i>	epsilon,sigma	type pairs
<i>lj/cut/coul/cut, lj/cut/coul/long, lj/cut/coul/msm</i>	epsilon,sigma,coulombic_cutoff	type pairs
<i>lj/cut/coul/cut/soft, lj/cut/coul/long/soft</i>	epsilon,sigma,lambda,coulombic_cutoff	type pairs
<i>lj/cut/coul/dsf</i>	cutoff	type global
<i>lj/cut/tip4p/cut</i>	epsilon,sigma,coulombic_cutoff	type pairs
<i>lj/cut/soft</i>	epsilon,sigma,lambda	type pairs
<i>lj/expand</i>	epsilon,sigma,delta	type pairs
<i>lj/mdf</i>	epsilon,sigma	type pairs
<i>lj/sf/dipole/sf</i>	epsilon,sigma,scale	type pairs
<i>lubricate</i>	mu	global
<i>mie/cut</i>	epsilon,sigma,gamma_repulsive,gamma_attractive	type pairs
<i>morse, morse/smooth/linear</i>	D0,R0,alpha	type pairs
<i>morse/soft</i>	D0,R0,alpha,lambda	type pairs
<i>nm/cut</i>	E0,R0,m,n	type pairs
<i>nm/cut/coul/cut, nm/cut/coul/long</i>	E0,R0,m,n,coulombic_cutoff	type pairs
<i>reaxff</i>	chi, eta, gamma	type global
<i>snap</i>	scale	type pairs
<i>spin/dmi</i>	coulombic_cutoff	type global
<i>spin/exchange</i>	coulombic_cutoff	type global
<i>spin/magelec</i>	coulombic_cutoff	type global
<i>spin/neel</i>	coulombic_cutoff	type global
<i>table</i>	table_cutoff	type pairs
<i>ufm</i>	epsilon,sigma	type pairs
<i>soft</i>	a	type pairs

**Note:** It is easy to add new pairwise potentials and their parameters to this list. All it typically takes is adding an `extract()` method to the `pair_*.cpp` file associated with the potential.

Some parameters are global settings for the pair style (e.g., the viscosity setting “mu” for *pair\_style lubricate*). Other parameters apply to atom type pairs within the pair style (e.g., the prefactor *a* for *pair\_style soft*).

Note that for many of the potentials, the parameter that can be varied is effectively a prefactor on the entire energy expression for the potential (e.g., the *lj/cut* epsilon). The parameters listed as “scale” are exactly that, since the energy expression for the *coul/cut* potential (for example) has no labeled prefactor in its formula. To apply an effective prefactor to some potentials, multiple parameters need to be altered. For example, the *Buckingham potential* needs both the *A* and *C* terms altered together. To scale the Buckingham potential, you should thus list the pair style twice, once for *A* and once for *C*.

If a type pair parameter is specified, the *I* and *J* settings should be specified to indicate which type pairs to apply it to. If a global parameter is specified, the *I* and *J* settings still need to be specified, but are ignored.

Similar to the *pair\_coeff* command,  $I$  and  $J$  can be specified in one of two ways. Explicit numeric values can be used for each, as in the first example above.  $I \leq J$  is required. LAMMPS sets the coefficients for the symmetric  $J, I$  interaction to the same values.

A wild-card asterisk can be used in place of or in conjunction with the  $I, J$  arguments to set the coefficients for multiple pairs of atom types. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the number of atom types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $m$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive). Note that only type pairs with  $I \leq J$  are considered; if asterisks imply type pairs where  $J < I$ , they are ignored.

IMPORTANT NOTE: If *pair\_style hybrid* or *hybrid/overlay* is being used, then the *pstyle* will be a sub-style name. You must specify  $I, J$  arguments that correspond to type pair values defined (via the *pair\_coeff* command) for that sub-style.

The *v\_name* argument for keyword *pair* is the name of an *equal-style variable* which will be evaluated each time this fix is invoked to set the parameter to a new value. It should be specified as *v\_name*, where *name* is the variable name. Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify parameters that change as a function of time or span consecutive runs in a continuous fashion. For the latter, see the *start* and *stop* keywords of the *run* command and the *elaplog* keyword of *thermo\_style custom* for details.

For example, these commands would change the prefactor coefficient of the *pair\_style soft* potential from 10.0 to 30.0 in a linear fashion over the course of a simulation:

```
variable prefactor equal ramp(10,30)
fix 1 all adapt 1 pair soft a * * v_prefactor
```

---

The *bond* keyword uses the specified variable to change the value of a bond coefficient over time, very similar to how the *pair* keyword operates. The only difference is that now a bond coefficient for a given bond type is adapted.

A wild-card asterisk can be used in place of or in conjunction with the bond type argument to set the coefficients for multiple bond types. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the number of bond types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $m$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive).

Currently *bond* does not support *bond\_style hybrid* nor *bond\_style hybrid/overlay* as bond styles. The bond styles that currently work with *fix\_adapt* are

<i>class2</i>	r0   type bonds
<i>fene</i>	k,r0   type bonds
<i>fene/nm</i>	k,r0   type bonds
<i>gromos</i>	k,r0   type bonds
<i>harmonic</i>	k,r0   type bonds
<i>morse</i>	r0   type bonds
<i>nonlinear</i>	epsilon,r0   type bonds

---

The *angle* keyword uses the specified variable to change the value of an angle coefficient over time, very similar to how the *pair* keyword operates. The only difference is that now an angle coefficient for a given angle type is adapted.

A wild-card asterisk can be used in place of or in conjunction with the angle type argument to set the coefficients for multiple angle types. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the number of angle types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $m$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive).

Currently *angle* does not support *angle\_style hybrid* nor *angle\_style hybrid/overlay* as angle styles. The angle styles that currently work with *fix\_adapt* are

<i>harmonic</i>   k,theta0   type angles
<i>cosine</i>   k   type angles

Note that internally, *theta0* is stored in radians, so the variable this fix uses to reset *theta0* needs to generate values in radians.

The *kspace* keyword used the specified variable as a scale factor on the energy, forces, virial calculated by whatever *k-space* solver is defined by the *kspace\_style* command. If the variable has a value of 1.0, then the solver is unaltered.

The *kspace* keyword works this way whether the *scale* keyword is set to *no* or *yes*.

The *atom* keyword enables various atom properties to be changed. The *aparam* argument is the name of the parameter to change. This is the current list of atom parameters that can be varied by this fix:

- charge = charge on particle
- diameter or diameter/disc = diameter of particle

The *v\_name* argument of the *atom* keyword is the name of an *equal-style variable* which will be evaluated each time this fix is invoked to set, or scale the parameter to a new value. It should be specified as *v\_name*, where *name* is the variable name. See the discussion above describing the formulas associated with equal-style variables. The new value is assigned to the corresponding attribute for all atoms in the fix group.

If the atom parameter is *diameter* and per-atom density and per-atom mass are defined for particles (e.g., *atom\_style granular*), then the mass of each particle is, by default, also changed when the diameter changes. The mass is set from the particle volume for 3d systems (density is assumed to stay constant). For 2d, the default is for LAMMPS to model particles with a radius attribute as spheres. However, if the atom parameter is *diameter/disc*, then the mass is set from the particle area (the density is assumed to be in mass/distance<sup>2</sup> units). The mass of the particle may also be kept constant if the *mass* keyword is set to *no*. This can be useful to account for diameter changes that do not involve mass changes (e.g., thermal expansion).

For example, these commands would shrink the diameter of all granular particles in the “center” group from 1.0 to 0.1 in a linear fashion over the course of a 1000-step simulation:

```
variable size equal ramp(1.0,0.1)
fix 1 center adapt 10 atom diameter v_size
```

This fix can be used in long simulations which are restarted one or more times to continuously adapt simulation parameters, but it must be done carefully. There are two issues to consider. The first is how to adapt the parameters in a continuous manner from one simulation to the next. The second is how, if desired, to reset the parameters to their original values at the end of the last restarted run.

Note that all the parameters changed by this fix are written into a restart file in their current changed state. A new restarted simulation does not know the original time=0 values, unless the input script explicitly resets the parameters (after the restart file is read) to the original values.

Also note that the time-dependent variable(s) used in the restart script should typically be written as a function of time elapsed since the original simulation began.

With this in mind, if the *scale* keyword is set to *no* (the default) in a restarted simulation, original parameters are not needed. The adapted parameters should seamlessly continue their variation relative to the preceding simulation.

If the *scale* keyword is set to *yes*, then the input script should typically reset the parameters being adapted to their original values, so that the scaling formula specified by the variable will operate correctly. An exception is if the *atom* keyword is being used with *scale yes*. In this case, information is added to the restart file so that per-atom properties in the new run will automatically be scaled relative to their original values. This will only work if the *fix adapt* command specified in the restart script has the same ID as the one used in the original script.

In a restarted run, if the *reset* keyword is set to *yes*, and the run ends in this script (as opposed to just writing more restart files), parameters will be restored to the values they were at the beginning of the run command in the restart script, which as explained above, may or may not be the original values of the parameters. Again, an exception is if the *atom* keyword is being used with *reset yes* (in all the runs). In that case, the original per-atom parameters are stored in the restart file, and will be restored when the restarted run finally completes.

---

### 17.3.4 Restart, *fix\_modify*, output, run start/stop, minimize info

If the *atom* keyword is used and the *scale* or *reset* keyword is set to *yes*, then this fix writes information to a restart file so that in a restarted run scaling can continue in a seamless manner and/or the per-atom values can be restored, as explained above.

None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

For *rRESPA time integration*, this fix changes parameters on the outermost rRESPA level.

### 17.3.5 Restrictions

none

### 17.3.6 Related commands

*compute ti*

### 17.3.7 Default

The option defaults are *scale = no*, *reset = no*, *mass = yes*.

## 17.4 *fix adapt/fep* command

### 17.4.1 Syntax

```
fix ID group-ID adapt/fep N attribute args ... keyword value ...
```

- ID, group-ID are documented in *fix* command
- adapt/fep = style name of this fix command
- N = adapt simulation settings every this many timesteps
- one or more attribute/arg pairs may be appended

- attribute = *pair* or *kspace* or *atom*

```
pair args = pstyle pparam I J v_name
  pstyle = pair style name (e.g., lj/cut)
  pparam = parameter to adapt over time
  I,J = type pair(s) to set parameter for
  v_name = variable with name that calculates value of pparam
kspace arg = v_name
  v_name = variable with name that calculates scale factor on K-space terms
atom args = aparam v_name
  aparam = parameter to adapt over time
  I = type(s) to set parameter for
  v_name = variable with name that calculates value of aparam
```

- zero or more keyword/value pairs may be appended

- keyword = *scale* or *reset* or *after*

```
scale value = no or yes
  no = the variable value is the new setting
  yes = the variable value multiplies the original setting
reset value = no or yes
  no = values will remain altered at the end of a run
  yes = reset altered values to their original values at the end
      of a run
after value = no or yes
  no = parameters are adapted at timestep N
  yes = parameters are adapted one timestep after N
```

## 17.4.2 Examples

```
fix 1 all adapt/fep 1 pair soft a 1 1 v_prefactor
fix 1 all adapt/fep 1 pair soft a 2* 3 v_prefactor
fix 1 all adapt/fep 1 pair lj/cut epsilon * * v_scale1 coul/cut scale 3 3 v_scale2 scale_
↪yes reset yes
fix 1 all adapt/fep 10 atom diameter 1 v_size
```

Example input scripts available: examples/PACKAGES/fep

## 17.4.3 Description

Change or adapt one or more specific simulation attributes or settings over time as a simulation runs.

This is an enhanced version of the *fix adapt* command with two differences:

- It is possible to modify the charges of chosen atom types only, instead of scaling all the charges in the system.
- There is a new option *after* for better compatibility with *fix ave/time*.

This version is suited for free energy calculations using *compute ti* or *compute fep*.

If  $N$  is specified as 0, the specified attributes are only changed once, before the simulation begins. This is all that is needed if the associated variables are not time-dependent. If  $N > 0$ , then changes are made every  $N$  steps during the simulation, presumably with a variable that is time-dependent.

Depending on the value of the *reset* keyword, attributes changed by this fix will or will not be reset back to their original values at the end of a simulation. Even if *reset* is specified as *yes*, a restart file written during a simulation will contain the modified settings.

If the *scale* keyword is set to *no*, then the value the parameter is set to will be whatever the variable generates. If the *scale* keyword is set to *yes*, then the value of the altered parameter will be the initial value of that parameter multiplied by whatever the variable generates (i.e., the variable is now a “scale factor” applied in (presumably) a time-varying fashion to the parameter). Internally, the parameters themselves are actually altered; make sure you use the *reset yes* option if you want the parameters to be restored to their initial values after the run.

If the *after* keyword is set to *yes*, then the parameters are changed one timestep after the multiple of N. In this manner, if a fix such as “fix ave/time” is used to calculate averages at every N timesteps, all the contributions to the average will be obtained with the same values of the parameters.

The *pair* keyword enables various parameters of potentials defined by the *pair\_style* command to be changed, if the pair style supports it. Note that the *pair\_style* and *pair\_coeff* commands must be used in the usual manner to specify these parameters initially; the fix adapt command simply overrides the parameters.

The *pstyle* argument is the name of the pair style. If *pair\_style hybrid or hybrid/overlay* is used, *pstyle* should be a sub-style name. For example, *pstyle* could be specified as “soft” or “lubricate”. The *pparam* argument is the name of the parameter to change. This is the current list of pair styles and parameters that can be varied by this fix. See the doc pages for individual pair styles and their energy formulas for the meaning of these parameters:

<i>born</i>	a,b,c	type pairs
<i>buck, buck/coul/cut, buck/coul/long, buck/coul/msm</i>	a,c	type pairs
<i>buck/mdf</i>	a,c	type pairs
<i>coul/cut</i>	scale	type pairs
<i>coul/cut/soft</i>	lambda	type pairs
<i>coul/long, coul/msm</i>	scale	type pairs
<i>coul/long/soft</i>	scale, lambda	type pairs
<i>eam</i>	scale	type pairs
<i>gauss</i>	a	type pairs
<i>lennard/mdf</i>	a,b	type pairs
<i>lj/class2</i>	epsilon,sigma	type pairs
<i>lj/class2/coul/cut, lj/class2/coul/long</i>	epsilon,sigma	type pairs
<i>lj/cut</i>	epsilon,sigma	type pairs
<i>lj/cut/soft</i>	epsilon,sigma,lambda	type pairs
<i>lj/cut/coul/cut, lj/cut/coul/long, lj/cut/coul/msm</i>	epsilon,sigma	type pairs
<i>lj/cut/coul/cut/soft, lj/cut/coul/long/soft</i>	epsilon,sigma,lambda	type pairs
<i>lj/cut/tip4p/cut, lj/cut/tip4p/long</i>	epsilon,sigma	type pairs
<i>lj/cut/tip4p/long/soft</i>	epsilon,sigma,lambda	type pairs
<i>lj/expand</i>	epsilon,sigma,delta	type pairs
<i>lj/mdf</i>	epsilon,sigma	type pairs
<i>lj/sf/dipole/sf</i>	epsilon,sigma,scale	type pairs
<i>mie/cut</i>	epsilon,sigma,gamR,gamA	type pairs
<i>morse, morse/smooth/linear</i>	d0,r0,alpha	type pairs
<i>morse/soft</i>	d0,r0,alpha,lambda	type pairs
<i>nm/cut</i>	e0,r0,nn,mm	type pairs
<i>nm/cut/coul/cut, nm/cut/coul/long</i>	e0,r0,nn,mm	type pairs
<i>snap</i>	scale	type pairs
<i>soft</i>	a	type pairs
<i>ufm</i>	epsilon,sigma,scale	type pairs

---

**Note:** It is easy to add new potentials and their parameters to this list. All it typically takes is adding an extract() method to the pair\_\*.cpp file associated with the potential.

---

Note that for many of the potentials, the parameter that can be varied is effectively a prefactor on the entire energy expression for the potential (e.g., the lj/cut epsilon). The parameters listed as “scale” are exactly that, since the energy expression for the *coul/cut* potential (for example) has no labeled prefactor in its formula. To apply an effective prefactor to some potentials, multiple parameters need to be altered. For example, the *Buckingham potential* needs both the A and C terms altered together. To scale the Buckingham potential, you should thus list the pair style twice, once for A and once for C.

If a type pair parameter is specified, the *I* and *J* settings should be specified to indicate which type pairs to apply it to. If a global parameter is specified, the *I* and *J* settings still need to be specified, but are ignored.

Similar to the *pair\_coeff command*, *I* and *J* can be specified in one of two ways. Explicit numeric values can be used for each, as in the first example above.  $I \leq J$  is required. LAMMPS sets the coefficients for the symmetric J,I interaction to the same values.

A wild-card asterisk can be used in place of or in conjunction with the *I*, *J* arguments to set the coefficients for multiple pairs of atom types. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If *N* is the number of atom types, then an asterisk with no numeric values means all types from 1 to *N*. A leading asterisk means all types from 1 to *n* (inclusive). A trailing asterisk means all types from *m* to *N* (inclusive). A middle asterisk means all types from *m* to *n* (inclusive). Note that only type pairs with  $I \leq J$  are considered; if asterisks imply type pairs where  $J < I$ , they are ignored.

IMPORTANT NOTE: If *pair\_style hybrid* or *hybrid/overlay* is being used, then the *pstyle* will be a sub-style name. You must specify *I*, *J* arguments that correspond to type pair values defined (via the *pair\_coeff* command) for that sub-style.

The *v\_name* argument for keyword *pair* is the name of an *equal-style variable* which will be evaluated each time this fix is invoked to set the parameter to a new value. It should be specified as *v\_name*, where name is the variable name. Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify parameters that change as a function of time or span consecutive runs in a continuous fashion. For the latter, see the *start* and *stop* keywords of the *run* command and the *elaplong* keyword of *thermo\_style custom* for details.

For example, these commands would change the prefactor coefficient of the *pair\_style soft* potential from 10.0 to 30.0 in a linear fashion over the course of a simulation:

```
variable prefactor equal ramp(10,30)
fix 1 all adapt 1 pair soft a * * v_prefactor
```

---

The *kspace* keyword used the specified variable as a scale factor on the energy, forces, virial calculated by whatever *k*-space solver is defined by the *kspace\_style* command. If the variable has a value of 1.0, then the solver is unaltered.

The *kspace* keyword works this way whether the *scale* keyword is set to *no* or *yes*.

---

The *atom* keyword enables various atom properties to be changed. The *aparam* argument is the name of the parameter to change. This is the current list of atom parameters that can be varied by this fix:

- charge = charge on particle
- diameter = diameter of particle

The *I* argument indicates which atom types are affected. A wild-card asterisk can be used in place of or in conjunction with the *I* argument to set the coefficients for multiple atom types.

The `v_name` argument of the `atom` keyword is the name of an *equal-style variable* which will be evaluated each time this fix is invoked to set the parameter to a new value. It should be specified as `v_name`, where `name` is the variable name. See the discussion above describing the formulas associated with equal-style variables. The new value is assigned to the corresponding attribute for all atoms in the fix group.

If the `atom` parameter is `diameter` and per-atom density and per-atom mass are defined for particles (e.g., *atom\_style granular*), then the mass of each particle is also changed when the diameter changes (density is assumed to stay constant).

For example, these commands would shrink the diameter of all granular particles in the “center” group from 1.0 to 0.1 in a linear fashion over the course of a 1000-step simulation:

```
variable size equal ramp(1.0,0.1)
fix 1 center adapt 10 atom diameter * v_size
```

For *rRESPA time integration*, this fix changes parameters on the outermost rRESPA level.

---

#### 17.4.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.4.5 Restrictions

none

#### 17.4.6 Related commands

*compute fep, fix adapt, compute ti, pair\_style \*/soft*

#### 17.4.7 Default

The option defaults are `scale = no`, `reset = no`, `after = no`.

### 17.5 fix addforce command

#### 17.5.1 Syntax

```
fix ID group-ID addforce fx fy fz keyword value ...
```

- ID, group-ID are documented in *fix* command
- addforce = style name of this fix command
- fx,fy,fz = force component values (force units)

any of `fx,fy,fz` can be a variable (see below)



- zero or more keyword/value pairs may be appended to args
- keyword = *every* or *region* or *energy*

*every* value = *Nevery*

*Nevery* = add force every this many time steps

*region* value = region-ID

region-ID = ID of region atoms must be in to have added force

*energy* value = *v\_name*

*v\_name* = variable with name that calculates the potential energy of each atom in  
 →the added force field

## 17.5.2 Examples

```
fix kick flow addforce 1.0 0.0 0.0
fix kick flow addforce 1.0 0.0 v_oscillate
fix ff boundary addforce 0.0 0.0 v_push energy v_espace
```

## 17.5.3 Description

Add  $(f_x, f_y, f_z)$  to the corresponding component of the force for each atom in the group. This command can be used to give an additional push to atoms in a simulation, such as for a simulation of Poiseuille flow in a channel.

Any of the three quantities defining the force components, namely  $f_x$ ,  $f_y$ , and  $f_z$ , can be specified as an equal-style or atom-style *variable*. If the value is a variable, it should be specified as *v\_name*, where name is the variable name. In this case, the variable will be evaluated each time step, and its value(s) will be used to determine the force component(s).

Equal-style variables can specify formulas with various mathematical functions and include *thermo\_style* command keywords for the simulation box parameters, time step, and elapsed time. Thus, it is easy to specify a time-dependent force field.

Atom-style variables can specify the same formulas as equal-style variables but can also include per-atom values, such as atom coordinates. Thus, it is easy to specify a spatially-dependent force field with optional time-dependence as well.

If the *every* keyword is used, the *Nevery* setting determines how often the forces are applied. The default value is 1, for every time step.

If the *region* keyword is used, the atom must also be in the specified geometric *region* in order to have force added to it.

Adding a force to atoms implies a change in their potential energy as they move due to the applied force field. For dynamics via the “run” command, this energy can be optionally added to the system’s potential energy for thermodynamic output (see below). For energy minimization via the “minimize” command, this energy must be added to the system’s potential energy to formulate a self-consistent minimization problem (see below).

The *energy* keyword is not allowed if the added force is a constant vector  $\vec{F} = (f_x, f_y, f_z)$ , with all components defined as numeric constants and not as variables. This is because LAMMPS can compute the energy for each atom directly as

$$E = -\vec{x} \cdot \vec{F} = -(xf_x + yf_y + zf_z),$$

so that  $-\vec{\nabla}E = \vec{F}$ .

The *energy* keyword is optional if the added force is defined with one or more variables, and if you are performing dynamics via the *run* command. If the keyword is not used, LAMMPS will set the energy to 0.0, which is typically fine for dynamics.

The *energy* keyword is required if the added force is defined with one or more variables, and you are performing energy minimization via the “minimize” command. The keyword specifies the name of an atom-style *variable* which is used to compute the energy of each atom as function of its position. Like variables used for  $f_x$ ,  $f_y$ ,  $f_z$ , the energy variable is specified as `v_name`, where name is the variable name.

Note that when the *energy* keyword is used during an energy minimization, you must insure that the formula defined for the atom-style *variable* is consistent with the force variable formulas (i.e., that  $-\vec{\nabla}E = \vec{F}$ ). For example, if the force were a spring-like,  $\vec{F} = -k\vec{x}$ , then the energy formula should be  $E = \frac{1}{2}kx^2$ . If you do not do this correctly, the minimization will not converge properly.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 17.5.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy inferred by the added force to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*. Note that this energy is a fictitious quantity but is needed so that the *minimize* command can include the forces added by this fix in a consistent manner (i.e., there is a decrease in potential energy when atoms move in the direction of the added force).

The *fix\_modify virial* option is supported by this fix to add the contribution due to the added forces on atoms to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global scalar and a global three-vector of forces, which can be accessed by various *output commands*. The scalar is the potential energy discussed above. The vector is the total force on the group of atoms before the forces on individual atoms are changed by the fix. The scalar and vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command. You should not specify force components with a variable that has time-dependence for use with a minimizer, since the minimizer increments the time step as the iteration count during the minimization.

---

**Note:** If you want the fictitious potential energy associated with the added forces to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

## 17.5.5 Restrictions

none

## 17.5.6 Related commands

*fix setforce*, *fix aveforce*

## 17.5.7 Default

The option default for the every keyword is every = 1.

# 17.6 fix addtorque command

## 17.6.1 Syntax

```
fix ID group-ID addtorque Tx Ty Tz
```

- ID, group-ID are documented in *fix* command
- addtorque = style name of this fix command
- Tx,Ty,Tz = torque component values (torque units)
- any of Tx,Ty,Tz can be a variable (see below)

## 17.6.2 Examples

```
fix kick bead addtorque 2.0 3.0 5.0
fix kick bead addtorque 0.0 0.0 v_oscillate
```

## 17.6.3 Description

Add a set of forces to each atom in the group such that:

- the components of the total torque applied on the group (around its center of mass) are  $T_x$ ,  $T_y$ , and  $T_z$
- the group would move as a rigid body in the absence of other forces.

This command can be used to drive a group of atoms into rotation.

Any of the three quantities defining the torque components can be specified as an equal-style *variable*, namely  $T_x$ ,  $T_y$ ,  $T_z$ . If the value is a variable, it should be specified as `v_name`, where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the torque component.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent torque.

## 17.6.4 Restart, `fix_modify`, `output`, `run start/stop`, `minimize` info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential “energy” inferred by the added torques to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*. Note that this is a fictitious quantity but is needed so that the *minimize* command can include the forces added by this fix in a consistent manner (i.e., there is a decrease in potential energy when atoms move in the direction of the added forces).

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its torque. Default is the outermost level.

This fix computes a global scalar and a global 3-vector, which can be accessed by various *output commands*. The scalar is the potential energy discussed above. The vector is the total torque on the group of atoms before the forces on individual atoms are changed by the fix. The scalar and vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the fictitious potential energy associated with the added forces to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

---

**Note:** You should not specify force components with a variable that has time-dependence for use with a minimizer, since the minimizer increments the timestep as the iteration count during the minimization.

---

## 17.6.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 17.6.6 Related commands

*fix addforce*

## 17.6.7 Default

none

## 17.7 `fix amoeba/bitorsion` command

### 17.7.1 Syntax

<code>fix ID group-ID amoeba/bitorsion filename</code>
--

- ID, group-ID are documented in *fix* command
- amoeba/bitorsion = style name of this fix command

- filename = force-field file with AMOEBA bitorsion coefficients

## 17.7.2 Examples

```
fix          bit all amoeba/bitorsion bitorsion.ubiquitin.data
read_data    proteinX.data fix bit bitorsions BiTorsions
fix_modify   bit energy yes
```

## 17.7.3 Description

This command enables 5-body torsion/torsion interactions to be added to simulations which use the AMOEBA and HIPPO force fields. It matches how the Tinker MD code computes its torsion/torsion interactions for the AMOEBA and HIPPO force fields. See the *Howto amoeba* doc page for more information about the implementation of AMOEBA and HIPPO in LAMMPS.

Bitorsion interactions add additional potential energy contributions to pairs of overlapping phi-psi dihedrals of amino-acids, which are important to properly represent their conformational behavior.

The examples/amoeba directory has a sample input script and data file for ubiquitin, which illustrates use of the fix amoeba/bitorsion command.

As in the example above, this fix should be used before reading a data file that contains a listing of bitorsion interactions. The *filename* specified should contain the bitorsion parameters for the AMOEBA or HIPPO force field.

The data file read by the *read\_data* command must contain the topology of all the bitorsion interactions, similar to the topology data for bonds, angles, dihedrals, etc. Specifically it should have a line like this in its header section:

```
N bitorsions
```

where  $N$  is the number of bitorsion 5-body interactions. It should also have a section in the body of the data file like this with  $N$  lines:

```
BiTorsions
  1      1      8      10     12     18     20
  2      5     18     20     22     25     27
  [...]
  N      3    314    315    317    318    330
```

The first column is an index from 1 to  $N$  to enumerate the bitorsion 5-atom tuples; it is ignored by LAMMPS. The second column is the *type* of the interaction; it is an index into the bitorsion force field file. The remaining 5 columns are the atom IDs of the atoms in the two 4-atom dihedrals that overlap to create the bitorsion 5-body interaction. Note that the *bitorsions* and *BiTorsions* keywords for the header and body sections match those specified in the *read\_data* command following the data file name.

The data file should be generated by using the tools/tinker/tinker2Imp.py conversion script which creates a LAMMPS data file from Tinker input files, including its PRM file which contains the parameters necessary for computing bitorsion interactions. The script must be invoked with the optional “-bitorsion” flag to do this; see the example for the ubiquitin system in the tools/tinker/README file. The same conversion script also creates the file of bitorsion coefficient data which is read by this command.

The potential energy associated with bitorsion interactions can be output as described below. It can also be included in the total potential energy of the system, as output by the *thermo\_style* command, if the *fix\_modify energy* command is used, as in the example above. See the note below about how to include the bitorsion energy when performing an *energy minimization*.

### 17.7.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the list of bitorsion interactions to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify energy* option is supported by this fix to add the potential energy of the bitorsion interactions to both the global potential energy and peratom potential energies of the system as part of *thermodynamic output* or output by the *compute pe/atom* command. The default setting for this fix is *fix\_modify energy yes*.

The *fix\_modify virial* option is supported by this fix to add the contribution due to the bitorsion interactions to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial yes*.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the potential energy discussed above. The scalar value calculated by this fix is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

---

**Note:** For energy minimization, if you want the potential energy associated with the bitorsion terms forces to be included in the total potential energy of the system (the quantity being minimized), you **MUST** not disable the *fix\_modify energy* option for this fix.

---

### 17.7.5 Restrictions

To function as expected this fix command must be issued *before* a *read\_data* command but *after* a *read\_restart* command.

This fix can only be used if LAMMPS was built with the AMOEBA package. See the *Build package* page for more info.

### 17.7.6 Related commands

*fix\_modify*, *read\_data*

### 17.7.7 Default

none

## 17.8 fix amoeba/pitorsion command

### 17.8.1 Syntax

```
fix ID group-ID amoeba/pitorsion
```

- ID, group-ID are documented in *fix* command
- amoeba/pitorsion = style name of this fix command

### 17.8.2 Examples

```
fix          pit all amoeba/pitorsion
read_data    proteinX.data fix pit "pitorsion types" "PiTorsion Coeffs" &
             fix pit pitorsions PiTorsions
fix_modify   pit energy yes
```

### 17.8.3 Description

This command enables 6-body pitorsion interactions to be added to simulations which use the AMOEBA and HIPPO force fields. It matches how the Tinker MD code computes its pitorsion interactions for the AMOEBA and HIPPO force fields. See the *Howto amoeba* doc page for more information about the implementation of AMOEBA and HIPPO in LAMMPS.

Pitorsion interactions add additional potential energy contributions to 6-tuples of atoms  $IJKLMN$  that have a bond between atoms  $K$  and  $L$ , where both  $K$  and  $L$  are additionally bonded to exactly two other atoms. Namely,  $K$  is also bonded to  $I$  and  $J$ , and  $L$  is also bonded to  $M$  and  $N$ .

The examples/amoeba directory has a sample input script and data file for ubiquitin, which illustrates use of the fix amoeba/pitorsion command.

As in the example above, this fix should be used before reading a data file that contains a listing of pitorsion interactions.

The data file read by the *read\_data* command must contain the topology of all the pitorsion interactions, similar to the topology data for bonds, angles, dihedrals, etc. Specifically, it should have two lines like these in its header section:

```
M pitorsion types
N pitorsions
```

where  $N$  is the number of pitorsion 5-body interactions and  $M$  is the number of pitorsion types. It should also have two sections in the body of the data file like these with  $M$  and  $N$  lines each:

```
PiTorsion Coeffs
```

```
  1 6.85
  2 10.2
  [...]
  M 6.85
```

```
PiTorsions
```

```
  1      1      8      10      12      18      20
```

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2	5	18	20	22	25	27
[...]						
N	3	314	315	317	318	330

For PiTorsion Coeffs, the first column is an index from 1 to  $M$  to enumerate the pitorsion types. The second column is the single prefactor coefficient needed for each type.

For PiTorsions, the first column is an index from 1 to  $N$  to enumerate the pitorsion 5-atom tuples; it is ignored by LAMMPS. The second column is the “type” of the interaction; it is an index into the PiTorsion Coeffs. The remaining 5 columns are the atom IDs of the atoms in the two 4-atom dihedrals that overlap to create the pitorsion 5-body interaction.

Note that the *pitorsion types* and *pitorsions* and *PiTorsion Coeffs* and *PiTorsions* keywords for the header and body sections of the data file match those specified in the *read\_data* command following the data file name.

The data file should be generated by using the tools/tinker/tinker2lmp.py conversion script which creates a LAMMPS data file from Tinker input files, including its PRM file which contains the parameters necessary for computing pitorsion interactions.

The potential energy associated with pitorsion interactions can be output as described below. It can also be included in the total potential energy of the system, as output by the *thermo\_style* command, if the *fix\_modify energy* command is used, as in the example above. See the note below about how to include the pitorsion energy when performing an *energy minimization*.

#### 17.8.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the list of pitorsion interactions to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify energy* option is supported by this fix to add the potential energy of the pitorsion interactions to both the global potential energy and peratom potential energies of the system as part of *thermodynamic output* or output by the *compute pe/atom* command. The default setting for this fix is *fix\_modify energy yes*.

The *fix\_modify virial* option is supported by this fix to add the contribution due to the pitorsion interactions to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial yes*.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the potential energy discussed above. The scalar value calculated by this fix is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

---

**Note:** For energy minimization, if you want the potential energy associated with the pitorsion terms forces to be included in the total potential energy of the system (the quantity being minimized), you MUST not disable the *fix\_modify energy* option for this fix.

---



## 17.8.5 Restrictions

To function as expected this fix command must be issued *before* a `read_data` command but *after* a `read_restart` command.

This fix can only be used if LAMMPS was built with the AMOEBA package. See the [Build package](#) page for more info.

## 17.8.6 Related commands

`fix_modify`, `read_data`

## 17.8.7 Default

none

# 17.9 fix append/atoms command

## 17.9.1 Syntax

```
fix ID group-ID append/atoms face ... keyword value ...
```

- ID, group-ID are documented in `fix` command
- append/atoms = style name of this fix command
- face = `zhi`
- zero or more keyword/value pairs may be appended
- keyword = `basis` or `size` or `freq` or `temp` or `random` or `units`

`basis` values = M itype

M = which basis atom

itype = atom type (1-N) to assign to this basis atom

`size` args = Lz

Lz = z size of lattice region appended in a single event(distance units)

`freq` args = freq

freq = the number of timesteps between append events

`temp` args = target damp seed extent

target = target temperature for the region between zhi-extent and zhi\_

↪(temperature units)

damp = damping parameter (time units)

seed = random number seed for langevin kicks

extent = extent of thermostatted region (distance units)

`random` args = xmax ymax zmax seed

xmax, ymax, zmax = maximum displacement in particular direction (distance units)

seed = random number seed for random displacement

`units` value = `lattice` or `box`

`lattice` = the wall position is defined in lattice units

`box` = the wall position is defined in simulation box units

## 17.9.2 Examples

```
fix 1 all append/atoms zhi size 5.0 freq 295 units lattice
fix 4 all append/atoms zhi size 15.0 freq 5 units box
fix A all append/atoms zhi size 1.0 freq 1000 units lattice
```

## 17.9.3 Description

This fix creates atoms on a lattice, appended on the zhi edge of the system box. This can be useful when a shock or wave is propagating from zlo. This allows the system to grow with time to accommodate an expanding wave. A simulation box must already exist, which is typically created via the *create\_box* command. Before using this command, a lattice must also be defined using the *lattice* command.

This fix will automatically freeze atoms on the zhi edge of the system, so that overlaps are avoided when new atoms are appended.

The *basis* keyword specifies an atom type that will be assigned to specific basis atoms as they are created. See the *lattice* command for specifics on how basis atoms are defined for the unit cell of the lattice. By default, all created atoms are assigned type = 1 unless this keyword specifies differently.

The *size* keyword defines the size in *z* of the chunk of material to be added.

The *random* keyword will give the atoms random displacements around their lattice points to simulate some initial temperature.

The *temp* keyword will cause a region to be thermostatted with a Langevin thermostat on the zhi boundary. The size of the region is measured from zhi and is set with the *extent* argument.

The *units* keyword determines the meaning of the distance units used to define a wall position, but only when a numeric constant is used. A *box* value selects standard distance units as defined by the *units* command (e.g., Å for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacings.

---

## 17.9.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.9.5 Restrictions

This fix style is part of the SHOCK package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The boundary on which atoms are added with *append/atoms* must be shrink/minimum. The opposite boundary may be any boundary type other than periodic.

## 17.9.6 Related commands

*fix wall/piston* command

## 17.9.7 Default

The keyword defaults are size = 0.0, freq = 0, units = lattice. All added atoms are of type 1 unless the basis keyword is used.

## 17.10 fix atc command

### 17.10.1 Syntax

```
fix <fixID> <group> atc <type> <parameter_file>
```

- fixID = name of fix
- group = name of group fix is to be applied
- type = *thermal* or *two\_temperature* or *hardy* or *field*

*thermal* = thermal coupling with fields: temperature

*two\_temperature* = electron-phonon coupling with field: temperature and electron\_↪temperature

*hardy* = on-the-fly post-processing using kernel localization functions

*field* = on-the-fly post-processing using mesh-based localization functions

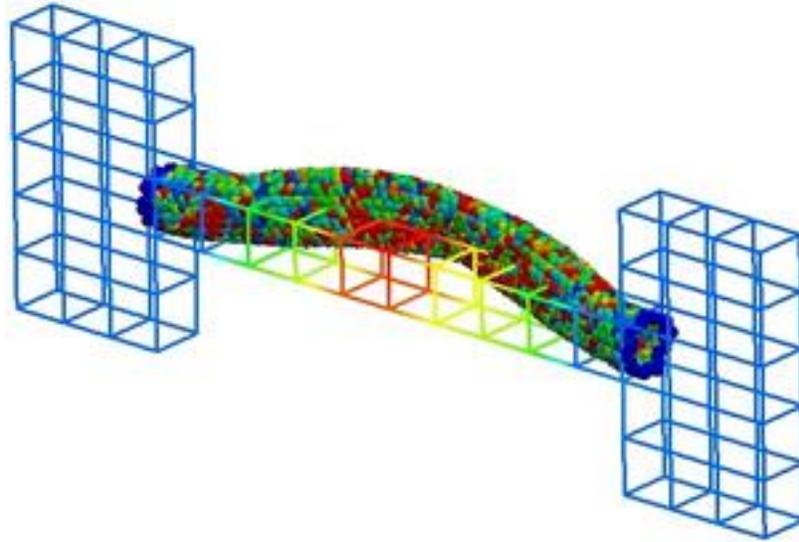
- parameter\_file = name of the file with material parameters. Note: Neither hardy nor field requires a parameter file

### 17.10.2 Examples

```
fix AtC internal atc thermal Ar_thermal.dat
fix AtC internal atc two_temperature Ar_ttm.mat
fix AtC internal atc hardy
fix AtC internal atc field
```

### 17.10.3 Description

This fix is the beginning to creating a coupled FE/MD simulation and/or an on-the-fly estimation of continuum fields. The coupled versions of this fix do Verlet integration and the post-processing does not. After instantiating this fix, several other `fix_modify` commands will be needed to set up the problem (i.e., define the finite element mesh and prescribe initial and boundary conditions).



The following coupling example is typical, but non-exhaustive:

```
# ... commands to create and initialize the MD system

# initial fix to designate coupling type and group to apply it to
# tag group physics material_file
fix AtC internal atc thermal Ar_thermal.mat

# create a uniform 12 x 2 x 2 mesh that covers region contain the group
# nx ny nz region periodicity
fix_modify AtC mesh create 12 2 2 mdRegion f p p

# specify the control method for the type of coupling
# physics control_type
fix_modify AtC thermal control flux

# specify the initial values for the empirical field "temperature"
# field node_group value
fix_modify AtC initial temperature all 30

# create an output stream for nodal fields
# filename output_frequency
fix_modify AtC output atc_fe_output 100

run 1000
```

likewise for this post-processing example:

```
# ... commands to create and initialize the MD system

# initial fix to designate post-processing and the group to apply it to
# no material file is allowed nor required
fix AtC internal atc hardy

# for hardy fix, specific kernel function (function type and range) to # be used as a
↪ localization function
```

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```

fix AtC kernel quartic_sphere 10.0

# create a uniform 1 x 1 x 1 mesh that covers region contain the group
# with periodicity this effectively creates a system average
fix_modify AtC mesh create 1 1 1 box p p p

# change from default lagrangian map to eulerian
# refreshed every 100 steps
fix_modify AtC atom_element_map eulerian 100

# start with no field defined
# add mass density, potential energy density, stress and temperature
fix_modify AtC fields add density energy stress temperature

# create an output stream for nodal fields
# filename output_frequency
fix_modify AtC output nvtFE 100 text

run 1000

```

the mesh's linear interpolation functions can be used as the localization function by using the field option:

```

fix AtC internal atc field
fix_modify AtC mesh create 1 1 1 box p p p
...

```

Note coupling and post-processing can be combined in the same simulations using separate fixes.

#### 17.10.4 Restart, `fix_modify`, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The `fix_modify energy` option is not supported by this fix, but this fix does add the kinetic energy imparted to atoms by the momentum coupling mode of the AtC package to the global potential energy of the system as part of *thermodynamic output*.

Additional `fix_modify` options relevant to this fix are listed below.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the energy discussed in the previous paragraph. The scalar value is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the `run` command. This fix is not invoked during *energy minimization*.

### 17.10.5 Restrictions

Thermal and two\_temperature (coupling) types use a Verlet time-integration algorithm. The hardy type does not contain its own time-integrator and must be used with a separate fix that does contain one (e.g., nve, nvt). In addition, currently:

- the coupling is restricted to thermal physics
- the FE computations are done in serial on each processor.

### 17.10.6 Related commands

After specifying this fix in your input script, several *fix\_modify AtC* commands are used to setup the problem (e.g., define the finite element mesh and prescribe initial and boundary conditions). Each of these options has its own doc page.

*fix\_modify* commands for setup:

- *fix\_modify AtC mesh create*
- *fix\_modify AtC mesh quadrature*
- *fix\_modify AtC mesh read*
- *fix\_modify AtC mesh write*
- *fix\_modify AtC mesh create\_nodeset*
- *fix\_modify AtC mesh add\_to\_nodeset*
- *fix\_modify AtC mesh create\_faceset box*
- *fix\_modify AtC mesh create\_faceset plane*
- *fix\_modify AtC mesh create\_elementset*
- *fix\_modify AtC mesh delete\_elements*
- *fix\_modify AtC mesh nodeset\_to\_elementset*
- *fix\_modify AtC boundary type*
- *fix\_modify AtC internal\_quadrature*
- *fix\_modify AtC time\_integration*
- *fix\_modify AtC extrinsic electron\_integration*
- *fix\_modify AtC internal\_element\_set*
- *fix\_modify AtC decomposition*

*fix\_modify* commands for boundary and initial conditions:

- *fix\_modify AtC initial*
- *fix\_modify AtC fix*
- *fix\_modify AtC unfix*
- *fix\_modify AtC fix\_flux*
- *fix\_modify AtC unfix\_flux*
- *fix\_modify AtC source*
- *fix\_modify AtC remove\_source*

*fix\_modify* commands for control and filtering:

- *fix\_modify AtC control thermal*
- *fix\_modify AtC control momentum*
- *fix\_modify AtC control localized\_lambda*
- *fix\_modify AtC control lumped\_lambda\_solve*
- *fix\_modify AtC control mask\_direction*
- *fix\_modify AtC filter*
- *fix\_modify AtC filter scale*
- *fix\_modify AtC filter type*
- *fix\_modify AtC equilibrium\_start*
- *fix\_modify AtC extrinsic exchange*
- *fix\_modify AtC poisson\_solver*

*fix\_modify* commands for output:

- *fix\_modify AtC output*
- *fix\_modify AtC output nodeset*
- *fix\_modify AtC output volume\_integral*
- *fix\_modify AtC output boundary\_integral*
- *fix\_modify AtC output contour\_integral*
- *fix\_modify AtC mesh output*
- *fix\_modify AtC write\_restart*
- *fix\_modify AtC read\_restart*

*fix\_modify* commands for post-processing:

- *fix\_modify AtC kernel*
- *fix\_modify AtC fields*
- *fix\_modify AtC gradients*
- *fix\_modify AtC rates*
- *fix\_modify AtC computes*
- *fix\_modify AtC on\_the\_fly*
- *fix\_modify AtC pair/bond\_interactions*
- *fix\_modify AtC sample\_frequency*
- *fix\_modify AtC set*

miscellaneous *fix\_modify* commands:

- *fix\_modify AtC atom\_element\_map*
- *fix\_modify AtC atom\_weight*
- *fix\_modify AtC write\_atom\_weights*
- *fix\_modify AtC kernel\_bandwidth*

- *fix\_modify AtC reset\_time*
- *fix\_modify AtC reset\_atomic\_reference\_positions*
- *fix\_modify AtC fe\_md\_boundary*
- *fix\_modify AtC boundary\_faceset*
- *fix\_modify AtC consistent\_fe\_initialization*
- *fix\_modify AtC mass\_matrix*
- *fix\_modify AtC material*
- *fix\_modify AtC atomic\_charge*
- *fix\_modify AtC source\_integration*
- *fix\_modify AtC temperature\_definition*
- *fix\_modify AtC track\_displacement*
- *fix\_modify AtC boundary\_dynamics*
- *fix\_modify AtC add\_species*
- *fix\_modify AtC add\_molecule*
- *fix\_modify AtC remove\_species*
- *fix\_modify AtC remove\_molecule*

Note: a set of example input files with the attendant material files are included in the `examples/PACKAGES/atc` folders.

### 17.10.7 Default

None

---

For detailed exposition of the theory and algorithms please see:

**(Wagner)** Wagner, GJ; Jones, RE; Templeton, JA; Parks, MA, “An atomistic-to-continuum coupling method for heat transfer in solids.” Special Issue of Computer Methods and Applied Mechanics (2008) 197:3351.

**(Zimmerman2004)** Zimmerman, JA; Webb, EB; Hoyt, JJ.; Jones, RE; Klein, PA; Bammann, DJ, “Calculation of stress in atomistic simulation.” Special Issue of Modelling and Simulation in Materials Science and Engineering (2004), 12:S319.

**(Zimmerman2010)** Zimmerman, JA; Jones, RE; Templeton, JA, “A material frame approach for evaluating continuum variables in atomistic simulations.” Journal of Computational Physics (2010), 229:2364.

**(Templeton2010)** Templeton, JA; Jones, RE; Wagner, GJ, “Application of a field-based method to spatially varying thermal transport problems in molecular dynamics.” Modelling and Simulation in Materials Science and Engineering (2010), 18:085007.

**(Jones)** Jones, RE; Templeton, JA; Wagner, GJ; Olmsted, D; Modine, JA, “Electron transport enhanced molecular dynamics for metals and semi-metals.” International Journal for Numerical Methods in Engineering (2010), 83:940.

**(Templeton2011)** Templeton, JA; Jones, RE; Lee, JW; Zimmerman, JA; Wong, BM, “A long-range electric field solver for molecular dynamics based on atomistic-to-continuum modeling.” Journal of Chemical Theory and Computation (2011), 7:1736.

**(Mandadapu)** Mandadapu, KK; Templeton, JA; Lee, JW, “Polarization as a field variable from molecular dynamics simulations.” Journal of Chemical Physics (2013), 139:054115.



Please refer to the standard finite element (FE) texts (e.g., T.J.R. Hughes, *The Finite Element Method*, Dover 2003) for the basics of FE simulations.

## 17.11 fix atom/swap command

### 17.11.1 Syntax

```
fix ID group-ID atom/swap N X seed T keyword values ...
```

- ID, group-ID are documented in *fix* command
- atom/swap = style name of this fix command
- N = invoke this fix every N steps
- X = number of swaps to attempt every N steps
- seed = random # seed (positive integer)
- T = scaling temperature of the MC swaps (temperature units)
- one or more keyword/value pairs may be appended to args
- keyword = *types* or *mu* or *ke* or *semi-grand* or *region*

*types* values = two or more atom types

*mu* values = chemical potential of swap types (energy units)

*ke* value = *no* or *yes*

*no* = no conservation of kinetic energy after atom swaps

*yes* = kinetic energy is conserved after atom swaps

*semi-grand* value = *no* or *yes*

*no* = particle type counts and fractions conserved

*yes* = semi-grand canonical ensemble, particle fractions not conserved

*region* value = region-ID

region-ID = ID of region to use as an exchange/move volume

### 17.11.2 Examples

```
fix 2 all atom/swap 1 1 29494 300.0 ke no types 1 2
fix myFix all atom/swap 100 1 12345 298.0 region my_swap_region types 5 6
fix SGMC all atom/swap 1 100 345 1.0 semi-grand yes types 1 2 3 mu 0.0 4.3 -5.0
```

### 17.11.3 Description

This fix performs Monte Carlo swaps of atoms of one given atom type with atoms of the other given atom types. The specified *T* is used in the Metropolis criterion dictating swap probabilities.

Perform *X* swaps of atoms of one type with atoms of another type according to a Monte Carlo probability. Swap candidates must be in the fix group, must be in the region (if specified), and must be of one of the listed types. Swaps are attempted between candidates that are chosen randomly with equal probability among the candidate atoms. Swaps are not attempted between atoms of the same type since nothing would happen.

All atoms in the simulation domain can be moved using regular time integration displacements (e.g., via *fix nvt*), resulting in a hybrid MC+MD simulation. A smaller-than-usual timestep size may be needed when running such a hybrid simulation, especially if the swapped atoms are not well equilibrated.

The *types* keyword is required. At least two atom types must be specified.

The *ke* keyword can be set to *no* to turn off kinetic energy conservation for swaps. The default is *yes*, which means that swapped atoms have their velocities scaled by the ratio of the masses of the swapped atom types. This ensures that the kinetic energy of each atom is the same after the swap as it was before the swap, even though the atom masses have changed.

The *semi-grand* keyword can be set to *yes* to switch to the semi-grand canonical ensemble as discussed in (*Sadigh*). This means that the total number of each particle type does not need to be conserved. The default is *no*, which means that the only kind of swap allowed exchanges an atom of one type with an atom of a different given type. In other words, the relative mole fractions of the swapped atoms remains constant. Whereas in the semi-grand canonical ensemble, the composition of the system can change. Note that when using *semi-grand*, atoms in the fix group whose type is not listed in the *types* keyword are ineligible for attempted conversion. An attempt is made to switch the selected atom (if eligible) to one of the other listed types with equal probability. Acceptance of each attempt depends upon the Metropolis criterion.

The *mu* keyword allows users to specify chemical potentials. This is required and allowed only when using *semi-grand*. All chemical potentials are absolute, so there is one for each swap type listed following the *types* keyword. In semi-grand canonical ensemble simulations the chemical composition of the system is controlled by the difference in these values. So shifting all values by a constant amount will have no effect on the simulation.

This command may optionally use the *region* keyword to define swap volume. The specified region must have been previously defined with a *region* command. It must be defined with *side = in*. Swap attempts occur only between atoms that are both within the specified region. Swaps are not otherwise attempted.

You should ensure you do not swap atoms belonging to a molecule, or LAMMPS will eventually generate an error when it tries to find those atoms. LAMMPS will warn you if any of the atoms eligible for swapping have a non-zero molecule ID, but does not check for this at the time of swapping.

If not using *semi-grand* this fix checks to ensure all atoms of the given types have the same atomic charge. LAMMPS does not enforce this in general, but it is needed for this fix to simplify the swapping procedure. Successful swaps will swap the atom type and charge of the swapped atoms. Conversely, when using *semi-grand*, it is assumed that all the atom types involved in switches have the same charge. Otherwise, charge would not be conserved. As a consequence, no checks on atomic charges are performed, and successful switches update the atom type but not the atom charge. While it is possible to use *semi-grand* with groups of atoms that have different charges, these charges will not be changed when the atom types change.

Since this fix computes total potential energies before and after proposed swaps, so even complicated potential energy calculations are OK, including the following:

- long-range electrostatics (*k*-space)
- many body pair styles
- hybrid pair styles
- eam pair styles
- triclinic systems
- need to include potential energy contributions from other fixes

Some fixes have an associated potential energy. Examples of such fixes include: *efield*, *gravity*, *addforce*, *langevin*, *restrain*, *temp/berendsen*, *temp/rescale*, and *wall fixes*. For that energy to be included in the total potential energy of the system (the quantity used when performing GCMC moves), you **must** enable the *fix\_modify energy* option for that fix. The doc pages for individual *fix* commands specify if this should be done.

### 17.11.4 Restart, `fix_modify`, `output`, `run start/stop`, `minimize info`

This fix writes the state of the fix to *binary restart files*. This includes information about the random number generator seed, the next timestep for MC exchanges, the number of exchange attempts and successes, etc. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

---

**Note:** For this to work correctly, the timestep must **not** be changed after reading the restart with *reset\_timestep*. The fix will try to detect it and stop with an error.

---

None of the *fix\_modify* options are relevant to this fix.

This fix computes a global vector of length 2, which can be accessed by various *output commands*. The vector values are the following global cumulative quantities:

- 1 = swap attempts
- 2 = swap accepts

The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.11.5 Restrictions

This fix is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

### 17.11.6 Related commands

*fix nvt*, *neighbor*, *fix deposit*, *fix evaporate*, *delete\_atoms*, *fix gcmc*, *fix mol/swap*

### 17.11.7 Default

The option defaults are ke = yes, semi-grand = no, mu = 0.0 for all atom types.

---

(Sadigh) B Sadigh, P Erhart, A Stukowski, A Caro, E Martinez, and L Zepeda-Ruiz, Phys. Rev. B, 85, 184203 (2012).

## 17.12 `fix ave/atom` command

### 17.12.1 Syntax

```
fix ID group-ID ave/atom Nevery Nrepeat Nfreq value1 value2 ...
```

- ID, group-ID are documented in *fix* command
- ave/atom = style name of this fix command
- Nevery = use input values every this many timesteps

- Nrepeat = # of times to use input values for calculating averages
- Nfreq = calculate averages every this many timesteps
- one or more input values can be listed
- value =  $x, y, z, vx, vy, vz, fx, fy, fz, c\_ID, c\_ID[i], f\_ID, f\_ID[i], v\_name$

```
x,y,z,vx,vy,vz,fx,fy,fz = atom attribute (position, velocity, force component)
c_ID = per-atom vector calculated by a compute with ID
c_ID[I] = Ith column of per-atom array calculated by a compute with ID, I can include
↳ wildcard (see below)
f_ID = per-atom vector calculated by a fix with ID
f_ID[I] = Ith column of per-atom array calculated by a fix with ID, I can include
↳ wildcard (see below)
v_name = per-atom vector calculated by an atom-style variable with name
```

## 17.12.2 Examples

```
fix 1 all ave/atom 1 100 100 vx vy vz
fix 1 all ave/atom 10 20 1000 c_my_stress[1]
fix 1 all ave/atom 10 20 1000 c_my_stress[*]
```

## 17.12.3 Description

Use one or more per-atom vectors as inputs every few timesteps, and average them atom by atom over longer timescales. The resulting per-atom averages can be used by other *output commands* such as the *fix ave/chunk* or *dump custom* commands.

The group specified with the command means only atoms within the group have their averages computed. Results are set to 0.0 for atoms not in the group.

Each input value can be an atom attribute (position, velocity, force component) or can be the result of a *compute* or *fix* or the evaluation of an atom-style *variable*. In the latter cases, the compute, fix, or variable must produce a per-atom vector, not a global quantity or local quantity. If you wish to time-average global quantities from a compute, fix, or variable, then see the *fix ave/time* command.

Each per-atom value of each input vector is averaged independently.

*Computes* that produce per-atom vectors or arrays are those which have the word *atom* in their style name. See the doc pages for individual *fixes* to determine which ones produce per-atom vectors or arrays. *Variables* of style *atom* are the only ones that can be used with this fix since they produce per-atom vectors.

Note that for values from a compute or fix, the bracketed index I can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the size of the vector (for *mode* = scalar) or the number of columns in the array (for *mode* = vector), then an asterisk with no numeric values means all indices from 1 to  $N$ . A leading asterisk means all indices from 1 to  $n$  (inclusive). A trailing asterisk means all indices from  $m$  to  $N$  (inclusive). A middle asterisk means all indices from  $m$  to  $n$  (inclusive).

Using a wildcard is the same as if the individual columns of the array had been listed one by one. For example, these two fix *ave/atom* commands are equivalent, since the *compute stress/atom* command creates a per-atom array with six columns:

```
compute my_stress all stress/atom NULL
fix 1 all ave/atom 10 20 1000 c_my_stress[*]
```

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```
fix 1 all ave/atom 10 20 1000 c_my_stress[1] c_my_stress[2] &
      c_my_stress[3] c_my_stress[4] &
      c_my_stress[5] c_my_stress[6]
```

The  $N_{\text{every}}$ ,  $N_{\text{repeat}}$ , and  $N_{\text{freq}}$  arguments specify on what timesteps the input values will be used in order to contribute to the average. The final averaged quantities are generated on timesteps that are a multiple of  $N_{\text{freq}}$ . The average is over  $N_{\text{repeat}}$  quantities, computed in the preceding portion of the simulation every  $N_{\text{every}}$  timesteps.  $N_{\text{freq}}$  must be a multiple of  $N_{\text{every}}$  and  $N_{\text{every}}$  must be non-zero even if  $N_{\text{repeat}}$  is 1. Also, the timesteps contributing to the average value cannot overlap; that is,  $N_{\text{repeat}}N_{\text{every}}$  cannot exceed  $N_{\text{freq}}$ .

For example, if  $N_{\text{every}} = 2$ ,  $N_{\text{repeat}} = 6$ , and  $N_{\text{freq}} = 100$ , then values on timesteps 90, 92, 94, 96, 98, and 100 will be used to compute the final average on time step 100. Similarly for timesteps 190, 192, 194, 196, 198, and 200 on time step 200, etc.

The atom attribute values ( $x$ ,  $y$ ,  $z$ ,  $vx$ ,  $vy$ ,  $vz$ ,  $fx$ ,  $fy$ , and  $fz$ ) are self-explanatory. Note that other atom attributes can be used as inputs to this fix by using the *compute property/atom* command and then specifying an input value from that compute.

**Note:** The  $x$ ,  $y$ , and  $z$  attributes are values that are re-wrapped inside the periodic box whenever an atom crosses a periodic boundary. Thus, if you time-average an atom that spends half of its time on either side of the periodic box, you will get a value in the middle of the box. If this is not what you want, consider averaging unwrapped coordinates, which can be provided by the *compute property/atom* command via its  $xu$ ,  $yu$ , and  $zu$  attributes.

If a value begins with “c\_”, a compute ID must follow which has been previously defined in the input script. If no bracketed term is appended, the per-atom vector calculated by the compute is used. If a bracketed term containing an index  $I$  is appended, the  $I^{\text{th}}$  column of the per-atom array calculated by the compute is used. Users can also write code for their own compute styles and *add them to LAMMPS*. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

If a value begins with “f\_”, a fix ID must follow which has been previously defined in the input script. If no bracketed term is appended, the per-atom vector calculated by the fix is used. If a bracketed term containing an index  $I$  is appended, the  $I^{\text{th}}$  column of the per-atom array calculated by the fix is used. Note that some fixes only produce their values on certain timesteps, which must be compatible with  $N_{\text{every}}$ , else an error will result. Users can also write code for their own fix styles and *add them to LAMMPS*. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

If a value begins with “v\_”, a variable name must follow which has been previously defined in the input script as an *atom-style variable*. Variables of style *atom* can reference thermodynamic keywords or invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of generating per-atom quantities to time average.

## 17.12.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global scalar or vector quantities are stored by this fix for access by various *output commands*.

This fix produces a per-atom vector or array which can be accessed by various *output commands*. A vector is produced if only a single quantity is averaged by this fix. If two or more quantities are averaged, then an array of values is produced. The per-atom values can only be accessed on timesteps that are multiples of  $N_{\text{freq}}$  since that is when averaging is performed.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.12.5 Restrictions

none

## 17.12.6 Related commands

*compute, fix ave/histo, fix ave/chunk, fix ave/time, variable,*

## 17.12.7 Default

none

# 17.13 fix ave/chunk command

## 17.13.1 Syntax

```
fix ID group-ID ave/chunk Nevery Nrepeat Nfreq chunkID value1 value2 ... keyword args ...
```

- ID, group-ID are documented in *fix* command
- ave/chunk = style name of this fix command
- Nevery = use input values every this many timesteps
- Nrepeat = # of times to use input values for calculating averages
- Nfreq = calculate averages every this many timesteps
- chunkID = ID of *compute chunk/atom* command
- one or more input values can be listed
- value = vx, vy, vz, fx, fy, fz, density/mass, density/number, mass, temp, c\_ID, c\_ID[I], f\_ID, f\_ID[I], v\_name

```
vx,vy,vz,fx,fy,fz,mass = atom attribute (velocity, force component, mass)
density/number, density/mass = number or mass density (per volume)
temp = temperature
c_ID = per-atom vector calculated by a compute with ID
c_ID[I] = Ith column of per-atom array calculated by a compute with ID, I can
↔include wildcard (see below)
```

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```
f_ID = per-atom vector calculated by a fix with ID
f_ID[I] = Ith column of per-atom array calculated by a fix with ID, I can include.
↳wildcard (see below)
v_name = per-atom vector calculated by an atom-style variable with name
```

- zero or more keyword/arg pairs may be appended
- keyword = *norm* or *ave* or *bias* or *adof* or *cdof* or *file* or *overwrite* or *title1* or *title2* or *title3*
  - norm* arg = *all* or *sample* or *none* = how output on *Nfreq* steps is normalized
    - all* = output is sum of atoms across all *Nrepeat* samples, divided by atom count
    - sample* = output is sum of *Nrepeat* sample averages, divided by *Nrepeat*
    - none* = output is sum of *Nrepeat* sample sums, divided by *Nrepeat*
  - ave* args = *one* or *running* or *window M*
    - one* = output new average value every *Nfreq* steps
    - running* = output cumulative average of all previous *Nfreq* steps
    - window M* = output average of *M* most recent *Nfreq* steps
  - bias* arg = *bias-ID*
    - bias-ID* = ID of a temperature compute that removes a velocity bias for.
      - ↳temperature calculation
  - adof* value = *dof\_per\_atom*
    - dof\_per\_atom* = define this many degrees-of-freedom per atom for temperature.
      - ↳calculation
  - cdof* value = *dof\_per\_chunk*
    - dof\_per\_chunk* = define this many degrees-of-freedom per chunk for temperature.
      - ↳calculation
  - file* arg = *filename*
    - filename* = file to write results to
  - overwrite* arg = *none* = overwrite output file with only latest output
  - format* arg = *string*
    - string* = C-style format string
  - title1* arg = *string*
    - string* = text to print as 1st line of output file
  - title2* arg = *string*
    - string* = text to print as 2nd line of output file
  - title3* arg = *string*
    - string* = text to print as 3rd line of output file

### 17.13.2 Examples

```
fix 1 all ave/chunk 10000 1 10000 binchunk c_myCentro title1 "My output values"
fix 1 flow ave/chunk 100 10 1000 molchunk vx vz norm sample file vel.profile
fix 1 flow ave/chunk 100 5 1000 binchunk density/mass ave running
fix 1 flow ave/chunk 100 5 1000 binchunk density/mass ave running
```

**Note:** Changed in version 31May2016.

If you are trying to replace a deprecated `fix ave/spatial` command with the newer, more flexible `fix ave/chunk` and `compute chunk/atom` commands, you simply need to split the `fix ave/spatial` arguments across the two new commands. For example, this command:

```
fix 1 flow ave/spatial 100 10 1000 y 0.0 1.0 vx vz norm sample file vel.profile
```

could be replaced by:

```
compute cc1 flow chunk/atom bin/1d y 0.0 1.0
fix 1 flow ave/chunk 100 10 1000 cc1 vx vz norm sample file vel.profile
```

### 17.13.3 Description

Use one or more per-atom vectors as inputs every few timesteps, sum the values over the atoms in each chunk at each timestep, then average the per-chunk values over longer timescales. The resulting chunk averages can be used by other *output commands* such as *thermo\_style custom*, and can also be written to a file.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* page and the *Howto chunk* page for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

Note that only atoms in the specified group contribute to the summing and averaging calculations. The *compute chunk/atom* command defines its own group as well as an optional region. Atoms will have a chunk ID = 0, meaning they belong to no chunk, if they are not in that group or region. Thus you can specify the “all” group for this command if you simply want to use the chunk definitions provided by chunkID.

Each specified per-atom value can be an atom attribute (position, velocity, force component), a number or mass density, a mass or temperature, or the result of a *compute* or *fix* or the evaluation of an atom-style *variable*. In the latter cases, the compute, fix, or variable must produce a per-atom quantity, not a global quantity. Note that the *compute property/atom* command provides access to any attribute defined and stored by atoms. If you wish to time-average global quantities from a compute, fix, or variable, then see the *fix ave/time* command.

The per-atom values of each input vector are summed and averaged independently of the per-atom values in other input vectors.

*Computes* that produce per-atom quantities are those which have the word *atom* in their style name. See the doc pages for individual *fixes* to determine which ones produce per-atom quantities. *Variables* of style *atom* are the only ones that can be used with this fix since all other styles of variable produce global quantities.

Note that for values from a compute or fix, the bracketed index I can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the size of the vector (for *mode* = scalar) or the number of columns in the array (for *mode* = vector), then an asterisk with no numeric values means all indices from 1 to  $N$ . A leading asterisk means all indices from 1 to  $n$  (inclusive). A trailing asterisk means all indices from  $m$  to  $N$  (inclusive). A middle asterisk means all indices from  $m$  to  $n$  (inclusive).

Using a wildcard is the same as if the individual columns of the array had been listed one by one. For example, these two fix ave/chunk commands are equivalent, since the *compute property/atom* command creates, in this case, a per-atom array with three columns:

```
compute myAng all property/atom angmomx angmomy angmomz
fix 1 all ave/chunk 100 1 100 cc1 c_myAng[*] file tmp.angmom
fix 2 all ave/chunk 100 1 100 cc1 c_myAng[1] c_myAng[2] c_myAng[3] file tmp.angmom
```

**Note:** This fix works by creating an array of size  $N_{\text{chunk}} \times N_{\text{values}}$  on each processor.  $N_{\text{chunk}}$  is the number of chunks, which is defined by the *compute chunk/atom* command.  $N_{\text{values}}$  is the number of input values specified. Each processor loops over its atoms, tallying its values to the appropriate chunk. Then the entire array is summed across all processors.



This means that using a large number of chunks will incur an overhead in memory and computational cost (summing across processors), so be careful to define a reasonable number of chunks.

The  $N_{\text{every}}$ ,  $N_{\text{repeat}}$ , and  $N_{\text{freq}}$  arguments specify on what time steps the input values will be accessed and contribute to the average. The final averaged quantities are generated on time steps that are a multiples of  $N_{\text{freq}}$ . The average is over  $N_{\text{repeat}}$  quantities, computed in the preceding portion of the simulation every  $N_{\text{every}}$  time steps.  $N_{\text{freq}}$  must be a multiple of  $N_{\text{every}}$  and  $N_{\text{every}}$  must be non-zero even if  $N_{\text{repeat}} = 1$ . Also, the time steps contributing to the average value cannot overlap (i.e.,  $N_{\text{repeat}}N_{\text{every}}$  cannot exceed  $N_{\text{freq}}$ ).

For example, if  $N_{\text{every}} = 2$ ,  $N_{\text{repeat}} = 6$ , and  $N_{\text{freq}} = 100$ , then values on time steps 90, 92, 94, 96, 98, 100 will be used to compute the final average on time step 100. Similarly for time steps 190, 192, 194, 196, 198, 200 on time step 200, etc. If  $N_{\text{repeat}} = 1$  and  $N_{\text{freq}} = 100$ , then no time averaging is done; values are simply generated on time steps 100, 200, etc.

Each input value can also be averaged over the atoms in each chunk. The way the averaging is done across the  $N_{\text{repeat}}$  time steps to produce output on the  $N_{\text{freq}}$  time steps, and across multiple  $N_{\text{freq}}$  outputs, is determined by the *norm* and *ave* keyword settings, as discussed below.

**Note:** To perform per-chunk averaging within a  $N_{\text{freq}}$  time window, the number of chunks  $N_{\text{chunk}}$  defined by the *compute chunk/atom* command must remain constant. If the *ave* keyword is set to *running* or *window* then  $N_{\text{chunk}}$  must remain constant for the duration of the simulation. This fix forces the chunk/atom compute specified by chunkID to hold  $N_{\text{chunk}}$  constant for the appropriate time windows, by not allowing it to re-calculate  $N_{\text{chunk}}$ , which can also affect how it assigns chunk IDs to atoms. This is particularly important to understand if the chunks defined by the *compute chunk/atom* command are spatial bins. If its *units* keyword is set to *box* or *lattice*, then the number of bins  $N_{\text{chunk}}$  and size of each bin will be fixed over the  $N_{\text{freq}}$  time window, which can affect which atoms are discarded if the simulation box size changes. If its *units* keyword is set to *reduced*, then the number of bins  $N_{\text{chunk}}$  will still be fixed, but the size of each bin can vary at each time step if the simulation box size changes (e.g., for an NPT simulation).

The atom attribute values (*vx*, *vy*, *vz*, *fx*, *fy*, *fz*, *mass*) are self-explanatory. As noted above, any other atom attributes can be used as input values to this fix by using the *compute property/atom* command and then specifying an input value from that compute.

The *density/number* value means the number density is computed for each chunk (i.e., number/volume). The *density/mass* value means the mass density is computed for each chunk (i.e., total-mass/volume). The output values are in units of 1/volume or mass density (mass/volume). See the *units* command page for the definition of density for each choice of units (e.g., g/cm<sup>3</sup>). If the chunks defined by the *compute chunk/atom* command are spatial bins, the volume is the bin volume. Otherwise, it is the volume of the entire simulation box.

The *temp* value means the temperature is computed for each chunk, by the formula

$$\text{KE} = \frac{\text{DOF}}{2} k_B T,$$

where KE is the total kinetic energy of the chunk of atoms (sum of  $\frac{1}{2}mv^2$ ), DOF is the the total number of degrees of freedom for all atoms in the chunk,  $k_B$  is the Boltzmann constant, and  $T$  is the absolute temperature.

The DOF is calculated as  $N*\text{adof} + \text{cdof}$ , where  $N$  is the number of atoms in the chunk, *adof* is the number of degrees of freedom per atom, and *cdof* is the number of degrees of freedom per chunk. By default, *adof* = 2 or 3 = dimensionality of system, as set via the *dimension* command, and *cdof* = 0.0. This gives the usual formula for temperature.

Note that currently this temperature only includes translational degrees of freedom for each atom. No rotational degrees of freedom are included for finite-size particles. Also, no degrees of freedom are subtracted for any velocity bias or constraints that are applied, such as *compute temp/partial*, *fix shake*, or *fix rigid*. This is because those degrees of

freedom (e.g., a constrained bond) could apply to sets of atoms that are both included and excluded from a specific chunk, and hence the concept is somewhat ill-defined. In some cases, you can use the *adof* and *cdof* keywords to adjust the calculated degrees of freedom appropriately, as explained below.

Also note that a bias can be subtracted from atom velocities before they are used in the above formula for KE, by using the *bias* keyword. This allows, for example, a thermal temperature to be computed after removal of a flow velocity profile.

Note that the per-chunk temperature calculated by this fix and the *compute temp/chunk* command can be different. The *compute* calculates the temperature for each chunk for a single snapshot. This fix can do that but can also time average those values over many snapshots, or it can compute a temperature as if the atoms in the chunk on different time steps were collected together as one set of atoms to calculate their temperature. The *compute* allows the center-of-mass velocity of each chunk to be subtracted before calculating the temperature; this fix does not.

If a value begins with “c\_”, a compute ID must follow which has been previously defined in the input script. If no bracketed integer is appended, the per-atom vector calculated by the compute is used. If a bracketed integer is appended, the Ith column of the per-atom array calculated by the compute is used. Users can also write code for their own compute styles and *add them to LAMMPS*. See the discussion above for how I can be specified with a wildcard asterisk to effectively specify multiple values.

If a value begins with “f\_”, a fix ID must follow which has been previously defined in the input script. If no bracketed integer is appended, the per-atom vector calculated by the fix is used. If a bracketed integer is appended, the Ith column of the per-atom array calculated by the fix is used. Note that some fixes only produce their values on certain time steps, which must be compatible with  $N_{\text{every}}$ , else an error results. Users can also write code for their own fix styles and *add them to LAMMPS*. See the discussion above for how I can be specified with a wildcard asterisk to effectively specify multiple values.

If a value begins with “v\_”, a variable name must follow which has been previously defined in the input script. Variables of style *atom* can reference thermodynamic keywords and various per-atom attributes, or invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of generating per-atom quantities to average within chunks.

---

Additional optional keywords also affect the operation of this fix and its outputs.

The *norm* keyword affects how averaging is done for the per-chunk values that are output every  $N_{\text{freq}}$  time steps.

If the *norm* setting is *all*, which is the default, a chunk value is summed over all atoms in all  $N_{\text{repeat}}$  samples, as is the count of atoms in the chunk. The averaged output value for the chunk on the  $N_{\text{freq}}$  time steps is Total-sum / Total-count. In other words it is an average over atoms across the entire  $N_{\text{freq}}$  timescale. For the *density/number* and *density/mass* values, the volume (bin volume or system volume) used in the final normalization will be the volume at the final  $N_{\text{freq}}$  time step. For the *temp* values, degrees of freedom and kinetic energy are summed separately across the entire  $N_{\text{freq}}$  timescale, and the output value is calculated by dividing those two sums.

If the *norm* setting is *sample*, the chunk value is summed over atoms for each sample, as is the count, and an “average sample value” is computed for each sample (i.e., Sample-sum / Sample-count). The output value for the chunk on the  $N_{\text{freq}}$  time steps is the average of the  $N_{\text{repeat}}$  “average sample values” (i.e., the sum of  $N_{\text{repeat}}$  “average sample values” divided by  $N_{\text{repeat}}$ ). In other words, it is an average of an average. For the *density/number* and *density/mass* values, the volume (bin volume or system volume) used in the per-sample normalization will be the current volume at each sampling step.

If the *norm* setting is *none*, a similar computation as for the *sample* setting is done, except the individual “average sample values” are “summed sample values”. A summed sample value is simply the chunk value summed over atoms in the sample, without dividing by the number of atoms in the sample. The output value for the chunk on the  $N_{\text{freq}}$  timesteps is the average of the  $N_{\text{repeat}}$  “summed sample values” (i.e., the sum of  $N_{\text{repeat}}$  “summed sample values” divided by  $N_{\text{repeat}}$ ). For the *density/number* and *density/mass* values, the volume (bin volume or system volume) used in the per-sample sum normalization will be the current volume at each sampling step.

The *ave* keyword determines how the per-chunk values produced every  $N_{\text{freq}}$  steps are averaged with values produced on previous steps that were multiples of  $N_{\text{freq}}$ , before they are accessed by another output command or written to a file.

If the *ave* setting is *one*, which is the default, then the chunk values produced on timesteps that are multiples of  $N_{\text{freq}}$  are independent of each other; they are output as-is without further averaging.

If the *ave* setting is *running*, then the chunk values produced on timesteps that are multiples of  $N_{\text{freq}}$  are summed and averaged in a cumulative sense before being output. Each output chunk value is thus the average of the chunk value produced on that timestep with all preceding values for the same chunk. This running average begins when the fix is defined; it can only be restarted by deleting the fix via the *unfix* command, or re-defining the fix by re-specifying it.

If the *ave* setting is *window*, then the chunk values produced on timesteps that are multiples of  $N_{\text{freq}}$  are summed and averaged within a moving “window” of time, so that the last  $M$  values for the same chunk are used to produce the output. For example, if  $M = 3$  and  $N_{\text{freq}} = 1000$ , then the output on step 10000 will be the average of the individual chunk values on time steps 8000, 9000, and 10000. Outputs on early steps will average over less than  $M$  values if they are not available.

The *bias* keyword specifies the ID of a temperature compute that removes a “bias” velocity from each atom, specified as *bias-ID*. It is only used when the *temp* value is calculated, to compute the thermal temperature of each chunk after the translational kinetic energy components have been altered in a prescribed way (e.g., to remove a flow velocity profile). See the doc pages for individual computes that calculate a temperature to see which ones implement a bias.

The *adof* and *cdof* keywords define the values used in the degree of freedom (DOF) formula described above for temperature calculation for each chunk. They are only used when the *temp* value is calculated. They can be used to calculate a more appropriate temperature for some kinds of chunks. Here are three examples:

If spatially binned chunks contain some number of water molecules and *fix shake* is used to make each molecule rigid, then you could calculate a temperature with six degrees of freedom (DOF) (three translational, three rotational) per molecule by setting *adof* to 2.0.

If *compute temp/partial* is used with the *bias* keyword to only allow the  $x$  component of velocity to contribute to the temperature, then *adof* = 1.0 would be appropriate.

If each chunk consists of a large molecule, with some number of its bonds constrained by *fix shake* or the entire molecule by *fix rigid/small*, *adof* = 0.0 and *cdof* could be set to the remaining degrees of freedom for the entire molecule (entire chunk in this case), that is, 6 for 3d or 3 for 2d for a rigid molecule.

The *file* keyword allows a filename to be specified. Every  $N_{\text{freq}}$  timesteps, a section of chunk info will be written to a text file in the following format. A line with the timestep and number of chunks is written. Then one line per chunk is written, containing the chunk ID ( $1 - N_{\text{chunk}}$ ), an optional original ID value, optional coordinate values for chunks that represent spatial bins, the number of atoms in the chunk, and one or more calculated values. More explanation of the optional values is given below. The number of values in each line corresponds to the number of values specified in the fix *ave/chunk* command. The number of atoms and the value(s) are summed or average quantities, as explained above.

The *overwrite* keyword will continuously overwrite the output file with the latest output, so that it only contains one timestep worth of output. This option can only be used with the *ave running* setting.

The *format* keyword sets the numeric format of each value when it is printed to a file via the *file* keyword. Note that all values are floating point quantities. The default format is %g. You can specify a higher precision if desired (e.g., %20.16g).

The *title1* and *title2* and *title3* keywords allow specification of the strings that will be printed as the first three lines of the output file, assuming the *file* keyword was used. LAMMPS uses default values for each of these, so they do not need to be specified.

By default, these header lines are as follows:

```
# Chunk-averaged data for fix ID and group name
# Timestep Number-of-chunks
# Chunk (OrigID) (Coord1) (Coord2) (Coord3) Ncount value1 value2 ...
```

In the first line, ID and name are replaced with the fix-ID and group name. The second line describes the two values that are printed at the first of each section of output. In the third line the values are replaced with the appropriate value names (e.g., *fx* or *c\_myCompute[2]*).

The words in parenthesis only appear with corresponding columns if the chunk style specified for the *compute chunk/atom* command supports them. The *OrigID* column is only used if the *compress* keyword was set to *yes* for the *compute chunk/atom* command. This means that the original chunk IDs (e.g., molecule IDs) will have been compressed to remove chunk IDs with no atoms assigned to them. Thus a compressed chunk ID of 3 may correspond to an original chunk ID or molecule ID of 415. The *OrigID* column will list 415 for the third chunk.

The *CoordN* columns only appear if a *binning* style was used in the *compute chunk/atom* command. For *bin/1d*, *bin/2d*, and *bin/3d* styles the column values are the center point of the bin in the corresponding dimension. Just *Coord1* is used for *bin/1d*, *Coord2* is added for *bin/2d*, *Coord3* is added for *bin/3d*. For *bin/sphere*, just *Coord1* is used, and it is the radial coordinate. For *bin/cylinder*, *Coord1* and *Coord2* are used. *Coord1* is the radial coordinate (away from the cylinder axis), and *coord2* is the coordinate along the cylinder axis.

Note that if the value of the *units* keyword used in the *compute chunk/atom* command is *box* or *lattice*, the coordinate values will be in distance *units*. If the value of the *units* keyword is *reduced*, the coordinate values will be in unitless reduced units (0–1). This is not true for the *Coord1* value of style *bin/sphere* or *bin/cylinder* which both represent radial dimensions. Those values are always in distance *units*.

---

#### 17.13.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global array of values which can be accessed by various *output commands*. The values can only be accessed on timesteps that are multiples of  $N_{\text{freq}}$ , since that is when averaging is performed. The global array has # of rows = the number of chunks  $N_{\text{chunk}}$ , as calculated by the specified *compute chunk/atom* command. The # of columns is  $M + 1 + N_{\text{values}}$ , where  $M \in \{1, \dots, 4\}$ , depending on whether the optional columns for *OrigID* and *CoordN* are used, as explained above. Following the optional columns, the next column contains the count of atoms in the chunk, and the remaining columns are the *Nvalue* quantities. When the array is accessed with a row *I* that exceeds the current number of chunks, then a 0.0 is returned by the fix instead of an error, since the number of chunks can vary as a simulation runs depending on how that value is computed by the *compute chunk/atom* command.

The array values calculated by this fix are treated as “intensive”, since they are typically already normalized by the count of atoms in each chunk.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.13.5 Restrictions

none

## 17.13.6 Related commands

*compute, fix ave/atom, fix ave/histo, fix ave/time, variable, fix ave/correlate*

## 17.13.7 Default

The option defaults are norm = all, ave = one, bias = none, no file output, and title 1,2,3 = strings as described above.

## 17.14 fix ave/correlate command

### 17.14.1 Syntax

```
fix ID group-ID ave/correlate Nevery Nrepeat Nfreq value1 value2 ... keyword args ...
```

- ID, group-ID are documented in *fix* command
- ave/correlate = style name of this fix command
- Nevery = use input values every this many timesteps
- Nrepeat = # of correlation time windows to accumulate
- Nfreq = calculate time window averages every this many timesteps
- one or more input values can be listed
- value = c\_ID, c\_ID[N], f\_ID, f\_ID[N], v\_name

```
c_ID = global scalar calculated by a compute with ID
c_ID[I] = Ith component of global vector calculated by a compute with ID, I can include wildcard (see below)
f_ID = global scalar calculated by a fix with ID
f_ID[I] = Ith component of global vector calculated by a fix with ID, I can include wildcard (see below)
v_name = global value calculated by an equal-style variable with name
v_name[I] = Ith component of a vector-style variable with name, I can include wildcard (see below)
```

- zero or more keyword/arg pairs may be appended
- keyword = *type* or *ave* or *start* or *prefactor* or *file* or *overwrite* or *title1* or *title2* or *title3*
  - type* arg = *auto* or *upper* or *lower* or *auto/upper* or *auto/lower* or *full*
    - auto* = correlate each value with itself
    - upper* = correlate each value with each succeeding value
    - lower* = correlate each value with each preceding value
    - auto/upper* = auto + upper
    - auto/lower* = auto + lower
    - full* = correlate each value with every other value, including itself = auto + upper + lower
  - ave* args = *one* or *running*
    - one* = zero the correlation accumulation every Nfreq steps
    - running* = accumulate correlations continuously
  - start* args = *Nstart*
    - Nstart* = start accumulating correlations on this timestep

*prefactor* args = value  
value = prefactor to scale all the correlation data by  
*file* arg = filename  
filename = name of file to output correlation data to  
*overwrite* arg = none = overwrite output file with only latest output  
*title1* arg = string  
string = text to print as 1st line of output file  
*title2* arg = string  
string = text to print as 2nd line of output file  
*title3* arg = string  
string = text to print as 3rd line of output file

### 17.14.2 Examples

```
fix 1 all ave/correlate 5 100 1000 c_myTemp file temp.correlate
fix 1 all ave/correlate 1 50 10000 &
      c_thermo_press[1] c_thermo_press[2] c_thermo_press[3] &
      type upper ave running title1 "My correlation data"

fix 1 all ave/correlate 1 50 10000 c_thermo_press[*]
```

### 17.14.3 Description

Use one or more global scalar values as inputs every few timesteps, calculate time correlations between them at varying time intervals, and average the correlation data over longer timescales. The resulting correlation values can be time integrated by *variables* or used by other *output commands* such as *thermo\_style custom*, and can also be written to a file. See the *fix ave/correlate/long* command for an alternate method for computing correlation functions efficiently over very long time windows.

The group specified with this command is ignored. However, note that specified values may represent calculations performed by computes and fixes which store their own “group” definitions.

Each listed value can be the result of a *compute* or *fix* or the evaluation of an equal-style or vector-style *variable*. In each case, the compute, fix, or variable must produce a global quantity, not a per-atom or local quantity. If you wish to spatial- or time-average or histogram per-atom quantities from a compute, fix, or variable, then see the *fix ave/chunk*, *fix ave/atom*, or *fix ave/histo* commands. If you wish to convert a per-atom quantity into a single global value, see the *compute reduce* command.

The input values must be all scalars. What kinds of correlations between input values are calculated is determined by the *type* keyword as discussed below.

*Computes* that produce global quantities are those which do not have the word *atom* in their style name. Only a few *fixes* produce global quantities. See the doc pages for individual fixes for info on which ones produce such values. *Variables* of style *equal* and *vector* are the only ones that can be used with this fix. Variables of style *atom* cannot be used, since they produce per-atom values.

---

For input values from a compute or fix or variable, the bracketed index *I* can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If *N* is the size of the vector, then an asterisk with no numeric values means all indices from 1 to *N*. A leading asterisk means all indices from 1 to *n* (inclusive). A trailing asterisk means all indices from *m* to *N* (inclusive). A middle asterisk means all indices from *m* to *n* (inclusive).

Using a wildcard is the same as if the individual elements of the vector had been listed one by one. For example, the following two `fix ave/correlate` commands are equivalent, since the `compute pressure` command creates a global vector with six values:

```
compute myPress all pressure NULL
fix 1 all ave/correlate 1 50 10000 c_myPress[*]
fix 1 all ave/correlate 1 50 10000 &
      c_myPress[1] c_myPress[2] c_myPress[3] &
      c_myPress[4] c_myPress[5] c_myPress[6]
```

**Note:** For a vector-style variable, only the wildcard forms “\*n” or “m\*n” are allowed. You must specify the upper bound, because vector-style variable lengths are not determined until the variable is evaluated. If n is specified larger than the vector length turns out to be, zeroes are output for missing vector values.

The  $N_{\text{every}}$ ,  $N_{\text{repeat}}$ , and  $N_{\text{freq}}$  arguments specify on what timesteps the input values will be used to calculate correlation data. The input values are sampled every  $N_{\text{every}}$  time steps. The correlation data for the preceding samples is computed on time steps that are a multiple of  $N_{\text{freq}}$ . Consider a set of samples from some initial time up to an output timestep. The initial time could be the beginning of the simulation or the last output time; see the `ave` keyword for options. For the set of samples, the correlation value  $C_{ij}$  is calculated as:

$$C_{ij}(\Delta t) = \langle V_i(t)V_j(t + \Delta t) \rangle,$$

which is the correlation value between input values  $V_i$  and  $V_j$ , separated by time  $\Delta t$ . Note that the second value  $V_j$  in the pair is always the one sampled at the later time. The average is an average over every pair of samples in the set that are separated by time  $\Delta t$ . The maximum  $\Delta t$  used is of size  $(N_{\text{repeat}} - 1)N_{\text{every}}$ . Thus the correlation between a pair of input values yields  $N_{\text{repeat}}$  correlation data:

$$C_{ij}(0), C_{ij}(N_{\text{every}}), C_{ij}(2N_{\text{every}}), \dots, C_{ij}((N_{\text{repeat}} - 1)N_{\text{every}})$$

For example, if  $N_{\text{every}} = 5$ ,  $N_{\text{repeat}} = 6$ , and  $N_{\text{freq}} = 100$ , then values on time steps 0, 5, 10, 15, ..., 100 will be used to compute the final averages on time step 100. Six averages will be computed:  $C_{ij}(0)$ ,  $C_{ij}(5)$ ,  $C_{ij}(10)$ ,  $C_{ij}(15)$ ,  $C_{ij}(20)$ , and  $C_{ij}(25)$ .  $C_{ij}(10)$  on time step 100 will be the average of 19 samples, namely  $V_i(0)V_j(10)$ ,  $V_i(5)V_j(15)$ ,  $V_i(10)V_j(20)$ ,  $V_i(15)V_j(25)$ , ...,  $V_i(85)V_j(95)$ , and  $V_i(90)V_j(100)$ .

$N_{\text{freq}}$  must be a multiple of  $N_{\text{every}}$ ;  $N_{\text{every}}$  and  $N_{\text{repeat}}$  must be non-zero. Also, if the `ave` keyword is set to `one` which is the default, then  $N_{\text{freq}} \geq (N_{\text{repeat}} - 1)N_{\text{every}}$  is required.

If a value begins with “c\_”, a compute ID must follow which has been previously defined in the input script. If no bracketed term is appended, the global scalar calculated by the compute is used. If a bracketed term is appended, the  $I^{\text{th}}$  element of the global vector calculated by the compute is used. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

Note that there is a `compute reduce` command that can sum per-atom quantities into a global scalar or vector which can then be accessed by `fix ave/correlate`. It can also be a compute defined not in your input script, but by `thermodynamic output` or other fixes such as `fix nvt` or `fix temp/rescale`. See the doc pages for these commands which give the IDs of these computes. Users can also write code for their own compute styles and `add them to LAMMPS`.

If a value begins with “f\_”, a fix ID must follow which has been previously defined in the input script. If no bracketed term is appended, the global scalar calculated by the fix is used. If a bracketed term is appended, the  $I^{\text{th}}$  element of the global vector calculated by the fix is used. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

Note that some fixes only produce their values on certain timesteps, which must be compatible with  $N_{\text{every}}$ , else an error will result. Users can also write code for their own fix styles and *add them to LAMMPS*.

If a value begins with “v\_”, a variable name must follow which has been previously defined in the input script. Only equal-style or vector-style variables can be referenced; the latter requires a bracketed term to specify the  $I^{\text{th}}$  element of the vector calculated by the variable. See the *variable* command for details. Note that variables of style *equal* or *vector* define a formula which can reference individual atom properties or thermodynamic keywords, or they can invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of specifying quantities to time correlate.

---

Additional optional keywords also affect the operation of this fix.

The *type* keyword determines which pairs of input values are correlated with each other. For  $N$  input values  $V_i$ , with  $i \in \{1, \dots, N\}$ , let the number of pairs be  $N_{\text{pair}}$ . Note that the second value in the pair,  $V_i(t)V_j(t + \Delta t)$ , is always the one sampled at the later time.

- If *type* is set to *auto* then each input value is correlated with itself (i.e.,  $C_{ii} = V_i^2$  for  $i \in \{1, \dots, N\}$ ), so  $N_{\text{pair}} = N$ .
- If *type* is set to *upper* then each input value is correlated with every succeeding value (i.e.,  $C_{ij} = V_i V_j$  for  $i < j$ , so  $N_{\text{pair}} = N(N - 1)/2$ ).
- If *type* is set to *lower* then each input value is correlated with every preceding value (i.e.,  $C_{ij} = V_i V_j$  for  $i > j$ , so  $N_{\text{pair}} = N(N - 1)/2$ ).
- If *type* is set to *auto/upper* then each input value is correlated with itself and every succeeding value (i.e.,  $C_{ij} = V_i V_j$  for  $i \geq j$ , so  $N_{\text{pair}} = N(N + 1)/2$ ).
- If *type* is set to *auto/lower* then each input value is correlated with itself and every preceding value (i.e.,  $C_{ij} = V_i V_j$  for  $i \leq j$ , so  $N_{\text{pair}} = N(N + 1)/2$ ).
- If *type* is set to *full* then each input value is correlated with itself and every other value (i.e.,  $C_{ij} = V_i V_j$  for  $\{i, j\} = \{1, N\}$ , so  $N_{\text{pair}} = N^2$ ).

The *ave* keyword determines what happens to the accumulation of correlation samples every  $N_{\text{freq}}$  timesteps. If the *ave* setting is *one*, then the accumulation is restarted or zeroed every  $N_{\text{freq}}$  timesteps. Thus the outputs on successive  $N_{\text{freq}}$  timesteps are essentially independent of each other. The exception is that the  $C_{ij}(0) = V_i(t)V_j(t)$  value at a time step  $t$ , where  $t$  is a multiple of  $N_{\text{freq}}$ , contributes to the correlation output both at time  $t$  and at time  $t + N_{\text{freq}}$ .

If the *ave* setting is *running*, then the accumulation is never zeroed. Thus the output of correlation data at any timestep is the average over samples accumulated every  $N_{\text{every}}$  steps since the fix was defined. It can only be restarted by deleting the fix via the *unfix* command, or by re-defining the fix by re-specifying it.

The *start* keyword specifies what time step the accumulation of correlation samples will begin on. The default is step 0. Setting it to a larger value can avoid adding non-equilibrated data to the correlation averages.

The *prefactor* keyword specifies a constant which will be used as a multiplier on the correlation data after it is averaged. It is effectively a scale factor on  $V_i V_j$ , which can be used to account for the size of the time window or other unit conversions.

The *file* keyword allows a filename to be specified. Every  $N_{\text{freq}}$  steps, an array of correlation data is written to the file. The number of rows is  $N_{\text{repeat}}$ , as described above. The number of columns is  $N_{\text{pair}} + 2$ , also as described above. Thus the file ends up to be a series of these array sections.

The *overwrite* keyword will continuously overwrite the output file with the latest output, so that it only contains one timestep worth of output. This option can only be used with the *ave running* setting.

The *title1*, *title2*, and *title3* keywords allow specification of the strings that will be printed as the first three lines of the output file, assuming the *file* keyword was used. LAMMPS uses default values for each of these, so they do not need to be specified.



By default, these header lines are as follows:

```
# Time-correlated data for fix ID
# TimeStep Number-of-time-windows
# Index TimeDelta Ncount valueI*valueJ valueI*valueJ ...
```

In the first line, ID is replaced with the fix-ID. The second line describes the two values that are printed at the first of each section of output. In the third line the value pairs are replaced with the appropriate fields from the fix ave/correlate command.

Let  $S_{ij}$  be a set of time correlation data for input values  $I$  and  $J$ , namely the  $N_{\text{repeat}}$  values:

$$S_{ij} = C_{ij}(0), C_{ij}(N_{\text{every}}), C_{ij}(2N_{\text{every}}), \dots, C_{ij}((N_{\text{repeat}} - 1)N_{\text{every}})$$

As explained below, these data are output as one column of a global array, which is effectively the correlation matrix.

The *trap* function defined for *equal-style variables* can be used to perform a time integration of this vector of data, using a trapezoidal rule. This is useful for calculating various quantities which can be derived from time correlation data. If a normalization factor is needed for the time integration, it can be included in the variable formula or via the *prefactor* keyword.

#### 17.14.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global array of values which can be accessed by various *output commands*. The values can only be accessed on timesteps that are multiples of  $N_{\text{freq}}$  since that is when averaging is performed. The global array has # of rows  $N_{\text{repeat}}$  and # of columns  $N_{\text{pair}} + 2$ . The first column has the time  $\Delta t$  (in time steps) between the pairs of input values used to calculate the correlation, as described above. The second column has the number of samples contributing to the correlation average, as described above. The remaining  $N_{\text{pair}}$  columns are for  $I, J$  pairs of the  $N$  input values, as determined by the *type* keyword, as described above.

- For *type = auto*, the  $N_{\text{pair}} = N$  columns are ordered:  $C_{11}, C_{22}, \dots, C_{NN}$
- For *type = upper*, the  $N_{\text{pair}} = N(N - 1)/2$  columns are ordered:  $C_{12}, C_{13}, \dots, C_{1N}, C_{23}, \dots, C_{2N}, C_{34}, \dots, C_{N-1,N}$
- For *type = lower*, the  $N_{\text{pair}} = N(N - 1)/2$  columns are ordered:  $C_{21}, C_{31}, C_{32}, C_{41}, C_{42}, C_{43}, \dots, C_{N1}, C_{N2}, \dots, C_{N,N-1}$
- For *type = auto/upper*, the  $N_{\text{pair}} = N(N + 1)/2$  columns are ordered:  $C_{11}, C_{12}, C_{13}, \dots, C_{1N}, C_{22}, C_{23}, \dots, C_{2N}, C_{33}, C_{34}, \dots, C_{N-1,N}, C_{NN}$
- For *type = auto/lower*, the  $N_{\text{pair}} = N(N + 1)/2$  columns are ordered:  $C_{11}, C_{21}, C_{22}, C_{31}, C_{32}, C_{33}, C_{41}, \dots, C_{44}, C_{N1}, C_{N2}, \dots, C_{N,N-1}, C_{NN}$
- For *type = full*, the  $N_{\text{pair}} = N^2$  columns are ordered:  $C_{11}, C_{12}, \dots, C_{1N}, C_{21}, C_{22}, \dots, C_{2N}, C_{31}, \dots, C_{3N}, \dots, C_{N1}, \dots, C_{N,N-1}, C_{NN}$

The array values calculated by this fix are treated as extensive. If you need to divide them by the number of atoms, you must do this in a later processing step (e.g., when using them in a *variable*).

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.14.5 Restrictions

none

### 17.14.6 Related commands

*fix ave/correlate/long, compute, fix ave/time, fix ave/atom, fix ave/chunk, fix ave/histo, variable*

### 17.14.7 Default

none

The option defaults are ave = one, type = auto, start = 0, no file output, title 1,2,3 = strings as described above, and prefactor = 1.0.

## 17.15 fix ave/correlate/long command

### 17.15.1 Syntax

```
fix ID group-ID ave/correlate/long Nevery Nfreq value1 value2 ... keyword args ...
```

- ID, group-ID are documented in *fix* command
- ave/correlate/long = style name of this fix command
- Nevery = use input values every this many time steps
- Nfreq = save state of the time correlation functions every this many time steps
- one or more input values can be listed
- value = c\_ID, c\_ID[N], f\_ID, f\_ID[N], v\_name

```
c_ID = global scalar calculated by a compute with ID
c_ID[I] = Ith component of global vector calculated by a compute with ID
f_ID = global scalar calculated by a fix with ID
f_ID[I] = Ith component of global vector calculated by a fix with ID
v_name = global value calculated by an equal-style variable with name
```

- zero or more keyword/arg pairs may be appended
- keyword = *type* or *start* or *file* or *overwrite* or *title1* or *title2* or *ncorr* or *nlen* or *ncount*  
*type* arg = *auto* or *upper* or *lower* or *auto/upper* or *auto/lower* or *full*  
auto = correlate each value with itself  
upper = correlate each value with each succeeding value  
lower = correlate each value with each preceding value  
auto/upper = auto + upper  
auto/lower = auto + lower  
full = correlate each value with every other value, including itself = auto +  
↔upper + lower  
*start* args = Nstart  
Nstart = start accumulating correlations on this time step  
*file* arg = filename

```

filename = name of file to output correlation data to
overwrite arg = none = overwrite output file with only latest output
title1 arg = string
  string = text to print as 1st line of output file
title2 arg = string
  string = text to print as 2nd line of output file
ncorr arg = Ncorrelators
  Ncorrelators = number of correlators to store
nlen args = Nlen
  Nlen = length of each correlator
ncount args = Ncount
  Ncount = number of values over which successive correlators are averaged

```

### 17.15.2 Examples

```

fix 1 all ave/correlate/long 5 1000 c_myTemp file temp.correlate
fix 1 all ave/correlate/long 1 10000 &
      c_thermo_press[1] c_thermo_press[2] c_thermo_press[3] &
      type upper title "My correlation data" nlen 15 ncount 3

```

### 17.15.3 Description

This fix is similar in spirit and syntax to the *fix ave/correlate*. However, this fix allows the efficient calculation of time correlation functions on-the-fly over extremely long time windows with little additional CPU overhead, using a multiple- $\tau$  method (*Ramirez*) that decreases the resolution of the stored correlation function with time. It is not a full drop-in replacement.

The group specified with this command is ignored. However, note that specified values may represent calculations performed by computes and fixes which store their own “group” definitions.

Each listed value can be the result of a compute or fix or the evaluation of an equal-style variable. See the *fix ave/correlate* page for details.

The *Nevery* and *Nfreq* arguments specify on what time steps the input values will be used to calculate correlation data and the frequency with which the time correlation functions will be output to a file, respectively. Note that there is no *Nrepeat* argument, unlike the *fix ave/correlate* command.

The optional keywords *ncorr*, *nlen*, and *ncount* are unique to this command and determine the number of correlation points calculated and the memory and CPU overhead used by this calculation. *Nlen* and *ncount* determine the amount of averaging done at longer correlation times. The default values *nlen* = 16 and *ncount* = 2 ensure that the systematic error of the multiple- $\tau$  correlator is always below the level of the statistical error of a typical simulation (which depends on the ensemble size and the simulation length).

The maximum correlation time (in time steps) that can be reached is given by the formula  $(nlen - 1)ncount^{(ncorr-1)}$ . Longer correlation times are discarded and not calculated. With the default values of the parameters (*ncorr* = 20, *nlen* = 16 and *ncount* = 2), this corresponds to 7864320 time steps. If longer correlation times are needed, the value of *ncorr* should be increased. Using *nlen* = 16 and *ncount* = 2, with *ncorr* = 30, the maximum number of steps that can be correlated is 80530636808. If *ncorr* = 40, correlation times in excess of  $8 \times 10^{12}$  time steps can be calculated.

The total memory needed for each correlation pair is roughly  $4 \times ncorr \times nlen \times 8$  bytes. With the default values of the parameters, this corresponds to about 10 KB.

For the meaning of the additional optional keywords, see the *fix ave/correlate* doc page.

### 17.15.4 Restart, fix\_modify, output, run start/stop, minimize info

Contrary to *fix ave/correlate* this fix does **not** provide access to its internal data to various output options. Since this fix is intended for the calculation of time correlation functions over very long MD simulations, the information about this fix is written automatically to binary restart files, so that the time correlation calculation can continue in subsequent simulations. None of the *fix\_modify* options are relevant to this fix.

No parameter of this fix can be used with the start/stop keywords of the run command. This fix is not invoked during energy minimization.

### 17.15.5 Restrictions

This compute is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.15.6 Related commands

*fix ave/correlate*

### 17.15.7 Default

none

The option defaults for keywords that are also keywords for the *fix ave/correlate* command are as follows: type = auto, start = 0, no file output, title 1,2 = strings as described on the *fix ave/correlate* doc page.

The option defaults for keywords unique to this command are as follows: ncorr=20, nlen=16, ncount=2.

---

(Ramirez) J. Ramirez, S.K. Sukumaran, B. Vorselaars and A.E. Likhtman, J. Chem. Phys. 133, 154103 (2010).

## 17.16 fix ave/histo command

## 17.17 fix ave/histo/weight command

### 17.17.1 Syntax

```
fix ID group-ID style Nevery Nrepeat Nfreq lo hi Nbin value1 value2 ... keyword args ...
```

- ID, group-ID are documented in *fix* command
- style = *ave/histo* or *ave/histo/weight* = style name of this fix command
- Nevery = use input values every this many timesteps
- Nrepeat = # of times to use input values for calculating histogram
- Nfreq = calculate histogram every this many timesteps
- lo,hi = lo/hi bounds within which to histogram
- Nbin = # of histogram bins

- one or more input values can be listed
- value =  $x, y, z, vx, vy, vz, fx, fy, fz, c\_ID, c\_ID[N], f\_ID, f\_ID[N], v\_name$

$x, y, z, vx, vy, vz, fx, fy, fz$  = atom attribute (position, velocity, force component)  
 $c\_ID$  = scalar or vector calculated by a compute with ID  
 $c\_ID[I]$  = Ith component of vector or Ith column of array calculated by a compute.  
 ↪with ID, I can include wildcard (see below)  
 $f\_ID$  = scalar or vector calculated by a fix with ID  
 $f\_ID[I]$  = Ith component of vector or Ith column of array calculated by a fix with.  
 ↪ID, I can include wildcard (see below)  
 $v\_name$  = value(s) calculated by an equal-style or vector-style or atom-style.  
 ↪variable with name  
 $v\_name[I]$  = value calculated by a vector-style variable with name, I can include.  
 ↪wildcard (see below)

- zero or more keyword/arg pairs may be appended
- keyword = *mode* or *file* or *ave* or *start* or *beyond* or *overwrite* or *title1* or *title2* or *title3*

*mode* arg = *scalar* or *vector*

scalar = all input values are scalars

vector = all input values are vectors

*kind* arg = *global* or *peratom* or *local*

*file* arg = filename

filename = name of file to output histogram(s) to

*ave* args = *one* or *running* or *window*

one = output a new average value every Nfreq steps

running = output cumulative average of all previous Nfreq steps

window M = output average of M most recent Nfreq steps

*start* args = Nstart

Nstart = start averaging on this timestep

*beyond* arg = *ignore* or *end* or *extra*

ignore = ignore values outside histogram lo/hi bounds

end = count values outside histogram lo/hi bounds in end bins

extra = create 2 extra bins for value outside histogram lo/hi bounds

*overwrite* arg = none = overwrite output file with only latest output

*title1* arg = string

string = text to print as 1st line of output file

*title2* arg = string

string = text to print as 2nd line of output file

*title3* arg = string

string = text to print as 3rd line of output file, only for vector mode

## 17.17.2 Examples

```
fix 1 all ave/histo 100 5 1000 0.5 1.5 50 c_myTemp file temp.histo ave running
fix 1 all ave/histo 100 5 1000 -5 5 100 c_thermo_press[2] c_thermo_press[3] title1 "My_
↪output values"
fix 1 all ave/histo 100 5 1000 -5 5 100 c_thermo_press[*]
fix 1 all ave/histo 1 100 1000 -2.0 2.0 18 vx vy vz mode vector ave running beyond extra
fix 1 all ave/histo/weight 1 1 1 10 100 2000 c_XRD[1] c_XRD[2]
```

### 17.17.3 Description

Use one or more values as inputs every few timesteps to create a single histogram. The histogram can then be averaged over longer timescales. The resulting histogram can be used by other *output commands*, and can also be written to a file. The `fix ave/histo/weight` command has identical syntax to `fix ave/histo`, except that exactly two values must be specified. See details below.

The group specified with this command is ignored for global and local input values. For per-atom input values, only atoms in the group contribute to the histogram. Note that regardless of the specified group, specified values may represent calculations performed by `compute`s and `fix`s which store their own “group” definition.

A histogram is simply a count of the number of values that fall within a histogram bin. *Nbins* are defined, with even spacing between *lo* and *hi*. Values that fall outside the *lo/hi* bounds can be treated in different ways; see the discussion of the *beyond* keyword below.

Each input value can be an atom attribute (position, velocity, force component) or can be the result of a *compute* or *fix* or the evaluation of an equal-style or vector-style or atom-style *variable*. The set of input values can be either all global, all per-atom, or all local quantities. Inputs of different kinds (e.g. global and per-atom) cannot be mixed. Atom attributes are per-atom vector values. See the page for individual “compute” and “fix” commands to see what kinds of quantities they generate. See the optional *kind* keyword below for how to force the `fix ave/histo` command to disambiguate if necessary.

Note that the output of this command is a single histogram for all input values combined together, not one histogram per input value. See below for details on the format of the output of this fix.

The input values must either be all scalars or all vectors (or arrays), depending on the setting of the *mode* keyword.

If *mode* = scalar, then the input values must be scalars, or vectors with a bracketed term appended, indicating the *I*th value of the vector is used.

If *mode* = vector, then the input values must be vectors, or arrays with a bracketed term appended, indicating the *I*th column of the array is used.

If the `fix ave/histo/weight` command is used, exactly two values must be specified. If the values are vectors, they must be the same length. The first value (a scalar or vector) is what is histogrammed into bins, in the same manner the `fix ave/histo` command operates. The second value (a scalar or vector) is used as a “weight”. This means that instead of each value tallying a “1” to its bin, the corresponding weight is tallied. For example, the  $N^{\text{th}}$  entry (weight) in the second vector is tallied to the bin corresponding to the  $N^{\text{th}}$  entry in the first vector.

---

For input values from a `compute` or `fix` or *variable*, the bracketed index *I* can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If *N* is the size of the vector (for *mode* = scalar) or the number of columns in the array (for *mode* = vector), then an asterisk with no numeric values means all indices from 1 to *N*. A leading asterisk means all indices from 1 to *n* (inclusive). A trailing asterisk means all indices from *m* to *N* (inclusive). A middle asterisk means all indices from *m* to *n* (inclusive).

Using a wildcard is the same as if the individual elements of the vector or columns of the array had been listed one by one. For example, the following two `fix ave/histo` commands are equivalent, since the *compute com/chunk* command creates a global array with three columns:

```
compute myCOM all com/chunk
fix 1 all ave/histo 100 1 100 -10.0 10.0 100 c_myCOM[*] file tmp1.com mode vector
fix 2 all ave/histo 100 1 100 -10.0 10.0 100 c_myCOM[1] c_myCOM[2] c_myCOM[3] file tmp2.
↪com mode vector
```

---

**Note:** For a vector-style variable, only the wildcard forms “\*n” or “m\*n” are allowed. You must specify the upper bound, because vector-style variable lengths are not determined until the variable is evaluated. If *n* is specified larger

than the vector length turns out to be, zeroes are output for missing vector values.

The  $N_{\text{every}}$ ,  $N_{\text{repeat}}$ , and  $N_{\text{freq}}$  arguments specify on what time steps the input values will be used in order to contribute to the histogram. The final histogram is generated on time steps that are multiple of  $N_{\text{freq}}$ . It is averaged over  $N_{\text{repeat}}$  histograms, computed in the preceding portion of the simulation every  $N_{\text{every}}$  time steps.  $N_{\text{freq}}$  must be a multiple of  $N_{\text{every}}$  and  $N_{\text{every}}$  must be non-zero even if  $N_{\text{repeat}}$  is 1. Also, the time steps contributing to the histogram value cannot overlap (i.e.,  $N_{\text{repeat}} \times N_{\text{every}}$  cannot exceed  $N_{\text{freq}}$ ).

For example, if  $N_{\text{every}} = 2$ ,  $N_{\text{repeat}} = 6$ , and  $N_{\text{freq}} = 100$ , then input values on time steps 90, 92, 94, 96, 98, and 100 will be used to compute the final histogram on timestep 100. Similarly for timesteps 190, 192, 194, 196, 198, and 200 on timestep 200, etc. If  $N_{\text{repeat}} = 1$  and  $N_{\text{freq}} = 100$ , then no time averaging of the histogram is done; a histogram is simply generated on timesteps 100, 200, etc.

The atom attribute values ( $x$ ,  $y$ ,  $z$ ,  $vx$ ,  $vy$ ,  $vz$ ,  $fx$ ,  $fy$ , and  $fz$ ) are self-explanatory. Note that other atom attributes can be used as inputs to this fix by using the `compute property/atom` command and then specifying an input value from that compute.

If a value begins with “c\_”, a compute ID must follow which has been previously defined in the input script. If *mode* = scalar, then if no bracketed term is appended, the global scalar calculated by the compute is used. If a bracketed term is appended, the  $I$ th element of the global vector calculated by the compute is used. If *mode* = vector, then if no bracketed term is appended, the global or per-atom or local vector calculated by the compute is used. If a bracketed term is appended, the  $I$ th column of the global or per-atom or local array calculated by the compute is used. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

Note that there is a `compute reduce` command that can sum per-atom quantities into a global scalar or vector, which can then be accessed by fix ave/histo. It can also be a compute defined not in your input script, but by `thermodynamic output` or other fixes such as `fix nvt` or `fix temp/rescale`. See the doc pages for these commands which give the IDs of these computes. Users can also write code for their own compute styles and `add them to LAMMPS`.

If a value begins with “f\_”, a fix ID must follow which has been previously defined in the input script. If *mode* = scalar, then if no bracketed term is appended, the global scalar calculated by the fix is used. If a bracketed term is appended, the  $I$ th element of the global vector calculated by the fix is used. If *mode* = vector, then if no bracketed term is appended, the global or per-atom or local vector calculated by the fix is used. If a bracketed term is appended, the  $I$ th column of the global or per-atom or local array calculated by the fix is used. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

Note that some fixes only produce their values on certain timesteps, which must be compatible with  $N_{\text{every}}$ , else an error will result. Users can also write code for their own fix styles and `add them to LAMMPS`.

If a value begins with “v\_”, a variable name must follow which has been previously defined in the input script. If *mode* = scalar, then only equal-style or vector-style variables can be used, which both produce global values. In this mode, a vector-style variable requires a bracketed term to specify the  $I$ th element of the vector calculated by the variable. If *mode* = vector, then only vector-style or atom-style variables can be used, which produce a global or per-atom vector respectively. The vector-style variable must be used without a bracketed term. See the `variable` command for details.

Note that variables of style `equal`, `vector`, and `atom` define a formula which can reference individual atom properties or thermodynamic keywords, or they can invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of specifying quantities to histogram.

Additional optional keywords also affect the operation of this fix.

If the *mode* keyword is set to `scalar`, then all input values must be global scalars, or elements of global vectors. If the *mode* keyword is set to `vector`, then all input values must be global or per-atom or local vectors, or columns of global or per-atom or local arrays.

The *kind* keyword only needs to be set if a compute or fix produces more than one kind of output (global, per-atom, local). If this is not the case, then LAMMPS will determine what kind of input is provided and whether all the input arguments are consistent. If a compute or fix produces more than one kind of output, the *kind* keyword should be used to specify which output will be used. The remaining input arguments must still be consistent.

The *beyond* keyword determines how input values that fall outside the *lo* to *hi* bounds are treated. Values such that  $lo \leq \text{value} \leq hi$  are assigned to one bin. Values on a bin boundary are assigned to the lower of the two bins. If *beyond* is set to *ignore* then values  $< lo$  and values  $> hi$  are ignored (i.e., they are not binned). If *beyond* is set to *end*, then values  $< lo$  are counted in the first bin and values  $> hi$  are counted in the last bin. If *beyond* is set to *extend*, then two extra bins are created so that there are  $N_{\text{bins}} + 2$  total bins. Values  $< lo$  are counted in the first bin and values  $> hi$  are counted in the last bin ( $N_{\text{bins}} + 2$ ). Values between *lo* and *hi* (inclusive) are counted in bins 2 through  $N_{\text{bins}} + 1$ . The “coordinate” stored and printed for these two extra bins is *lo* and *hi*.

The *ave* keyword determines how the histogram produced every  $N_{\text{freq}}$  steps are averaged with histograms produced on previous steps that were multiples of  $N_{\text{freq}}$ , before they are accessed by another output command or written to a file.

If the *ave* setting is *one*, then the histograms produced on timesteps that are multiples of  $N_{\text{freq}}$  are independent of each other; they are output as-is without further averaging.

If the *ave* setting is *running*, then the histograms produced on timesteps that are multiples of  $N_{\text{freq}}$  are summed and averaged in a cumulative sense before being output. Each bin value in the histogram is thus the average of the bin value produced on that timestep with all preceding values for the same bin. This running average begins when the fix is defined; it can only be restarted by deleting the fix via the *unfix* command, or by re-defining the fix by re-specifying it.

If the *ave* setting is *window*, then the histograms produced on timesteps that are multiples of  $N_{\text{freq}}$  are summed within a moving “window” of time, so that the last  $M$  histograms are used to produce the output (e.g., if  $M = 3$  and  $N_{\text{freq}} = 1000$ , then the output on step 10000 will be the combined histogram of the individual histograms on steps 8000, 9000, and 10000. Outputs on early steps will be sums over less than  $M$  histograms if they are not available.

The *start* keyword specifies what timestep histogramming will begin on. The default is step 0. Often input values can be 0.0 at time 0, so setting *start* to a larger value can avoid including a 0.0 in a running or windowed histogram.

The *file* keyword allows a filename to be specified. Every  $N_{\text{freq}}$  steps, one histogram is written to the file. This includes a leading line that contains the timestep, number of bins, the total count of values contributing to the histogram, the count of values that were not histogrammed (see the *beyond* keyword), the minimum value encountered, and the maximum value encountered. The min/max values include values that were not histogrammed. Following the leading line, one line per bin is written into the file. Each line contains the bin #, the coordinate for the center of the bin (between *lo* and *hi*), the count of values in the bin, and the normalized count. The normalized count is the bin count divided by the total count (not including values not histogrammed), so that the normalized values sum to 1.0 across all bins.

The *overwrite* keyword will continuously overwrite the output file with the latest output, so that it only contains one timestep worth of output. This option can only be used with the *ave running* setting.

The *title1*, *title2*, and *title3* keywords allow specification of the strings that will be printed as the first three lines of the output file, assuming the *file* keyword was used. LAMMPS uses default values for each of these, so they do not need to be specified.

By default, these header lines are as follows:

```
# Histogram for fix ID
# TimeStep Number-of-bins Total-counts Missing-counts Min-value Max-value
# Bin Coord Count Count/Total
```

In the first line, ID is replaced with the fix-ID. The second line describes the six values that are printed at the first of each section of output. The third describes the four values printed for each bin in the histogram.



### 17.17.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix produces a global vector and global array which can be accessed by various *output commands*. The values can only be accessed on timesteps that are multiples of  $N_{\text{freq}}$  since that is when a histogram is generated. The global vector has four values:

- 1 = total counts in the histogram
- 2 = values that were not histogrammed (see *beyond* keyword)
- 3 = min value of all input values, including ones not histogrammed
- 4 = max value of all input values, including ones not histogrammed

The global array has  $N_{\text{bins}}$  rows and three columns. The first column has the bin coordinate, the second column has the count of values in that histogram bin, and the third column has the bin count divided by the total count (not including missing counts), so that the values in the third column sum to 1.0.

The vector and array values calculated by this fix are all treated as intensive. If this is not the case (e.g., due to histogramming per-atom input values), then you will need to account for that when interpreting the values produced by this fix.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.17.5 Restrictions

none

### 17.17.6 Related commands

*compute*, *fix ave/atom*, *fix ave/chunk*, *fix ave/time*, *variable*, *fix ave/correlate*,

### 17.17.7 Default

none

The option defaults are mode = scalar, kind = figured out from input arguments, ave = one, start = 0, no file output, beyond = ignore, and title 1,2,3 = strings as described above.

## 17.18 fix ave/time command

### 17.18.1 Syntax

```
fix ID group-ID ave/time Nevery Nrepeat Nfreq value1 value2 ... keyword args ...
```

- ID, group-ID are documented in *fix* command
- ave/time = style name of this fix command
- Nevery = use input values every this many time steps
- Nrepeat = # of times to use input values for calculating averages

- Nfreq = calculate averages every this many time steps
- one or more input values can be listed
- value = c\_ID, c\_ID[N], f\_ID, f\_ID[N], v\_name

```
c_ID = global scalar or vector calculated by a compute with ID
c_ID[I] = Ith component of global vector or Ith column of global array calculated.
↳by a compute with ID, I can include wildcard (see below)
f_ID = global scalar or vector calculated by a fix with ID
f_ID[I] = Ith component of global vector or Ith column of global array calculated.
↳by a fix with ID, I can include wildcard (see below)
v_name = value(s) calculated by an equal-style or vector-style variable with name
v_name[I] = value calculated by a vector-style variable with name, I can include.
↳wildcard (see below)
```

- zero or more keyword/arg pairs may be appended
- keyword = *mode* or *file* or *ave* or *start* or *off* or *overwrite* or *title1* or *title2* or *title3*
  - mode* arg = *scalar* or *vector*
    - scalar* = all input values are global scalars
    - vector* = all input values are global vectors or global arrays
  - ave* args = *one* or *running* or *window M*
    - one* = output a new average value every Nfreq steps
    - running* = output cumulative average of all previous Nfreq steps
    - window M* = output average of M most recent Nfreq steps
  - start* args = *Nstart*
    - Nstart* = start averaging on this time step
  - off* arg = *M* = do not average this value
    - M* = value # from 1 to Nvalues
  - file* arg = *filename*
    - filename* = name of file to output time averages to
  - overwrite* arg = *none* = overwrite output file with only latest output
  - format* arg = *string*
    - string* = C-style format string
  - title1* arg = *string*
    - string* = text to print as 1st line of output file
  - title2* arg = *string*
    - string* = text to print as 2nd line of output file
  - title3* arg = *string*
    - string* = text to print as 3rd line of output file, only for vector mode

## 17.18.2 Examples

```
fix 1 all ave/time 100 5 1000 c_myTemp c_thermo_temp file temp.profile
fix 1 all ave/time 100 5 1000 c_thermo_press[2] ave window 20 &
      title1 "My output values"
fix 1 all ave/time 100 5 1000 c_thermo_press[*]
fix 1 all ave/time 1 100 1000 f_indent f_indent[1] file temp.indent off 1
```

### 17.18.3 Description

Use one or more global values as inputs every few time steps, and average them over longer timescales. The resulting averages can be used by other *output commands* such as *thermo\_style custom*, and can also be written to a file. Note that if no time averaging is done, this command can be used as a convenient way to simply output one or more global values to a file.

The group specified with this command is ignored. However, note that specified values may represent calculations performed by computes and fixes which store their own “group” definitions.

Each listed value can be the result of a *compute* or *fix* or the evaluation of an equal-style or vector-style *variable*. In each case, the compute, fix, or variable must produce a global quantity, not a per-atom or local quantity. If you wish to spatial- or time-average or histogram per-atom quantities from a compute, fix, or variable, then see the *fix ave/chunk*, *fix ave/atom*, or *fix ave/histo* commands. If you wish to sum a per-atom quantity into a single global quantity, see the *compute reduce* command.

*Computes* that produce global quantities are those which do not have the word *atom* in their style name. Only a few *fixes* produce global quantities. See the doc pages for individual fixes for info on which ones produce such values. *Variables* of style *equal* and *vector* are the only ones that can be used with this fix. Variables of style *atom* cannot be used, since they produce per-atom values.

The input values must either be all scalars or all vectors depending on the setting of the *mode* keyword. In both cases, the averaging is performed independently on each input value (i.e., each input scalar is averaged independently or each element of each input vector is averaged independently).

If *mode* = scalar, then the input values must be scalars, or vectors with a bracketed term appended, indicating the  $I^{\text{th}}$  value of the vector is used.

If *mode* = vector, then the input values must be vectors, or arrays with a bracketed term appended, indicating the  $I^{\text{th}}$  column of the array is used. All vectors must be the same length, which is the length of the vector or number of rows in the array.

For input values from a compute or fix or variable, the bracketed index  $I$  can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the size of the vector (for *mode* = scalar) or the number of columns in the array (for *mode* = vector), then an asterisk with no numeric values means all indices from 1 to  $N$ . A leading asterisk means all indices from 1 to  $n$  (inclusive). A trailing asterisk means all indices from  $n$  to  $N$  (inclusive). A middle asterisk means all indices from  $m$  to  $n$  (inclusive).

Using a wildcard is the same as if the individual elements of the vector or columns of the array had been listed one by one. For example, the following two *fix ave/time* commands are equivalent, since the *compute rdf* command creates, in this case, a global array with three columns, each of length 50:

```
compute myRDF all rdf 50 1 2
fix 1 all ave/time 100 1 100 c_myRDF[*] file tmp1.rdf mode vector
fix 2 all ave/time 100 1 100 c_myRDF[1] c_myRDF[2] c_myRDF[3] file tmp2.rdf mode vector
```

**Note:** For a vector-style variable, only the wildcard forms “\*n” or “m\*n” are allowed. You must specify the upper bound, because vector-style variable lengths are not determined until the variable is evaluated. If  $n$  is specified larger than the vector length turns out to be, zeroes are output for missing vector values.

The  $N_{\text{every}}$ ,  $N_{\text{repeat}}$ , and  $N_{\text{freq}}$  arguments specify on what time steps the input values will be used in order to contribute to the average. The final averaged quantities are generated on time steps that are a multiple of  $N_{\text{freq}}$ . The average is over  $N_{\text{repeat}}$  quantities, computed in the preceding portion of the simulation every  $N_{\text{every}}$  time steps.  $N_{\text{freq}}$  must be a multiple

of  $N_{\text{every}}$  and  $N_{\text{repeat}}$  must be non-zero even if  $N_{\text{repeat}} = 1$ . Also, the time steps contributing to the average value cannot overlap, i.e.  $N_{\text{repeat}} * N_{\text{every}}$  can not exceed  $N_{\text{freq}}$ .

For example, if  $N_{\text{every}} = 2$ ,  $N_{\text{repeat}} = 6$ , and  $N_{\text{freq}} = 100$ , then values on time steps 90, 92, 94, 96, 98, and 100 will be used to compute the final average on time step 100. Similarly for time steps 190, 192, 194, 196, 198, and 200 on time step 200, etc. If  $N_{\text{repeat}} = 1$  and  $N_{\text{freq}} = 100$ , then no time averaging is done; values are simply generated on time steps 100, 200, etc.

---

If a value begins with “c\_”, a compute ID must follow which has been previously defined in the input script. If *mode* = scalar, then if no bracketed term is appended, the global scalar calculated by the compute is used. If a bracketed term is appended, the *I*th element of the global vector calculated by the compute is used. If *mode* = vector, then if no bracketed term is appended, the global vector calculated by the compute is used. If a bracketed term is appended, the *I*th column of the global array calculated by the compute is used. See the discussion above for how *I* can be specified with a wildcard asterisk to effectively specify multiple values.

Note that there is a *compute reduce* command that can sum per-atom quantities into a global scalar or vector, which can then be accessed by *fix ave/time*. It can also be a compute defined not in your input script, but by *thermodynamic output* or other fixes such as *fix nvt* or *fix temp/rescale*. See the doc pages for these commands which give the IDs of these computes. Users can also write code for their own compute styles and *add them to LAMMPS*.

If a value begins with “f\_”, a fix ID must follow which has been previously defined in the input script. If *mode* = scalar, then if no bracketed term is appended, the global scalar calculated by the fix is used. If a bracketed term is appended, the *I*th element of the global vector calculated by the fix is used. If *mode* = vector, then if no bracketed term is appended, the global vector calculated by the fix is used. If a bracketed term is appended, the *I*th column of the global array calculated by the fix is used. See the discussion above for how *I* can be specified with a wildcard asterisk to effectively specify multiple values.

Note that some fixes only produce their values on certain time steps, which must be compatible with *Nevery*, else an error will result. Users can also write code for their own fix styles and *add them to LAMMPS*.

If a value begins with “v\_”, a variable name must follow which has been previously defined in the input script. If *mode* = scalar, then only equal-style or vector-style variables can be used, which both produce global values. In this mode, a vector-style variable requires a bracketed term to specify the *I*th element of the vector calculated by the variable. If *mode* = vector, then only a vector-style variable can be used, without a bracketed term. See the *variable* command for details.

Note that variables of style *equal* and *vector* define a formula which can reference individual atom properties or thermodynamic keywords, or they can invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of specifying quantities to time average.

---

Additional optional keywords also affect the operation of this fix.

If the *mode* keyword is set to *scalar*, then all input values must be global scalars, or elements of global vectors. If the *mode* keyword is set to *vector*, then all input values must be global vectors, or columns of global arrays. They can also be global arrays, which are converted into a series of global vectors (one per column), as explained above.

The *ave* keyword determines how the values produced every  $N_{\text{freq}}$  steps are averaged with values produced on previous steps that were multiples of  $N_{\text{freq}}$ , before they are accessed by another output command or written to a file.

If the *ave* setting is *one*, then the values produced on time steps that are multiples of  $N_{\text{freq}}$  are independent of each other; they are output as-is without further averaging.

If the *ave* setting is *running*, then the values produced on time steps that are multiples of  $N_{\text{freq}}$  are summed and averaged in a cumulative sense before being output. Each output value is thus the average of the value produced on that time step with all preceding values. This running average begins when the fix is defined; it can only be restarted by deleting the fix via the *unfix* command, or by re-defining the fix by re-specifying it.

If the *ave* setting is *window*, then the values produced on time steps that are multiples of  $N_{freq}$  are summed and averaged within a moving “window” of time, so that the last  $M$  values are used to produce the output. For example, if  $M = 3$  and  $N_{freq} = 1000$ , then the output on step 10000 will be the average of the individual values on steps 8000, 9000, and 10000. Outputs on early steps will average over less than  $M$  values if they are not available.

The *start* keyword specifies what time step averaging will begin on. The default is step 0. Often input values can be 0.0 at time 0, so setting *start* to a larger value can avoid including a 0.0 in a running or windowed average.

The *off* keyword can be used to flag any of the input values. If a value is flagged, it will not be time averaged. Instead the most recent input value will always be stored and output. This is useful if one or more of the inputs produced by a compute or fix or variable are effectively constant or are simply current values (e.g., they are being written to a file with other time-averaged values for purposes of creating well-formatted output).

The *file* keyword allows a filename to be specified. Every  $N_{freq}$  steps, one quantity or vector of quantities is written to the file for each input value specified in the fix *ave/time* command. For *mode* = scalar, this means a single line is written each time output is performed. Thus the file ends up to be a series of lines, i.e. one column of numbers for each input value. For *mode* = vector, an array of numbers is written each time output is performed. The number of rows is the length of the input vectors, and the number of columns is the number of values. Thus the file ends up to be a series of these array sections.

New in version 4May2022.

If the filename ends in ‘.yaml’ or ‘.yml’ then the output format conforms to the [YAML standard](#) which allows easy import that data into tools and scripts that support reading YAML files. The [structured data Howto](#) contains examples for parsing and plotting such data with very little programming effort in Python using the *pyyaml*, *pandas*, and *matplotlib* packages.

The *overwrite* keyword will continuously overwrite the output file with the latest output, so that it only contains one time step worth of output. This option can only be used with the *ave running* setting.

The *format* keyword sets the numeric format of each value when it is printed to a file via the *file* keyword. Note that all values are floating point quantities. The default format is %g. You can specify a higher precision if desired (e.g., %20.16g).

The *title1* and *title2* and *title3* keywords allow specification of the strings that will be printed as the first 2 or 3 lines of the output file, assuming the *file* keyword was used. LAMMPS uses default values for each of these, so they do not need to be specified.

By default, these header lines are as follows for *mode* = scalar:

```
# Time-averaged data for fix ID
# TimeStep value1 value2 ...
```

In the first line, ID is replaced with the fix-ID. In the second line the values are replaced with the appropriate fields from the fix *ave/time* command. There is no third line in the header of the file, so the *title3* setting is ignored when *mode* = scalar.

By default, these header lines are as follows for *mode* = vector:

```
# Time-averaged data for fix ID
# TimeStep Number-of-rows
# Row value1 value2 ...
```

In the first line, ID is replaced with the fix-ID. The second line describes the two values that are printed at the first of each section of output. In the third line the values are replaced with the appropriate fields from the fix *ave/time* command.

## 17.18.4 Restart, fix\_modify, output, run start/stop, minimize info

New in version 4May2022.

No information about this fix is written to *binary restart files*. The *fix\_modify colname* option can be used to change the name of the column in the output file. When writing a YAML format file this name will be in the list of keywords.

This fix produces a global scalar or global vector or global array which can be accessed by various *output commands*. The values can only be accessed on time steps that are multiples of  $N_{\text{freq}}$  since that is when averaging is performed.

A scalar is produced if only a single input value is averaged and *mode* = scalar. A vector is produced if multiple input values are averaged for *mode* = scalar, or a single input value for *mode* = vector. In the first case, the length of the vector is the number of inputs. In the second case, the length of the vector is the same as the length of the input vector. An array is produced if multiple input values are averaged and *mode* = vector. The global array has # of rows = length of the input vectors and # of columns = number of inputs.

If the fix produces a scalar or vector, then the scalar and each element of the vector can be either “intensive” or “extensive”, depending on whether the values contributing to the scalar or vector element are “intensive” or “extensive”. If the fix produces an array, then all elements in the array must be the same, either “intensive” or “extensive”. If a compute or fix provides the value being time averaged, then the compute or fix determines whether the value is intensive or extensive; see the page for that compute or fix for further info. Values produced by a variable are treated as intensive.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.18.5 Restrictions

none

## 17.18.6 Related commands

*compute*, *fix ave/atom*, *fix ave/chunk*, *fix ave/histo*, *variable*, *fix ave/correlate*,

## 17.18.7 Default

The option defaults are *mode* = scalar, *ave* = one, *start* = 0, no file output, *format* = %g, *title* 1,2,3 = strings as described above, and no off settings for any input values.

## 17.19 fix aveforce command

### 17.19.1 Syntax

```
fix ID group-ID aveforce fx fy fz keyword value ...
```

- ID, group-ID are documented in *fix* command
- aveforce = style name of this fix command
- fx,fy,fz = force component values (force units)

any of fx,fy,fz can be a variable (see below)

- zero or more keyword/value pairs may be appended to args

- keyword = *region*

*region* value = region-ID

region-ID = ID of region atoms must be in to have added force

## 17.19.2 Examples

```
fix pressdown topwall aveforce 0.0 -1.0 0.0
fix 2 bottomwall aveforce NULL -1.0 0.0 region top
fix 2 bottomwall aveforce NULL -1.0 v_oscillate region top
```

## 17.19.3 Description

Apply an additional external force to a group of atoms in such a way that every atom experiences the same force. This is useful for pushing on wall or boundary atoms so that the structure of the wall does not change over time.

The existing force is averaged for the group of atoms, component by component. The actual force on each atom is then set to the average value plus the component specified in this command. This means each atom in the group receives the same force.

Any of the  $f_x$ ,  $f_y$ , or  $f_z$  values can be specified as NULL, which means the force in that dimension is not changed. Note that this is not the same as specifying a 0.0 value, since that sets all forces to the same average value without adding in any additional force.

Any of the three quantities defining the force components, namely  $f_x$ ,  $f_y$ , and  $f_z$ , can be specified as an equal-style *variable*. If the value is a variable, it should be specified as  $v\_name$ , where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the average force.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent average force.

If the *region* keyword is used, the atom must also be in the specified geometric *region* in order to have force added to it.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 17.19.4 Restart, `fix_modify`, `output`, `run start/stop`, `minimize info`

No information about this fix is written to *binary restart files*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global three-vector of forces, which can be accessed by various *output commands*. This is the total force on the group of atoms before the forces on individual atoms are changed by the fix. The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command. You should not specify force components with a variable that has time-dependence for use with a minimizer, since the minimizer increments the timestep as the iteration count during the minimization.

### 17.19.5 Restrictions

none

### 17.19.6 Related commands

*fix setforce*, *fix addforce*

### 17.19.7 Default

none

## 17.20 `fix balance` command

### 17.20.1 Syntax

```
fix ID group-ID balance Nfreq thresh style args keyword args ...
```

- ID, group-ID are documented in *fix* command
- balance = style name of this fix command
- Nfreq = perform dynamic load balancing every this many steps
- thresh = imbalance threshold that must be exceeded to perform a re-balance
- style = *shift* or *rcb*

`shift args = dimstr Niter stopthresh`

`dimstr = sequence of letters containing x or y or z, each not more than once`

`Niter = # of times to iterate within each dimension of dimstr sequence`

`stopthresh = stop balancing when this imbalance threshold is reached`

`rcb args = none`

- zero or more keyword/arg pairs may be appended
- keyword = *weight* or *out*



*weight* style args = use weighted particle counts for the balancing  
*style* = *group* or *neigh* or *time* or *var* or *store*  
*group* args = Ngroup group1 weight1 group2 weight2 ...  
 Ngroup = number of groups with assigned weights  
 group1, group2, ... = group IDs  
 weight1, weight2, ... = corresponding weight factors  
*neigh* factor = compute weight based on number of neighbors  
 factor = scaling factor (> 0)  
*time* factor = compute weight based on time spend computing  
 factor = scaling factor (> 0)  
*var* name = take weight from atom-style variable  
 name = name of the atom-style variable  
*store* name = store weight in custom atom property defined by *fix property/atom*  
 ↪command  
 name = atom property name (without d\_ prefix)  
 out arg = filename  
 filename = write each processor's sub-domain to a file, at each re-balancing

## 17.20.2 Examples

```
fix 2 all balance 1000 1.05 shift x 10 1.05
fix 2 all balance 100 0.9 shift xy 20 1.1 out tmp.balance
fix 2 all balance 100 0.9 shift xy 20 1.1 weight group 3 substrate 3.0 solvent 1.0
↪solute 0.8 out tmp.balance
fix 2 all balance 100 1.0 shift x 10 1.1 weight time 0.8
fix 2 all balance 100 1.0 shift xy 5 1.1 weight var myweight weight neigh 0.6 weight
↪store allweight
fix 2 all balance 1000 1.1 rcb
```

## 17.20.3 Description

This command adjusts the size and shape of processor sub-domains within the simulation box, to attempt to balance the number of particles and thus the computational cost (load) evenly across processors. The load balancing is “dynamic” in the sense that re-balancing is performed periodically during the simulation. To perform “static” balancing, before or between runs, see the *balance* command.

Load-balancing is typically most useful if the particles in the simulation box have a spatially-varying density distribution or where the computational cost varies significantly between different atoms (e.g., a model of a vapor/liquid interface, or a solid with an irregular-shaped geometry containing void regions, or *hybrid pair style simulations* that combine pair styles with different computational cost). In these cases, the LAMMPS default of dividing the simulation box volume into a regular-spaced grid of 3d bricks, with one equal-volume sub-domain per processor, may assign numbers of particles per processor in a way that the computational effort varies significantly. This can lead to poor performance when the simulation is run in parallel.

The balancing can be performed with or without per-particle weighting. With no weighting, the balancing attempts to assign an equal number of particles to each processor. With weighting, the balancing attempts to assign an equal aggregate computational weight to each processor, which typically induces a different number of atoms assigned to each processor.

**Note:** The weighting options listed above are documented with the *balance* command in *this section of the balance command* doc page. That section describes the various weighting options and gives a few examples of how they can

be used. The weighting options are the same for both the `fix balance` and `balance` commands.

---

Note that the `processors` command allows some control over how the box volume is split across processors. Specifically, for a  $P_x \times P_y \times P_z$  grid of processors, it allows choices of  $P_x$ ,  $P_y$ , and  $P_z$  subject to the constraint that  $P_x P_y P_z = P$ , the total number of processors. This is sufficient to achieve good load-balance for some problems on some processor counts. However, all the processor sub-domains will still have the same shape and the same volume.

On a particular time step, a load-balancing operation is only performed if the current “imbalance factor” in particles owned by each processor exceeds the specified `thresh` parameter. The imbalance factor is defined as the maximum number of particles (or weight) owned by any processor, divided by the average number of particles (or weight) per processor. Thus, an imbalance factor of 1.0 is perfect balance.

As an example, for 10000 particles running on 10 processors, if the most heavily loaded processor has 1200 particles, then the imbalance factor is 1.2, meaning there is a 20% imbalance. Note that re-balances can be forced even if the current balance is perfect (1.0) by specifying a `thresh < 1.0`.

---

**Note:** This command attempts to minimize the imbalance factor, as defined above. But depending on the method a perfect balance (1.0) may not be achieved. For example, “grid” methods (defined below) that create a logical 3d grid cannot achieve perfect balance for many irregular distributions of particles. Likewise, if a portion of the system is a perfect, non-rotated lattice (e.g., the initial system is generated by the `create_atoms` command with no rotations), then “grid” methods may be unable to achieve exact balance. This is because entire lattice planes will be owned or not owned by a single processor.

---

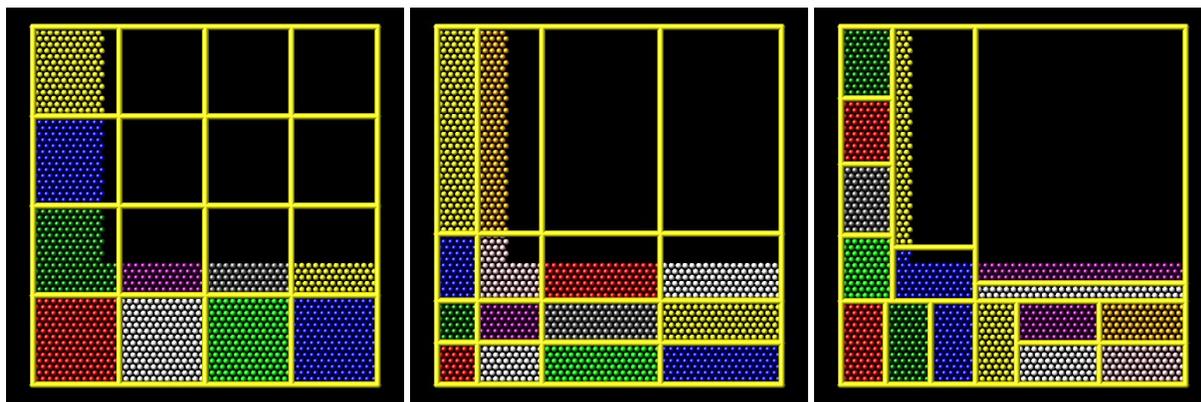
---

**Note:** The imbalance factor is also an estimate of the maximum speed-up you can hope to achieve by running a perfectly balanced simulation versus an imbalanced one. In the example above, the 10000-particle simulation could run up to 20% faster if it were perfectly balanced, versus when imbalanced. However, computational cost is not strictly proportional to particle count, and changing the relative size and shape of processor sub-domains may lead to additional computational and communication overheads (e.g., in the PPPM solver used via the `k-space_style` command). Thus, you should benchmark the run times of a simulation before and after balancing.

---

The method used to perform a load balance is specified by one of the listed styles, which are described in detail below. There are two kinds of styles.

The `shift` style is a “grid” method which produces a logical 3d grid of processors. It operates by changing the cutting planes (or lines) between processors in 3d (or 2d), to adjust the volume (area in 2d) assigned to each processor, as in the following 2d diagram where processor sub-domains are shown and atoms are colored by the processor that owns them.



The leftmost diagram is the default partitioning of the simulation box across processors (one sub-box for each of 16 processors); the middle diagram is after a “grid” method has been applied. The *rcb* style is a “tiling” method which does not produce a logical 3d grid of processors. Rather it tiles the simulation domain with rectangular sub-boxes of varying size and shape in an irregular fashion so as to have equal numbers of particles (or weight) in each sub-box, as in the rightmost diagram above.

The “grid” methods can be used with either of the *comm\_style* command options, *brick* or *tiled*. The “tiling” methods can only be used with *comm\_style tiled*.

When a “grid” method is specified, the current domain partitioning can be either a logical 3d grid or a tiled partitioning. In the former case, the current logical 3d grid is used as a starting point and changes are made to improve the imbalance factor. In the latter case, the tiled partitioning is discarded and a logical 3d grid is created with uniform spacing in all dimensions. This is the starting point for the balancing operation.

When a “tiling” method is specified, the current domain partitioning (“grid” or “tiled”) is ignored, and a new partitioning is computed from scratch.

The *group-ID* is ignored. However the impact of balancing on different groups of atoms can be affected by using the *group weight* style as described below.

The  $N_{\text{freq}}$  setting determines how often a re-balance is performed. If  $N_{\text{freq}} > 0$ , then re-balancing will occur every  $N_{\text{freq}}$  steps. Each time a re-balance occurs, a reneighboring is triggered, so  $N_{\text{freq}}$  should not be too small. If  $N_{\text{freq}} = 0$ , then re-balancing will be done every time reneighboring normally occurs, as determined by the *neighbor* and *neigh\_modify* command settings.

On re-balance steps, re-balancing will only be attempted if the current imbalance factor, as defined above, exceeds the *thresh* setting.

The *shift* style invokes a “grid” method for balancing, as described above. It changes the positions of cutting planes between processors in an iterative fashion, seeking to reduce the imbalance factor.

The *dimstr* argument is a string of characters, each of which must be *x* or *y* or *z*. Each character can appear zero or one time, since there is no advantage to balancing on a dimension more than once. You should normally only list dimensions where you expect there to be a density variation in the particles.

Balancing proceeds by adjusting the cutting planes in each of the dimensions listed in *dimstr*, one dimension at a time. For a single dimension, the balancing operation (described below) is iterated on up to  $N_{\text{iter}}$  times. After each dimension finishes, the imbalance factor is re-computed, and the balancing operation halts if the *stopthresh* criterion is met.

A re-balance operation in a single dimension is performed using a density-dependent recursive multisectioning algorithm, where the position of each cutting plane (line in 2d) in the dimension is adjusted independently. This is similar to a recursive bisectioning for a single value, except that the bounds used for each bisectioning take advantage of information from neighboring cuts if possible, as well as counts of particles at the bounds on either side of each cuts, which themselves were cuts in previous iterations. The latter is used to infer a density of particles near each of the current cuts. At each iteration, the count of particles on either side of each plane is tallied. If the counts do not match the target value for the plane, the position of the cut is adjusted based on the local density. The low and high bounds are adjusted on each iteration, using new count information, so that they become closer together over time. Thus as the recursion progresses, the count of particles on either side of the plane gets closer to the target value.

The density-dependent part of this algorithm is often an advantage when you re-balance a system that is already nearly balanced. It typically converges more quickly than the geometric bisectioning algorithm used by the *balance* command. However, it can be a disadvantage if you attempt to re-balance a system that is far from balanced, and converge more slowly. In this case you probably want to use the *balance* command before starting a run, so that you begin the run with a balanced system.

Once the re-balancing is complete and final processor sub-domains assigned, particles migrate to their new owning processor as part of the normal reneighboring procedure.

**Note:** At each re-balance operation, the bisectioning for each cutting plane (line in 2d) typically starts with low and high bounds separated by the extent of a processor's sub-domain in one dimension. The size of this bracketing region shrinks based on the local density, as described above, which should typically be 1/2 or more every iteration. Thus if  $N_{\text{iter}}$  is specified as 10, the cutting plane will typically be positioned to better than 1 part in 1000 accuracy (relative to the perfect target position). For  $N_{\text{iter}} = 20$ , it will be accurate to better than 1 part in a million. Thus there is no need to set  $N_{\text{iter}}$  to a large value. This is especially true if you are re-balancing often enough that each time you expect only an incremental adjustment in the cutting planes is necessary. LAMMPS will check if the threshold accuracy is reached (in a dimension) is less iterations than  $N_{\text{iter}}$  and exit early.

---

The *rcb* style invokes a “tiled” method for balancing, as described above. It performs a recursive coordinate bisectioning (RCB) of the simulation domain. The basic idea is as follows.

The simulation domain is cut into two boxes by an axis-aligned cut in the longest dimension, leaving one new box on either side of the cut. All the processors are also partitioned into two groups, half assigned to the box on the lower side of the cut, and half to the box on the upper side. If the processor count is odd, one side gets an extra processor. The cut is positioned so that the number of atoms in the lower box is exactly the number that the processors assigned to that box should own for load balance to be perfect. This also makes load balance for the upper box perfect. The positioning is done iteratively, by a bisectioning method. Note that counting atoms on either side of the cut requires communication between all processors at each iteration.

That is the procedure for the first cut. Subsequent cuts are made recursively, in exactly the same manner. The subset of processors assigned to each box make a new cut in the longest dimension of that box, splitting the box, the subset of processors, and the atoms in the box in two. The recursion continues until every processor is assigned a sub-box of the entire simulation domain, and owns the atoms in that sub-box.

---

The *out* keyword writes text to the specified *filename* with the results of each re-balancing operation. The file contains the bounds of the sub-domain for each processor after the balancing operation completes. The format of the file is compatible with the [Pizza.py mdump](#) tool which has support for manipulating and visualizing mesh files. An example is shown here for a balancing by four processors for a 2d problem:

```
ITEM: TIMESTEP
0
ITEM: NUMBER OF NODES
16
ITEM: BOX BOUNDS
0 10
0 10
0 10
ITEM: NODES
1 1 0 0 0
2 1 5 0 0
3 1 5 5 0
4 1 0 5 0
5 1 5 0 0
6 1 10 0 0
7 1 10 5 0
8 1 5 5 0
9 1 0 5 0
10 1 5 5 0
11 1 5 10 0
```

(continues on next page)

(continued from previous page)

```

12 1 10 5 0
13 1 5 5 0
14 1 10 5 0
15 1 10 10 0
16 1 5 10 0
ITEM: TIMESTEP
0
ITEM: NUMBER OF SQUARES
4
ITEM: SQUARES
1 1 1 2 3 4
2 1 5 6 7 8
3 1 9 10 11 12
4 1 13 14 15 16

```

The coordinates of all the vertices are listed in the `NODES` section, five per processor. Note that the four sub-domains share vertices, so there will be duplicate nodes in the list.

The “`SQUARES`” section lists the node IDs of the four vertices in a rectangle for each processor (1 to 4).

For a 3d problem, the syntax is similar but with eight vertices listed for each processor instead of four, and “`SQUARES`” replaced by “`CUBES`”.

#### 17.20.4 Restart, `fix_modify`, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global scalar which is the imbalance factor after the most recent re-balance and a global vector of length 3 with additional information about the most recent re-balancing. The three values in the vector are as follows:

- 1 = max # of particles per processor
- 2 = total # iterations performed in last re-balance
- 3 = imbalance factor right before the last re-balance was performed

As explained above, the imbalance factor is the ratio of the maximum number of particles (or total weight) on any processor to the average number of particles (or total weight) per processor.

These quantities can be accessed by various *output commands*. The scalar and vector values calculated by this fix are “intensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.20.5 Restrictions

For 2d simulations, the *z* style cannot be used, nor can *z* appear in *dimstr* for the *shift* style.

Balancing through recursive bisectioning (*rcb* style) requires *comm\_style tiled*.

## 17.20.6 Related commands

*group*, *processors*, *balance*, *comm\_style*

## 17.20.7 Default

none

## 17.21 fix bocs command

### 17.21.1 Syntax

```
fix ID group-ID bocs keyword values ...
```

keyword = *temp* or *cgiso* or *analytic* or *linear\_spline* or *cubic\_spline*

*temp* values = Tstart Tstop Tdamp

*cgiso* values = Pstart Pstop Pdamp

*basis set*

*analytic* values = V\_avg N\_particles N\_coeff Coeff\_1 Coeff\_2 ... Coeff\_N

*linear\_spline* values = input\_filename

*cubic\_spline* values = input\_filename

### 17.21.2 Examples

```
fix 1 all bocs temp 300.0 300.0 100.0 cgiso 0.986 0.986 1000.0 analytic 66476.015 968 2.↵  
↵245030.10 8962.20
```

```
fix 1 all bocs temp 300.0 300.0 100.0 cgiso 0.986 0.986 1000.0 cubic_spline input_Fv.dat
```

```
thermo_modify press 1_press
```

### 17.21.3 Description

These commands incorporate a pressure correction as described by Dunn and Noid in (*Dunn1*) to the standard MTTK barostat by Martyna et. al. in (*Martyna*). The first half of the command mimics a standard fix npt command:

```
fix 1 all bocs temp Tstart Tstop Tcoupl cgiso Pstart Pstop Pdamp
```

The two differences are replacing *npt* with *bocs*, and replacing *isolaniso/etc* with *cgiso*. The rest of the command details what form you would like to use for the pressure correction equation. The choices are: *analytic*, *linear\_spline*, or *cubic\_spline*.

With either spline method, the only argument that needs to follow it is the name of a file that contains the desired pressure correction as a function of volume. The file must be formatted so each line has:

```
Volume_i, PressureCorrection_i
```

Note both the COMMA and the SPACE separating the volume's value and its corresponding pressure correction. The volumes in the file must be uniformly spaced. Both the volumes and the pressure corrections should be provided in the proper units, e.g. if you are using *units real*, the volumes should all be in cubic angstroms, and the pressure corrections should all be in atmospheres. Furthermore, the table should start/end at a volume considerably smaller/larger than you expect your system to sample during the simulation. If the system ever reaches a volume outside of the range provided, the simulation will stop.

With the *analytic* option, the arguments are as follows:

```
... analytic V_avg N_particles N_coeff Coeff_1 Coeff_2 ... Coeff_N
```

Note that *V\_avg* and *Coeff\_i* should all be in the proper units, e.g. if you are using *units real*, *V\_avg* should be in cubic angstroms, and the coefficients should all be in atmospheres \* cubic angstroms.

#### 17.21.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the cumulative global energy change to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the fix continues in an uninterrupted fashion.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a temperature *compute* you have defined to this fix which will be used in its thermostating procedure, as described above. For consistency, the group used by this fix and by the compute should be the same.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is "extensive".

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

#### 17.21.5 Restrictions

As this is computing a (modified) pressure, group-ID should be *all*.

The pressure correction has only been tested for use with an isotropic pressure coupling in 3 dimensions.

By default, LAMMPS will still report the normal value for the pressure if the pressure is printed via a *thermo* command, or if the pressures are written to a file every so often. In order to have LAMMPS report the modified pressure, you must include the *thermo\_modify* command given in the examples. For the last argument in the command, you should put XXXX\_press, where XXXX is the ID given to the fix bocs command (in the example, the ID of the fix bocs command is 1).

This fix is part of the BOCS package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.21.6 Further information

For more details about the pressure correction and the entire BOCS software package, visit the [BOCS package on GitHub](#) and read the release paper by Dunn et. al. ([Dunn2](#)).

---

(**Dunn1**) Dunn and Noid, J Chem Phys, 143, 243148 (2015).

(**Martyna**) Martyna, Tobias, and Klein, J Chem Phys, 101, 4177 (1994).

(**Dunn2**) Dunn, Lebold, DeLysler, Rudzinski, and Noid, J. Phys. Chem. B, 122, 3363 (2018).

## 17.22 fix bond/break command

### 17.22.1 Syntax

```
fix ID group-ID bond/break Nevery bondtype Rmax keyword values ...
```

- ID, group-ID are documented in *fix* command
- bond/break = style name of this fix command
- Nevery = attempt bond breaking every this many steps
- bondtype = type of bonds to break
- Rmax = bond longer than Rmax can break (distance units)
- zero or more keyword/value pairs may be appended to args
- keyword = *prob*

*prob* values = fraction seed

fraction = break a bond with this probability if otherwise eligible

seed = random number seed (positive integer)

### 17.22.2 Examples

```
fix 5 all bond/break 10 2 1.2  
fix 5 polymer bond/break 1 1 2.0 prob 0.5 49829
```

### 17.22.3 Description

Break bonds between pairs of atoms as a simulation runs according to specified criteria. This can be used to model the dissolution of a polymer network due to stretching of the simulation box or other deformations. In this context, a bond means an interaction between a pair of atoms computed by the *bond\_style* command. Once the bond is broken it will be permanently deleted, as will all angle, dihedral, and improper interactions that bond is part of.

This is different than a *pairwise* bond-order potential such as Tersoff or AIREBO which infers bonds and many-body interactions based on the current geometry of a small cluster of atoms and effectively creates and destroys bonds and higher-order many-body interactions from timestep to timestep as atoms move.

A check for possible bond breakage is performed every *Nevery* timesteps. If two bonded atoms I,J are further than a distance *Rmax* of each other, if the bond is of type *bondtype*, and if both I and J are in the specified fix group, then I,J is labeled as a “possible” bond to break.



If several bonds involving an atom are stretched, it may have multiple possible bonds to break. Every atom checks its list of possible bonds to break and labels the longest such bond as its “sole” bond to break. After this is done, if atom I is bonded to atom J in its sole bond, and atom J is bonded to atom I in its sole bond, then the I,J bond is “eligible” to be broken.

Note that these rules mean an atom will only be part of at most one broken bond on a given timestep. It also means that if atom I chooses atom J as its sole partner, but atom J chooses atom K as its sole partner (due to  $R_{jk} > R_{ij}$ ), then this means atom I will not be part of a broken bond on this timestep, even if it has other possible bond partners.

The *prob* keyword can effect whether an eligible bond is actually broken. The *fraction* setting must be a value between 0.0 and 1.0. A uniform random number between 0.0 and 1.0 is generated and the eligible bond is only broken if the random number  $<$  fraction.

When a bond is broken, data structures within LAMMPS that store bond topology are updated to reflect the breakage. Likewise, if the bond is part of a 3-body (angle) or 4-body (dihedral, improper) interaction, that interaction is removed as well. These changes typically affect pairwise interactions between atoms that used to be part of bonds, angles, etc.

---

**Note:** One data structure that is not updated when a bond breaks are the molecule IDs stored by each atom. Even though one molecule becomes two molecules due to the broken bond, all atoms in both new molecules retain their original molecule IDs.

---

Computationally, each timestep this fix operates, it loops over all the bonds in the system and computes distances between pairs of bonded atoms. It also communicates between neighboring processors to coordinate which bonds are broken. Moreover, if any bonds are broken, neighbor lists must be immediately updated on the same timestep. This is to insure that any pairwise interactions that should be turned “on” due to a bond breaking, because they are no longer excluded by the presence of the bond and the settings of the *special\_bonds* command, will be immediately recognized. All of these operations increase the cost of a timestep. Thus you should be cautious about invoking this fix too frequently.

You can dump out snapshots of the current bond topology via the *dump local* command.

---

**Note:** Breaking a bond typically alters the energy of a system. You should be careful not to choose bond breaking criteria that induce a dramatic change in energy. For example, if you define a very stiff harmonic bond and break it when 2 atoms are separated by a distance far from the equilibrium bond length, then the 2 atoms will be dramatically released when the bond is broken. More generally, you may need to thermostat your system to compensate for energy changes resulting from broken bonds (and angles, dihedrals, impropers).

---

See the *Howto* page on broken bonds for more information on related features in LAMMPS.

---

#### 17.22.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes two statistics which it stores in a global vector of length 2, which can be accessed by various *output commands*. The vector values calculated by this fix are “intensive”.

These are the 2 quantities:

- (1) # of bonds broken on the most recent breakage timestep
- (2) cumulative # of bonds broken

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.22.5 Restrictions

This fix is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

### 17.22.6 Related commands

*fix bond/create*, *fix bond/react*, *fix bond/swap*, *dump local*, *special\_bonds*

### 17.22.7 Default

The option defaults are `prob = 1.0`.

## 17.23 fix bond/create command

## 17.24 fix bond/create/angle command

### 17.24.1 Syntax

```
fix ID group-ID bond/create Nevery itype jtype Rmin bondtype keyword values ...
```

- ID, group-ID are documented in *fix* command
- bond/create = style name of this fix command
- Nevery = attempt bond creation every this many steps
- itype,jtype = atoms of itype can bond to atoms of jtype
- Rmin = 2 atoms separated by less than Rmin can bond (distance units)
- bondtype = type of created bonds
- zero or more keyword/value pairs may be appended to args
- keyword = *iparam* or *jparam* or *prob* or *atype* or *dtype* or *itype* or *aconstrain*

*iparam* values = maxbond, newtype

maxbond = max # of bonds of bondtype the itype atom can have

newtype = change the itype atom to this type when maxbonds exist

*jparam* values = maxbond, newtype

maxbond = max # of bonds of bondtype the jtype atom can have

newtype = change the jtype atom to this type when maxbonds exist

*prob* values = fraction seed

fraction = create a bond with this probability if otherwise eligible

seed = random number seed (positive integer)

*atype* value = angletype

angletype = type of created angles

*dtype* value = dihedraltype

dihedraltype = type of created dihedrals

*itype* value = improprtype

improprtype = type of created impropers

*aconstrain* value = amin amax

amin = minimal angle at which new bonds can be created  
 amax = maximal angle at which new bonds can be created

## 17.24.2 Examples

```
fix 5 all bond/create 10 1 2 0.8 1
fix 5 all bond/create 1 3 3 0.8 1 prob 0.5 85784 iparam 2 3
fix 5 all bond/create 1 3 3 0.8 1 prob 0.5 85784 iparam 2 3 atype 1 dtype 2
fix 5 all bond/create/angle 10 1 2 1.122 1 aconstrain 120 180 prob 1 4928459 iparam 2 1
↪ jparam 2 2
```

## 17.24.3 Description

Create bonds between pairs of atoms as a simulation runs according to specified criteria. This can be used to model cross-linking of polymers, the formation of a percolation network, etc. In this context, a bond means an interaction between a pair of atoms computed by the *bond\_style* command. Once the bond is created it will be permanently in place. Optionally, the creation of a bond can also create angle, dihedral, and improper interactions that bond is part of. See the discussion of the *atype*, *dtype*, and *itype* keywords below.

This is different than a *pairwise* bond-order potential such as Tersoff or AIREBO which infers bonds and many-body interactions based on the current geometry of a small cluster of atoms and effectively creates and destroys bonds and higher-order many-body interactions from timestep to timestep as atoms move.

A check for possible new bonds is performed every *Nevery* timesteps. If two atoms I,J are within a distance *Rmin* of each other, if I is of atom type *itype*, if J is of atom type *jtype*, if both I and J are in the specified fix group, if a bond does not already exist between I and J, and if both I and J meet their respective *maxbond* requirement (explained below), then I,J is labeled as a “possible” bond pair.

If several atoms are close to an atom, it may have multiple possible bond partners. Every atom checks its list of possible bond partners and labels the closest such partner as its “sole” bond partner. After this is done, if atom I has atom J as its sole partner, and atom J has atom I as its sole partner, then the I,J bond is “eligible” to be formed.

Note that these rules mean an atom will only be part of at most one created bond on a given timestep. It also means that if atom I chooses atom J as its sole partner, but atom J chooses atom K as its sole partner (due to  $R_{jk} < R_{ij}$ ), then this means atom I will not form a bond on this timestep, even if it has other possible bond partners.

It is permissible to have *itype* = *jtype*. *Rmin* must be  $\leq$  the pairwise cutoff distance between *itype* and *jtype* atoms, as defined by the *pair\_style* command.

The *iparam* and *jparam* keywords can be used to limit the bonding functionality of the participating atoms. Each atom keeps track of how many bonds of *bondtype* it already has. If atom I of *itype* already has *maxbond* bonds (as set by the *iparam* keyword), then it will not form any more. Likewise for atom J. If *maxbond* is set to 0, then there is no limit on the number of bonds that can be formed with that atom.

The *newtype* value for *iparam* and *jparam* can be used to change the atom type of atom I or J when it reaches *maxbond* number of bonds of type *bondtype*. This means it can now interact in a pairwise fashion with other atoms in a different way by specifying different *pair\_coeff* coefficients. If you do not wish the atom type to change, simply specify *newtype* as *itype* or *jtype*.

The *prob* keyword can also effect whether an eligible bond is actually created. The *fraction* setting must be a value between 0.0 and 1.0. A uniform random number between 0.0 and 1.0 is generated and the eligible bond is only created if the random number  $<$  fraction.

The *aconstrain* keyword is only available with the fix bond/create/angle command. It allows to specify a minimal and maximal angle *amin* and *amax* between the two prospective bonding partners and a third particle that is already bonded to one of the two partners. Such a criterion can be important when new angles are defined together with the formation

of a new bond. Without a restriction on the permissible angle, and for stiffer angle potentials, very large energies can arise and lead to uncontrolled behavior.

Any bond that is created is assigned a bond type of *bondtype*.

When a bond is created, data structures within LAMMPS that store bond topology are updated to reflect the creation. If the bond is part of new 3-body (angle) or 4-body (dihedral, improper) interactions, you can choose to create new angles, dihedrals, impropers as well, using the *atype*, *dtype*, and *itype* keywords. All of these changes typically affect pairwise interactions between atoms that are now part of new bonds, angles, etc.

---

**Note:** One data structure that is not updated when a bond breaks are the molecule IDs stored by each atom. Even though two molecules become one molecule due to the created bond, all atoms in the new molecule retain their original molecule IDs.

---

If the *atype* keyword is used and if an angle potential is defined via the *angle\_style* command, then any new 3-body interactions inferred by the creation of a bond will create new angles of type *angletype*, with parameters assigned by the corresponding *angle\_coeff* command. Likewise, the *dtype* and *itype* keywords will create new dihedrals and impropers of type *dihedraltype* and *improptype*.

---

**Note:** To create a new bond, the internal LAMMPS data structures that store this information must have space for it. When LAMMPS is initialized from a data file, the list of bonds is scanned and the maximum number of bonds per atom is tallied. If some atom will acquire more bonds than this limit as this fix operates, then the “extra bond per atom” parameter must be set to allow for it. Ditto for “extra angle per atom”, “extra dihedral per atom”, and “extra improper per atom” if angles, dihedrals, or impropers are being added when bonds are created. See the *read\_data* or *create\_box* command for more details. Note that a data file with no atoms can be used if you wish to add non-bonded atoms via the *create\_atoms* command, e.g. for a percolation simulation.

---

---

**Note:** LAMMPS stores and maintains a data structure with a list of the first, second, and third neighbors of each atom (within the bond topology of the system) for use in weighting pairwise interactions for bonded atoms. Note that adding a single bond always adds a new first neighbor but may also induce \*many\* new second and third neighbors, depending on the molecular topology of your system. The “extra special per atom” parameter must typically be set to allow for the new maximum total size (first + second + third neighbors) of this per-atom list. There are 2 ways to do this. See the *read\_data* or *create\_box* commands for details.

---

---

**Note:** Even if you do not use the *atype*, *dtype*, or *itype* keywords, the list of topological neighbors is updated for atoms affected by the new bond. This in turn affects which neighbors are considered for pairwise interactions, using the weighting rules set by the *special\_bonds* command. Consider a new bond created between atoms I,J. If J has a bonded neighbor K, then K becomes a second neighbor of I. Even if the *atype* keyword is not used to create angle I-J-K, the pairwise interaction between I and K will be potentially turned off or weighted by the 1-3 weighting specified by the *special\_bonds* command. This is the case even if the “angle yes” option was used with that command. The same is true for third neighbors (1-4 interactions), the *dtype* keyword, and the “dihedral yes” option used with the *special\_bonds* command.

---

Note that even if your simulation starts with no bonds, you must define a *bond\_style* and use the *bond\_coeff* command to specify coefficients for the *bondtype*. Similarly, if new atom types are specified by the *iparam* or *jparam* keywords, they must be within the range of atom types allowed by the simulation and pairwise coefficients must be specified for the new types.

Computationally, each timestep this fix operates, it loops over neighbor lists and computes distances between pairs of atoms in the list. It also communicates between neighboring processors to coordinate which bonds are created.

Moreover, if any bonds are created, neighbor lists must be immediately updated on the same timestep. This is to insure that any pairwise interactions that should be turned “off” due to a bond creation, because they are now excluded by the presence of the bond and the settings of the *special\_bonds* command, will be immediately recognized. All of these operations increase the cost of a timestep. Thus you should be cautious about invoking this fix too frequently.

You can dump out snapshots of the current bond topology via the *dump local* command.

---

**Note:** Creating a bond typically alters the energy of a system. You should be careful not to choose bond creation criteria that induce a dramatic change in energy. For example, if you define a very stiff harmonic bond and create it when 2 atoms are separated by a distance far from the equilibrium bond length, then the 2 atoms will oscillate dramatically when the bond is formed. More generally, you may need to thermostat your system to compensate for energy changes resulting from created bonds (and angles, dihedrals, impropers).

---

#### 17.24.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes two statistics which it stores in a global vector of length 2, which can be accessed by various *output commands*. The vector values calculated by this fix are “intensive”.

These are the 2 quantities:

- (1) # of bonds created on the most recent creation timestep
- (2) cumulative # of bonds created

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.24.5 Restrictions

This fix is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

#### 17.24.6 Related commands

*fix bond/break, fix bond/react, fix bond/swap, dump local, special\_bonds*

#### 17.24.7 Default

The option defaults are *iparam* = (0,*itype*), *jparam* = (0,*jtype*), and *prob* = 1.0.

## 17.25 fix bond/react command

### 17.25.1 Syntax

```
fix ID group-ID bond/react common_keyword values ...
  react react-ID react-group-ID Nevery Rmin Rmax template-ID(pre-reacted) template-
  ↪ID(post-reacted) map_file individual_keyword values ...
  react react-ID react-group-ID Nevery Rmin Rmax template-ID(pre-reacted) template-
  ↪ID(post-reacted) map_file individual_keyword values ...
  react react-ID react-group-ID Nevery Rmin Rmax template-ID(pre-reacted) template-
  ↪ID(post-reacted) map_file individual_keyword values ...
  ...
```

- ID, group-ID are documented in *fix* command.
- bond/react = style name of this fix command
- the common keyword/values may be appended directly after ‘bond/react’
- common keywords apply to all reaction specifications
- common\_keyword = *stabilization* or *reset\_mol\_ids*
  - stabilization* values = *no* or *yes* *group-ID* *xmax*
    - no* = no reaction site stabilization (default)
    - yes* = perform reaction site stabilization
    - group-ID* = user-assigned prefix for the dynamic group of atoms not currently ↪involved in a reaction
    - xmax* = *xmax* value that is used by an internally-created *nve/limit* integrator
  - reset\_mol\_ids* values = *yes* or *no*
    - yes* = update molecule IDs based on new global topology (default)
    - no* = do not update molecule IDs
- react = mandatory argument indicating new reaction specification
- react-ID = user-assigned name for the reaction
- react-group-ID = only atoms in this group are considered for the reaction
- Nevery = attempt reaction every this many steps
- Rmin = initiator atoms must be separated by more than Rmin to initiate reaction (distance units)
- Rmax = initiator atoms must be separated by less than Rmax to initiate reaction (distance units)
- template-ID(pre-reacted) = ID of a molecule template containing pre-reaction topology
- template-ID(post-reacted) = ID of a molecule template containing post-reaction topology
- map\_file = name of file specifying corresponding atom-IDs in the pre- and post-reacted templates
- zero or more individual keyword/value pairs may be appended to each react argument
- individual\_keyword = *prob* or *max\_rxn* or *stabilize\_steps* or *custom\_charges* or *molecule* or *modify\_create*
  - prob* values = fraction seed
    - fraction = initiate reaction with this probability if otherwise eligible
    - seed = random number seed (positive integer)
  - max\_rxn* value = N
    - N = maximum number of reactions allowed to occur
  - stabilize\_steps* value = timesteps

```

    timesteps = number of timesteps to apply the internally-created nve/limit fix to.
↪reacting atoms
custom_charges value = no or fragmentID
    no = update all atomic charges (default)
    fragmentID = ID of molecule fragment whose charges are updated
molecule value = off or inter or intra
    off = allow both inter- and intramolecular reactions (default)
    inter = search for reactions between molecules with different IDs
    intra = search for reactions within the same molecule
modify_create keyword values
    fit value = all or fragmentID
        all = use all eligible atoms for create-atoms fit (default)
        fragmentID = ID of molecule fragment used for create-atoms fit
    overlap value = R
        R = only insert atom/molecule if further than R from existing particles.
↪(distance units)

```

## 17.25.2 Examples

For unabridged example scripts and files, see `examples/PACKAGES/reaction`.

```

molecule mol1 pre_reacted_topology.txt
molecule mol2 post_reacted_topology.txt
fix 5 all bond/react react myrxn1 all 1 0 3.25 mol1 mol2 map_file.txt

molecule mol1 pre_reacted_rxn1.txt
molecule mol2 post_reacted_rxn1.txt
molecule mol3 pre_reacted_rxn2.txt
molecule mol4 post_reacted_rxn2.txt
fix 5 all bond/react stabilization yes nvt_grp .03 &
    react myrxn1 all 1 0 3.25 mol1 mol2 map_file_rxn1.txt prob 0.50 12345 &
    react myrxn2 all 1 0 2.75 mol3 mol4 map_file_rxn2.txt prob 0.25 12345
fix 6 nvt_grp_REACT nvt temp 300 300 100 # set thermostat after bond/react

```

## 17.25.3 Description

Initiate complex covalent bonding (topology) changes. These topology changes will be referred to as ‘reactions’ throughout this documentation. Topology changes are defined in pre- and post-reaction molecule templates and can include creation and deletion of bonds, angles, dihedrals, impropers, bond types, angle types, dihedral types, atom types, or atomic charges. In addition, reaction by-products or other molecules can be identified and deleted. Finally, atoms can be created and inserted at specific positions relative to the reaction site.

Fix `bond/react` does not use quantum mechanical (eg. `fix qmmm`) or pairwise bond-order potential (eg. Tersoff or AIREBO) methods to determine bonding changes a priori. Rather, it uses a distance-based probabilistic criteria to effect predetermined topology changes in simulations using standard force fields.

This fix was created to facilitate the dynamic creation of polymeric, amorphous or highly cross-linked systems. A suggested workflow for using this fix is: 1) identify a reaction to be simulated 2) build a molecule template of the reaction site before the reaction has occurred 3) build a molecule template of the reaction site after the reaction has occurred 4) create a map that relates the template-atom-IDs of each atom between pre- and post-reaction molecule templates 5) fill a simulation box with molecules and run a simulation with `fix bond/react`.

Only one ‘fix bond/react’ command can be used at a time. Multiple reactions can be simultaneously applied by specifying multiple *react* arguments to a single ‘fix bond/react’ command. This syntax is necessary because the ‘common keywords’ are applied to all reactions.

The *stabilization* keyword enables reaction site stabilization. Reaction site stabilization is performed by including reacting atoms in an internally-created fix *nve/limit* time integrator for a set number of timesteps given by the *stabilize\_steps* keyword. While reacting atoms are being time integrated by the internal *nve/limit*, they are prevented from being involved in any new reactions. The *xmax* value keyword should typically be set to the maximum distance that non-reacting atoms move during the simulation.

Fix bond/react creates and maintains two important dynamic groups of atoms when using the *stabilization* keyword. The first group contains all atoms currently involved in a reaction; this group is automatically thermostatted by an internally-created *nve/limit* integrator. The second group contains all atoms currently not involved in a reaction. This group should be used by a thermostat in order to time integrate the system. The name of this group of non-reacting atoms is created by appending ‘\_REACT’ to the group-ID argument of the *stabilization* keyword, as shown in the second example above.

---

**Note:** When using reaction stabilization, you should generally not have a separate thermostat which acts on the ‘all’ group.

---

The group-ID set using the *stabilization* keyword can be an existing static group or a previously-unused group-ID. It cannot be specified as ‘all’. If the group-ID is previously unused, the fix bond/react command creates a *dynamic group* that is initialized to include all atoms. If the group-ID is that of an existing static group, the group is used as the parent group of new, internally-created dynamic group. In both cases, this new dynamic group is named by appending ‘\_REACT’ to the group-ID, e.g. *nvt\_grp\_REACT*. By specifying an existing group, you may thermostat constant-topology parts of your system separately. The dynamic group contains only atoms not involved in a reaction at a given timestep, and therefore should be used by a subsequent system-wide time integrator such as *nvt*, *npt*, or *nve*, as shown in the second example above (full examples can be found at [examples/PACKAGES/reaction](#)). The time integration command should be placed after the fix bond/react command due to the internal dynamic grouping performed by fix bond/react.

---

**Note:** If the group-ID is an existing static group, react-group-IDs should also be specified as this static group, or a subset.

---

The *reset\_mol\_ids* keyword invokes the *reset\_mol\_ids* command after a reaction occurs, to ensure that molecule IDs are consistent with the new bond topology. The group-ID used for *reset\_mol\_ids* is the group-ID for this fix. Resetting molecule IDs is necessarily a global operation, and so can be slow for very large systems.

The following comments pertain to each *react* argument (in other words, can be customized for each reaction, or reaction step):

A check for possible new reaction sites is performed every *Nevery* timesteps. *Nevery* can be specified with an equal-style *variable*, whose value is rounded up to the nearest integer.

Three physical conditions must be met for a reaction to occur. First, an initiator atom pair must be identified within the reaction distance cutoffs. Second, the topology surrounding the initiator atom pair must match the topology of the pre-reaction template. Only atom types and bond connectivity are used to identify a valid reaction site (not bond types, etc.). Finally, any reaction constraints listed in the map file (see below) must be satisfied. If all of these conditions are met, the reaction site is eligible to be modified to match the post-reaction template.

An initiator atom pair will be identified if several conditions are met. First, a pair of atoms I,J within the specified react-group-ID of type *itype* and *jtype* must be separated by a distance between *Rmin* and *Rmax*. *Rmin* and *Rmax* can be specified with equal-style *variables*. For example, these reaction cutoffs can be a function of the reaction conversion using the following commands:



```

variable rmax equal 0 # initialize variable before bond/react
fix myrxn all bond/react react myrxn1 all 1 0 v_rmax mol1 mol2 map_file.txt
variable rmax equal 3+f_myrxn[1]/100 # arbitrary function of reaction count

```

The following criteria are used if multiple candidate initiator atom pairs are identified within the cutoff distance: 1) If the initiator atoms in the pre-reaction template are not 1-2 neighbors (i.e. not directly bonded) the closest potential partner is chosen. 2) Otherwise, if the initiator atoms in the pre-reaction template are 1-2 neighbors (i.e. directly bonded) the farthest potential partner is chosen. 3) Then, if both an atom I and atom J have each other as their initiator partners, these two atoms are identified as the initiator atom pair of the reaction site. Note that it can be helpful to select unique atom types for the initiator atoms: if an initiator atom pair is identified, as described in the previous steps, but does not correspond to the same pair specified in the pre-reaction template, an otherwise eligible reaction could be prevented from occurring. Once this unique initiator atom pair is identified for each reaction, there could be two or more reactions that involve the same atom on the same timestep. If this is the case, only one such reaction is permitted to occur. This reaction is chosen randomly from all potential reactions involving the overlapping atom. This capability allows e.g. for different reaction pathways to proceed from identical reaction sites with user-specified probabilities.

The pre-reacted molecule template is specified by a molecule command. This molecule template file contains a sample reaction site and its surrounding topology. As described below, the initiator atom pairs of the pre-reacted template are specified by atom ID in the map file. The pre-reacted molecule template should contain as few atoms as possible while still completely describing the topology of all atoms affected by the reaction (which includes all atoms that change atom type or connectivity, and all bonds that change bond type). For example, if the force field contains dihedrals, the pre-reacted template should contain any atom within three bonds of reacting atoms.

Some atoms in the pre-reacted template that are not reacting may have missing topology with respect to the simulation. For example, the pre-reacted template may contain an atom that, in the simulation, is currently connected to the rest of a long polymer chain. These are referred to as edge atoms, and are also specified in the map file. All pre-reaction template atoms should be linked to an initiator atom, via at least one path that does not involve edge atoms. When the pre-reaction template contains edge atoms, not all atoms, bonds, charges, etc. specified in the reaction templates will be updated. Specifically, topology that involves only atoms that are ‘too near’ to template edges will not be updated. The definition of ‘too near the edge’ depends on which interactions are defined in the simulation. If the simulation has defined dihedrals, atoms within two bonds of edge atoms are considered ‘too near the edge.’ If the simulation defines angles, but not dihedrals, atoms within one bond of edge atoms are considered ‘too near the edge.’ If just bonds are defined, only edge atoms are considered ‘too near the edge.’

---

**Note:** Small molecules, i.e. ones that have all their atoms contained within the reaction templates, never have edge atoms.

---

Note that some care must be taken when building a molecule template for a given simulation. All atom types in the pre-reacted template must be the same as those of a potential reaction site in the simulation. A detailed discussion of matching molecule template atom types with the simulation is provided on the [molecule](#) command page.

The post-reacted molecule template contains a sample of the reaction site and its surrounding topology after the reaction has occurred. It must contain the same number of atoms as the pre-reacted template (unless there are created atoms). A one-to-one correspondence between the atom IDs in the pre- and post-reacted templates is specified in the map file as described below. Note that during a reaction, an atom, bond, etc. type may change to one that was previously not present in the simulation. These new types must also be defined during the setup of a given simulation. A discussion of correctly handling this is also provided on the [molecule](#) command page.

---

**Note:** When a reaction occurs, it is possible that the resulting topology/atom (e.g. special bonds, dihedrals, etc.) exceeds that of the existing system and reaction templates. As when inserting molecules, enough space for this increased topology/atom must be reserved by using the relevant “extra” keywords to the [read\\_data](#) or [create\\_box](#) commands.

---

The map file is a text document with the following format:

A map file has a header and a body. The header of map file the contains one mandatory keyword and five optional keywords. The mandatory keyword is 'equivalences':

```
N equivalences = # of atoms N in the reaction molecule templates
```

The optional keywords are 'edgeIDs', 'deleteIDs', 'chiralIDs' and 'constraints':

```
N edgeIDs = # of edge atoms N in the pre-reacted molecule template
```

```
N deleteIDs = # of atoms N that are deleted
```

```
N createIDs = # of atoms N that are created
```

```
N chiralIDs = # of chiral centers N
```

```
N constraints = # of reaction constraints N
```

The body of the map file contains two mandatory sections and five optional sections. The first mandatory section begins with the keyword 'InitiatorIDs' and lists the two atom IDs of the initiator atom pair in the pre-reacted molecule template. The second mandatory section begins with the keyword 'Equivalences' and lists a one-to-one correspondence between atom IDs of the pre- and post-reacted templates. The first column is an atom ID of the pre-reacted molecule template, and the second column is the corresponding atom ID of the post-reacted molecule template. The first optional section begins with the keyword 'EdgeIDs' and lists the atom IDs of edge atoms in the pre-reacted molecule template. The second optional section begins with the keyword 'DeleteIDs' and lists the atom IDs of pre-reaction template atoms to delete. The third optional section begins with the keyword 'CreateIDs' and lists the atom IDs of the post-reaction template atoms to create. The fourth optional section begins with the keyword 'ChiralIDs' lists the atom IDs of chiral atoms whose handedness should be enforced. The fifth optional section begins with the keyword 'Constraints' and lists additional criteria that must be satisfied in order for the reaction to occur. Currently, there are six types of constraints available, as discussed below: 'distance', 'angle', 'dihedral', 'arrhenius', 'rmsd', and 'custom'.

A sample map file is given below:

---

```
# this is a map file

7 equivalences
2 edgeIDs

InitiatorIDs

3
5

EdgeIDs

1
7

Equivalences

1 1
2 2
3 3
4 4
5 5
6 6
7 7
```

---

A user-specified set of atoms can be deleted by listing their pre-reaction template IDs in the DeleteIDs section. A deleted atom must still be included in the post-reaction molecule template, in which it cannot be bonded to an atom

that is not deleted. In addition to deleting unwanted reaction by-products, this feature can be used to remove specific topologies, such as small rings, that may be otherwise indistinguishable.

Atoms can be created by listing their post-reaction template IDs in the CreateIDs section. A created atom should not be included in the pre-reaction template. The inserted positions of created atoms are determined by the coordinates of the post-reaction template, after optimal translation and rotation of the post-reaction template to the reaction site (using a fit with atoms that are neither created nor deleted). The *modify\_create* keyword can be used to modify the default behavior when creating atoms. The *modify\_create* keyword has two sub-keywords, *fit* and *overlap*. One or more of the sub-keywords may be used after the *modify\_create* keyword. The *fit* sub-keyword can be used to specify which post-reaction atoms are used for the optimal translation and rotation of the post-reaction template. The *fragmentID* value of the *fit* sub-keyword must be the name of a molecule fragment defined in the post-reaction *molecule* template, and only atoms in this fragment are used for the fit. Atoms are created only if no current atom in the simulation is within a distance R of any created atom, including the effect of periodic boundary conditions if applicable. R is defined by the *overlap* sub-keyword. Note that the default value for R is 0.0, which will allow atoms to strongly overlap if you are inserting where other atoms are present. The velocity of each created atom is initialized in a random direction with a magnitude calculated from the instantaneous temperature of the reaction site.

The handedness of atoms that are chiral centers can be enforced by listing their IDs in the ChiralIDs section. A chiral atom must be bonded to four atoms with mutually different atom types. This feature uses the coordinates and types of the involved atoms in the pre-reaction template to determine handedness. Three atoms bonded to the chiral center are arbitrarily chosen, to define an oriented plane, and the relative position of the fourth bonded atom determines the chiral center's handedness.

Any number of additional constraints may be specified in the Constraints section of the map file. The constraint of type 'distance' has syntax as follows:

```
distance ID1 ID2 rmin rmax
```

where 'distance' is the required keyword, *ID1* and *ID2* are pre-reaction atom IDs (or molecule-fragment IDs, see below), and these two atoms must be separated by a distance between *rmin* and *rmax* for the reaction to occur.

The constraint of type 'angle' has the following syntax:

```
angle ID1 ID2 ID3 amin amax
```

where 'angle' is the required keyword, *ID1*, *ID2* and *ID3* are pre-reaction atom IDs (or molecule-fragment IDs, see below), and these three atoms must form an angle between *amin* and *amax* for the reaction to occur (where *ID2* is the central atom). Angles must be specified in degrees. This constraint can be used to enforce a certain orientation between reacting molecules.

The constraint of type 'dihedral' has the following syntax:

```
dihedral ID1 ID2 ID3 ID4 amin amax amin2 amax2
```

where 'dihedral' is the required keyword, and *ID1*, *ID2*, *ID3* and *ID4* are pre-reaction atom IDs (or molecule-fragment IDs, see below). Dihedral angles are calculated in the interval (-180,180]. Refer to the *dihedral style* documentation for further details on convention. If *amin* is less than *amax*, these four atoms must form a dihedral angle greater than *amin* **and** less than *amax* for the reaction to occur. If *amin* is greater than *amax*, these four atoms must form a dihedral angle greater than *amin* **or** less than *amax* for the reaction to occur. Angles must be specified in degrees. Optionally, a second range of permissible angles *amin2*-*amax2* can be specified.

For the 'distance', 'angle', and 'dihedral' constraints (explained above), atom IDs can be replaced by pre-reaction molecule-fragment IDs. The molecule-fragment ID must begin with a letter. The location of the ID is the geometric center of all atom positions in the fragment. The molecule fragment must have been defined in the *molecule* command for the pre-reaction template.

The constraint of type 'arrhenius' imposes an additional reaction probability according to the temperature-dependent Arrhenius equation:

$$k = AT^n e^{\frac{-E_a}{k_B T}}$$

The Arrhenius constraint has the following syntax:

```
arrhenius A n E_a seed
```

where ‘arrhenius’ is the required keyword,  $A$  is the pre-exponential factor,  $n$  is the exponent of the temperature dependence,  $E_a$  is the activation energy (*units* of energy), and *seed* is a random number seed. The temperature is defined as the instantaneous temperature averaged over all atoms in the reaction site, and is calculated in the same manner as for example *compute temp/chunk*. Currently, there are no options for additional temperature averaging or velocity-biased temperature calculations. A uniform random number between 0 and 1 is generated using *seed*; if this number is less than the result of the Arrhenius equation above, the reaction is permitted to occur.

The constraint of type ‘rmsd’ has the following syntax:

```
rmsd RMSDmax molfragment
```

where ‘rmsd’ is the required keyword, and  $RMSD_{max}$  is the maximum root-mean-square deviation between atom positions of the pre-reaction template and the local reaction site (distance units), after optimal translation and rotation of the pre-reaction template. Optionally, the name of a molecule fragment (of the pre-reaction template) can be specified by *molfragment*. If a molecule fragment is specified, only atoms that are part of this molecule fragment are used to determine the RMSD. A molecule fragment must have been defined in the *molecule* command for the pre-reaction template. For example, the molecule fragment could consist of only the backbone atoms of a polymer chain. This constraint can be used to enforce a specific relative position and orientation between reacting molecules.

The constraint of type ‘custom’ has the following syntax:

```
custom varstring
```

where ‘custom’ is the required keyword, and *varstring* is a variable expression. The expression must be a valid equal-style variable formula that can be read by the *variable* command, after any special reaction functions are evaluated. If the resulting expression is zero, the reaction is prevented from occurring; otherwise, it is permitted to occur. There are two special reaction functions available, ‘rxnsum’ and ‘rxnave’. These functions operate over the atoms in a given reaction site, and have one mandatory argument and one optional argument. The mandatory argument is the identifier for an atom-style variable. The second, optional argument is the name of a molecule fragment in the pre-reaction template, and can be used to operate over a subset of atoms in the reaction site. The ‘rxnsum’ function sums the atom-style variable over the reaction site, while the ‘rxnave’ returns the average value. For example, a constraint on the total potential energy of atoms involved in the reaction can be imposed as follows:

```
compute 1 all pe/atom # in LAMMPS input script
variable my_pe atom c_1 # in LAMMPS input script
```

```
custom "rxnsum(v_my_pe) > 100" # in Constraints section of map file
```

The above example prevents the reaction from occurring unless the total potential energy of the reaction site is above 100. The variable expression can be interpreted as the probability of the reaction occurring by using an inequality and the ‘random(x,y,z)’ function available as an equal-style variable input, similar to the ‘arrhenius’ constraint above.

By default, all constraints must be satisfied for the reaction to occur. In other words, constraints are evaluated as a series of logical values using the logical AND operator “&&”. More complex logic can be achieved by explicitly adding the logical AND operator “&&” or the logical OR operator “||” after a given constraint command. If a logical operator is specified after a constraint, it must be placed after all constraint parameters, on the same line as the constraint (one per line). Similarly, parentheses can be used to group constraints. The expression that results from concatenating all constraints should be a valid logical expression that can be read by the *variable* command after converting each constraint to a logical value. Because exactly one constraint is allowed per line, having a valid logical expression implies that left parentheses “(” should only appear before a constraint, and right parentheses “)” should only appear after a constraint and before any logical operator.

Once a reaction site has been successfully identified, data structures within LAMMPS that store bond topology are updated to reflect the post-reacted molecule template. All force fields with fixed bonds, angles, dihedrals or impropers are supported.

A few capabilities to note: 1) You may specify as many *react* arguments as desired. For example, you could break down a complicated reaction mechanism into several reaction steps, each defined by its own *react* argument. 2) While typically a bond is formed or removed between the initiator atoms specified in the pre-reacted molecule template, this is not required. 3) By reversing the order of the pre- and post- reacted molecule templates in another *react* argument, you can allow for the possibility of one or more reverse reactions.

The optional keywords deal with the probability of a given reaction occurring as well as the stable equilibration of each reaction site as it occurs:

The *prob* keyword can affect whether or not an eligible reaction actually occurs. The fraction setting must be a value between 0.0 and 1.0, and can be specified with an equal-style *variable*. A uniform random number between 0.0 and 1.0 is generated and the eligible reaction only occurs if the random number is less than the fraction. Up to N reactions are permitted to occur, as optionally specified by the *max\_rxn* keyword.

The *stabilize\_steps* keyword allows for the specification of how many timesteps a reaction site is stabilized before being returned to the overall system thermostat. In order to produce the most physical behavior, this ‘reaction site equilibration time’ should be tuned to be as small as possible while retaining stability for a given system or reaction step. After a limited number of case studies, this number has been set to a default of 60 timesteps. Ideally, it should be individually tuned for each fix reaction step. Note that in some situations, decreasing rather than increasing this parameter will result in an increase in stability.

The *custom\_charges* keyword can be used to specify which atoms’ atomic charges are updated. When the value is set to ‘no’, all atomic charges are updated to those specified by the post-reaction template (default). Otherwise, the value should be the name of a molecule fragment defined in the pre-reaction molecule template. In this case, only the atomic charges of atoms in the molecule fragment are updated.

The *molecule* keyword can be used to force the reaction to be intermolecular, intramolecular or either. When the value is set to ‘off’, molecule IDs are not considered when searching for reactions (default). When the value is set to ‘inter’, the initiator atoms must have different molecule IDs in order to be considered for the reaction. When the value is set to ‘intra’, only initiator atoms with the same molecule ID are considered for the reaction.

A few other considerations:

Optionally, you can enforce additional behaviors on reacting atoms. For example, it may be beneficial to force reacting atoms to remain at a certain temperature. For this, you can use the internally-created dynamic group named “bond\_react\_MASTER\_group”, which consists of all atoms currently involved in a reaction. For example, adding the following command would add an additional thermostat to the group of all currently-reacting atoms:

```
fix 1 bond_react_MASTER_group temp/rescale 1 300 300 10 1
```

**Note:** This command must be added after the fix bond/react command, and will apply to all reactions.

Computationally, each timestep this fix operates, it loops over neighbor lists (for bond-forming reactions) and computes distances between pairs of atoms in the list. It also communicates between neighboring processors to coordinate which bonds are created and/or removed. All of these operations increase the cost of a timestep. Thus you should be cautious about invoking this fix too frequently.

You can dump out snapshots of the current bond topology via the dump local command.

### 17.25.4 Restart, fix\_modify, output, run start/stop, minimize info

Cumulative reaction counts for each reaction are written to *binary restart files*. These values are associated with the reaction name (react-ID). Additionally, internally-created per-atom properties are stored to allow for smooth restarts. None of the *fix\_modify* options are relevant to this fix.

This fix computes one statistic for each *react* argument that it stores in a global vector, of length ‘number of react arguments’, that can be accessed by various *output commands*. The vector values calculated by this fix are “intensive”.

There is 1 quantity for each react argument:

- (1) cumulative # of reactions occurred

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

When *fix bond/react* is ‘unfixed’, all internally-created groups are deleted. Therefore, *fix bond/react* can only be unfixed after unfixing all other fixes that use any group created by *fix bond/react*.

### 17.25.5 Restrictions

This fix is part of the REACTION package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.25.6 Related commands

*fix bond/create, fix bond/break, fix bond/swap, dump local, special\_bonds*

### 17.25.7 Default

The option defaults are *stabilization = no, prob = 1.0, stabilize\_steps = 60, reset\_mol\_ids = yes, custom\_charges = no, molecule = off, modify\_create = no*

---

(Gissinger2017) Gissinger, Jensen and Wise, *Polymer*, 128, 211-217 (2017).

(Gissinger2020) Gissinger, Jensen and Wise, *Macromolecules*, 53, 22, 9953-9961 (2020).

## 17.26 fix bond/swap command

### 17.26.1 Syntax

```
fix ID group-ID bond/swap Nevery fraction cutoff seed
```

- ID, group-ID are documented in *fix* command
- bond/swap = style name of this fix command
- Nevery = attempt bond swapping every this many steps
- fraction = fraction of group atoms to consider for swapping
- cutoff = distance at which swapping will be considered (distance units)
- seed = random # seed (positive integer)

## 17.26.2 Examples

```
fix 1 all bond/swap 50 0.5 1.3 598934
```

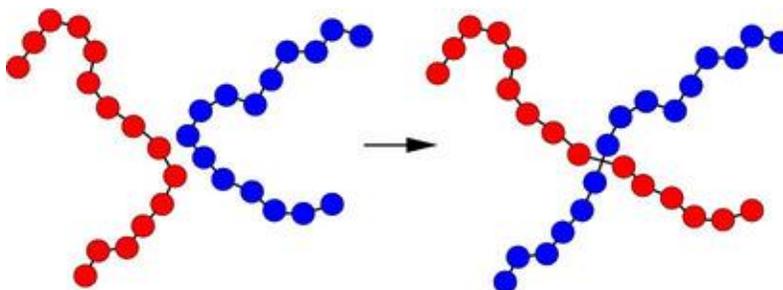
### 17.26.3 Description

In a simulation of polymer chains this command attempts to swap a pair of bonds, as illustrated below. This is done via Monte Carlo rules using the Boltzmann acceptance criterion, typically with the goal of equilibrating the polymer system more quickly. This fix is designed for use with idealized bead-spring polymer chains where each polymer is a linear chain of monomers, but LAMMPS does not check that is the case for your system.

Here are two use cases for this fix.

The first use case is for swapping bonds on two different chains, effectively grafting the end of one chain onto the other chain and vice versa. The purpose is to equilibrate the polymer chain conformations more rapidly than dynamics alone would do it, by enabling instantaneous large conformational changes in a dense polymer melt. The polymer chains should thus more rapidly converge to the proper end-to-end distances and radii of gyration.

A schematic of the kinds of bond swaps that can occur in this use case is shown here:



On the left, the red and blue chains have two monomers A1 and B1 close to each other, which are currently bonded to monomers A2 and B2 respectively within their own chains. The bond swap operation will attempt to delete the A1-A2 and B1-B2 bonds and replace them with A1-B2 and B1-A2 bonds. If the swap is energetically favorable, the two chains on the right are the result and each polymer chain has undergone a dramatic conformational change. This reference, (*Sides*) provides more details on the algorithm's effectiveness for this use case.

The second use case is a collection of polymer chains with some fraction of their sites identified as “sticker” sites. Initially each polymer chain is isolated from the others in a topological sense, and there is an intra-chain bond between every pair of sticker sites on the same chain. Over time, bonds swap so that inter-molecular sticker bonds are created. This models a vitrification-style process whereby the polymer chains all become interconnected. For this use case, if angles are defined they should not include bonds between sticker sites.

---

**Note:** For the sticker site model, you should set the newton flag for bonds to “off”, via the *newton on off* command (“on” is the default for the 2nd argument). This is to ensure appropriate randomness in bond selection because the I,J bond will be stored by both atom I and atom J. LAMMPS cannot check for this, because it is not aware that a sticker site model is being used.

---

The bond swapping operation is invoked once every *Nevery* timesteps. If any bond in the entire system is swapped, a re-build of the neighbor lists is triggered, since a swap alters the list of which neighbors are considered for pairwise interaction. At each invocation, each processor considers a random specified *fraction* of its atoms as potential swapping monomers for this timestep. Choosing a small *fraction* value can reduce the likelihood of a reverse swap occurring soon after an initial swap.

For each monomer A1, its neighbors are looped over as B1 monomers. For each A1,B1 an additional double loop of bond partners A2 of A1, and bond partners B2 of B1 is performed. For each pair of A1-A2 and B1-B2 bonds to be eligible for swapping, the following 4 criteria must be met:

1. All 4 monomers must be in the fix group.
2. All 4 monomers must be owned by the processor (not ghost atoms). This insures that another processor does not attempt to swap bonds involving the same atoms on the same timestep. Note that this also means that bond pairs which straddle processor boundaries are not eligible for swapping on this step.
3. The distances between 4 pairs of atoms – (A1,A2), (B1,B2), (A1,B2), (B1,A2) – must all be less than the specified *cutoff*.
4. The molecule IDs of A1 and B1 must be the same (see below).

If an eligible B1 partner is found, the energy change due to swapping the 2 bonds is computed. This includes changes in pairwise, bond, and angle energies due to the altered connectivity of the 2 chains. Dihedral and improper interactions are not allowed to be defined when this fix is used.

If the energy decreases due to the swap operation, the bond swap is accepted. If the energy increases it is accepted with probability  $\exp(-\delta/kT)$  where  $\delta$  is the increase in energy,  $k$  is the Boltzmann constant, and  $T$  is the current temperature of the system.

---

**Note:** Whether the swap is accepted or rejected, no other swaps are attempted by this processor on this timestep. No other eligible 4-tuples of atoms are considered. This means that each processor will perform either a single swap or none on timesteps this fix is invoked.

---

The criterion for matching molecule IDs is how the first use case described above can be simulated while conserving chain lengths. This is done by setting up the molecule IDs for the polymer chains in a specific way, typically in the data file, read by the *read\_data* command.

Consider a system of 6-mer chains. You have 2 choices. If the molecule IDs for monomers on each chain are set to 1,2,3,4,5,6 then swaps will conserve chain length. For a particular monomer there will be only one other monomer on another chain which is a potential swap partner. If the molecule IDs for monomers on each chain are set to 1,2,3,3,2,1 then swaps will conserve chain length but swaps will be able to occur at either end of a chain. Thus for a particular monomer there will be 2 possible swap partners on another chain. In this scenario, swaps can also occur within a single chain, i.e. the two ends of a chain swap with each other.

---

**Note:** If your simulation uses molecule IDs in the usual way, where all monomers on a single chain are assigned the same ID (different for each chain), then swaps will only occur within the same chain. If you assign the same molecule ID to all monomers in all chains then inter-chain swaps will occur, but they will not conserve chain length. Neither of these scenarios is probably what you want for this fix.

---

---

**Note:** When a bond swap occurs the image flags of monomers in the new polymer chains can become inconsistent. See the *dump* command for a discussion of image flags. This is not an issue for running dynamics, but can affect calculation of some diagnostic quantities or the printing of unwrapped coordinates to a dump file.

---

For the second use case described above, the molecule IDs for all sticker sites should be the same.

---

This fix computes a temperature each time it is invoked for use by the Boltzmann criterion. To do this, the fix creates its own compute of style *temp*, as if this command had been issued:



```
compute fix-ID_temp all temp
```

See the *compute temp* command for details. Note that the ID of the new compute is the fix-ID with underscore + “temp” appended and the group for the new compute is “all”, so that the temperature of the entire system is used.

Note that this is NOT the compute used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp*. This means you can change the attributes of this fix’s temperature (e.g. its degrees-of-freedom) via the *compute\_modify* command or print this temperature during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* will have no effect on this fix.

#### 17.26.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. Because the state of the random number generator is not saved in restart files, this means you cannot do “exact” restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior. Also note that each processor generates possible swaps independently of other processors. Thus if you repeat the same simulation on a different number of processors, the specific swaps performed will be different.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a *compute* you have defined to this fix which will be used to compute the temperature for the Boltzmann criterion.

This fix computes two statistical quantities as a global 2-vector of output, which can be accessed by various *output commands*. The first component of the vector is the cumulative number of swaps performed by all processors. The second component of the vector is the cumulative number of swaps attempted (whether accepted or rejected). Note that a swap “attempt” only occurs when swap partners meeting the criteria described above are found on a particular timestep. The vector values calculated by this fix are “intensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.26.5 Restrictions

This fix is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

The settings of the “special\_bond” command must be 0,1,1 in order to use this fix, which is typical of bead-spring chains with FENE or harmonic bonds. This means that pairwise interactions between bonded atoms are turned off, but are turned on between atoms two or three hops away along the chain backbone.

Currently, energy changes in dihedral and improper interactions due to a bond swap are not considered. Thus a simulation that uses this fix cannot use a dihedral or improper potential.

#### 17.26.6 Related commands

*fix atom/swap*

## 17.26.7 Default

none

---

(Sides) Sides, Grest, Stevens, Plimpton, J Polymer Science B, 42, 199-208 (2004).

## 17.27 fix box/relax command

### 17.27.1 Syntax

```
fix ID group-ID box/relax keyword value ...
```

- ID, group-ID are documented in *fix* command
- box/relax = style name of this fix command

one or more keyword value pairs may be appended

keyword = *iso* or *aniso* or *tri* or *x* or *y* or *z* or *xy* or *yz* or *xz* or *couple* or *nreset*.

→ or *vmax* or *dilate* or *scaleyz* or *scalexz* or *scalexy* or *fixedpoint*

*iso* or *aniso* or *tri* value = Ptarget = desired pressure (pressure units)

*x* or *y* or *z* or *xy* or *yz* or *xz* value = Ptarget = desired pressure (pressure units)

*couple* = *none* or *xyz* or *xy* or *yz* or *xz*

*nreset* value = reset reference cell every this many minimizer iterations

*vmax* value = fraction = max allowed volume change in one iteration

*dilate* value = *all* or *partial*

*scaleyz* value = *yes* or *no* = scale yz with lz

*scalexz* value = *yes* or *no* = scale xz with lz

*scalexy* value = *yes* or *no* = scale xy with ly

*fixedpoint* values = x y z

x,y,z = perform relaxation dilation/contraction around this point (distance.  
→units)

### 17.27.2 Examples

```
fix 1 all box/relax iso 0.0 vmax 0.001
fix 2 water box/relax aniso 0.0 dilate partial
fix 2 ice box/relax tri 0.0 couple xy nreset 100
```

### 17.27.3 Description

Apply an external pressure or stress tensor to the simulation box during an *energy minimization*. This allows the box size and shape to vary during the iterations of the minimizer so that the final configuration will be both an energy minimum for the potential energy of the atoms, and the system pressure tensor will be close to the specified external tensor. Conceptually, specifying a positive pressure is like squeezing on the simulation box; a negative pressure typically allows the box to expand.

---

The external pressure tensor is specified using one or more of the *iso*, *aniso*, *tri*, *x*, *y*, *z*, *xy*, *xz*, *yz*, and *couple* keywords. These keywords give you the ability to specify all 6 components of an external stress tensor, and to couple various of these components together so that the dimensions they represent are varied together during the minimization.

Orthogonal simulation boxes have 3 adjustable dimensions ( $x,y,z$ ). Triclinic (non-orthogonal) simulation boxes have 6 adjustable dimensions ( $x,y,z,xy,xz,yz$ ). The `create_box`, `read_data`, and `read_restart` commands specify whether the simulation box is orthogonal or non-orthogonal (triclinic) and explain the meaning of the  $xy,xz,yz$  tilt factors.

The target pressures  $P_{target}$  for each of the 6 components of the stress tensor can be specified independently via the  $x$ ,  $y$ ,  $z$ ,  $xy$ ,  $xz$ ,  $yz$  keywords, which correspond to the 6 simulation box dimensions. For example, if the  $y$  keyword is used, the  $y$ -box length will change during the minimization. If the  $xy$  keyword is used, the  $xy$  tilt factor will change. A box dimension will not change if that component is not specified.

Note that in order to use the  $xy$ ,  $xz$ , or  $yz$  keywords, the simulation box must be triclinic, even if its initial tilt factors are 0.0.

When the size of the simulation box changes, all atoms are re-scaled to new positions, unless the keyword `dilate` is specified with a value of `partial`, in which case only the atoms in the `fix` group are re-scaled. This can be useful for leaving the coordinates of atoms in a solid substrate unchanged and controlling the pressure of a surrounding fluid.

The `scaleyz`, `scalexz`, and `scalexy` keywords control whether or not the corresponding tilt factors are scaled with the associated box dimensions when relaxing triclinic periodic cells. The default values `yes` will turn on scaling, which corresponds to adjusting the linear dimensions of the cell while preserving its shape. Choosing `no` ensures that the tilt factors are not scaled with the box dimensions. See below for restrictions and default values in different situations. In older versions of LAMMPS, scaling of tilt factors was not performed. The old behavior can be recovered by setting all three scale keywords to `no`.

The `fixedpoint` keyword specifies the fixed point for cell relaxation. By default, it is the center of the box. Whatever point is chosen will not move during the simulation. For example, if the lower periodic boundaries pass through (0,0,0), and this point is provided to `fixedpoint`, then the lower periodic boundaries will remain at (0,0,0), while the upper periodic boundaries will move twice as far. In all cases, the particle positions at each iteration are unaffected by the chosen value, except that all particles are displaced by the same amount, different on each iteration.

---

**Note:** Applying an external pressure to tilt dimensions  $xy$ ,  $xz$ ,  $yz$  can sometimes result in arbitrarily large values of the tilt factors, i.e. a dramatically deformed simulation box. This typically indicates that there is something badly wrong with how the simulation was constructed. The two most common sources of this error are applying a shear stress to a liquid system or specifying an external shear stress tensor that exceeds the yield stress of the solid. In either case the minimization may converge to a bogus conformation or not converge at all. Also note that if the box shape tilts to an extreme shape, LAMMPS will run less efficiently, due to the large volume of communication needed to acquire ghost atoms around a processor's irregular-shaped sub-domain. For extreme values of tilt, LAMMPS may also lose atoms and generate an error.

---

---

**Note:** Performing a minimization with this `fix` is not a mathematically well-defined minimization problem. This is because the objective function being minimized changes if the box size/shape changes. In practice this means the minimizer can get "stuck" before you have reached the desired tolerance. The solution to this is to restart the minimizer from the new adjusted box size/shape, since that creates a new objective function valid for the new box size/shape. Repeat as necessary until the box size/shape has reached its new equilibrium.

---

The `couple` keyword allows two or three of the diagonal components of the pressure tensor to be "coupled" together. The value specified with the keyword determines which are coupled. For example,  $xz$  means the  $P_{xx}$  and  $P_{zz}$  components of the stress tensor are coupled.  $xyz$  means all 3 diagonal components are coupled. Coupling means two things: the instantaneous stress will be computed as an average of the corresponding diagonal components, and the coupled box dimensions will be changed together in lockstep, meaning coupled dimensions will be dilated or contracted by the same percentage every timestep. The  $P_{target}$  values for any coupled dimensions must be identical. `Couple xyz` can be used for a 2d simulation; the  $z$  dimension is simply ignored.

---

The *iso*, *aniso*, and *tri* keywords are simply shortcuts that are equivalent to specifying several other keywords together.

The keyword *iso* means couple all 3 diagonal components together when pressure is computed (hydrostatic pressure), and dilate/contract the dimensions together. Using “iso Ptarget” is the same as specifying these 4 keywords:

```
x Ptarget
y Ptarget
z Ptarget
couple xyz
```

The keyword *aniso* means *x*, *y*, and *z* dimensions are controlled independently using the  $P_{xx}$ ,  $P_{yy}$ , and  $P_{zz}$  components of the stress tensor as the driving forces, and the specified scalar external pressure. Using “aniso Ptarget” is the same as specifying these 4 keywords:

```
x Ptarget
y Ptarget
z Ptarget
couple none
```

The keyword *tri* means *x*, *y*, *z*, *xy*, *xz*, and *yz* dimensions are controlled independently using their individual stress components as the driving forces, and the specified scalar pressure as the external normal stress. Using “tri Ptarget” is the same as specifying these 7 keywords:

```
x Ptarget
y Ptarget
z Ptarget
xy 0.0
yz 0.0
xz 0.0
couple none
```

---

The *vmax* keyword can be used to limit the fractional change in the volume of the simulation box that can occur in one iteration of the minimizer. If the pressure is not settling down during the minimization this can be because the volume is fluctuating too much. The specified fraction must be greater than 0.0 and should be  $\ll 1.0$ . A value of 0.001 means the volume cannot change by more than 1/10 of a percent in one iteration when *couple xyz* has been specified. For any other case it means no linear dimension of the simulation box can change by more than 1/10 of a percent.

---

With this fix, the potential energy used by the minimizer is augmented by an additional energy provided by the fix. The overall objective function then is:

$$E = U + P_t (V - V_0) + E_{strain}$$

where  $U$  is the system potential energy,  $P_t$  is the desired hydrostatic pressure,  $V$  and  $V_0$  are the system and reference volumes, respectively.  $E_{strain}$  is the strain energy expression proposed by Parrinello and Rahman ([Parrinello1981](#)). Taking derivatives of  $E$  w.r.t. the box dimensions, and setting these to zero, we find that at the minimum of the objective function, the global system stress tensor  $\mathbf{P}$  will satisfy the relation:

$$\mathbf{P} = P_t \mathbf{I} + \mathbf{S}_t (\mathbf{h}_0^{-1})^t \mathbf{h}_{0d}$$

where  $\mathbf{I}$  is the identity matrix,  $\mathbf{h}_0$  is the box dimension tensor of the reference cell, and  $:\mathbf{h}_{0d}$  is the diagonal part of  $\mathbf{h}_0$ .  $\mathbf{S}_t$  is a symmetric stress tensor that is chosen by LAMMPS so that the upper-triangular components of  $\mathbf{P}$  equal the stress tensor specified by the user.

This equation only applies when the box dimensions are equal to those of the reference dimensions. If this is not the case, then the converged stress tensor will not equal that specified by the user. We can resolve this problem by periodically resetting the reference dimensions. The keyword *nreset* controls how often this is done. If this keyword is not used, or is given a value of zero, then the reference dimensions are set to those of the initial simulation domain and are never changed. A value of *nstep* means that every *nstep* minimization steps, the reference dimensions are set to those of the current simulation domain. Note that resetting the reference dimensions changes the objective function and gradients, which sometimes causes the minimization to fail. This can be resolved by changing the value of *nreset*, or simply continuing the minimization from a restart file.

---

**Note:** As normally computed, pressure includes a kinetic- energy or temperature-dependent component; see the *compute pressure* command. However, atom velocities are ignored during a minimization, and the applied pressure(s) specified with this command are assumed to only be the virial component of the pressure (the non-kinetic portion). Thus if atoms have a non-zero temperature and you print the usual thermodynamic pressure, it may not appear the system is converging to your specified pressure. The solution for this is to either (a) zero the velocities of all atoms before performing the minimization, or (b) make sure you are monitoring the pressure without its kinetic component. The latter can be done by outputting the pressure from the *pressure compute* this command creates (see below) or a *pressure compute* you define yourself.

---

**Note:** Because pressure is often a very sensitive function of volume, it can be difficult for the minimizer to equilibrate the system the desired pressure with high precision, particularly for solids. Some techniques that seem to help are (a) use the “min\_modify line quadratic” option when minimizing with box relaxations, (b) minimize several times in succession if need be, to drive the pressure closer to the target pressure, (c) relax the atom positions before relaxing the box, and (d) relax the box to the target hydrostatic pressure before relaxing to a target shear stress state. Also note that some systems (e.g. liquids) will not sustain a non-hydrostatic applied pressure, which means the minimizer will not converge.

---

This fix computes a temperature and pressure each timestep. The temperature is used to compute the kinetic contribution to the pressure, even though this is subsequently ignored by default. To do this, the fix creates its own computes of style “temp” and “pressure”, as if these commands had been issued:

```
compute fix-ID_temp group-ID temp
compute fix-ID_press group-ID pressure fix-ID_temp virial
```

See the *compute temp* and *compute pressure* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix\_ID + underscore + “press”, and the group for the new computes is the same as the fix group. Also note that the pressure compute does not include a kinetic component.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of this fix’s temperature or pressure via the *compute\_modify* command or print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this fix.

---

### 17.27.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify temp* and *press* options are supported by this fix. You can use them to assign a *compute* you have defined to this fix which will be used in its temperature and pressure calculation, as described above. Note that as described above, if you assign a pressure compute to this fix that includes a kinetic energy component it will affect the minimization, most likely in an undesirable way.

---

**Note:** If both the *temp* and *press* keywords are used in a single *thermo\_modify* command (or in two separate commands), then the order in which the keywords are specified is important. Note that a *pressure compute* defines its own temperature compute as an argument when it is specified. The *temp* keyword will override this (for the pressure compute being used by *fix box/relax*), but only if the *temp* keyword comes after the *press* keyword. If the *temp* keyword comes before the *press* keyword, then the new pressure compute specified by the *press* keyword will be unaffected by the *temp* setting.

---

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the pressure-volume energy, plus the strain energy, if it exists, as described above. The energy values reported at the end of a minimization run under “Minimization stats” include this energy, and so differ from what LAMMPS normally reports as potential energy. This fix does not support the *fix\_modify energy* option, because that would result in double-counting of the fix energy in the minimization energy. Instead, the fix energy can be explicitly added to the potential energy using one of these two variants:

```
variable emin equal pe+f_1
variable emin equal pe+f_1/atoms
```

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

This fix is invoked during *energy minimization*, but not for the purpose of adding a contribution to the energy or forces being minimized. Instead it alters the simulation box geometry as described above.

### 17.27.5 Restrictions

Only dimensions that are available can be adjusted by this fix. Non-periodic dimensions are not available. *z*, *xz*, and *yz*, are not available for 2D simulations. *xy*, *xz*, and *yz* are only available if the simulation domain is non-orthogonal. The *create\_box*, *read\_data*, and *read\_restart* commands specify whether the simulation box is orthogonal or non-orthogonal (triclinic) and explain the meaning of the *xy,xz,yz* tilt factors.

The *scaleyz yes* and *scalexz yes* keyword/value pairs can not be used for 2D simulations. *scaleyz yes*, *scalexz yes*, and *scalexy yes* options can only be used if the second dimension in the keyword is periodic, and if the tilt factor is not coupled to the barostat via keywords *tri*, *yz*, *xz*, and *xy*.

### 17.27.6 Related commands

*fix npt*, *minimize*

### 17.27.7 Default

The keyword defaults are dilate = all, vmax = 0.0001, nreset = 0.

(Parrinello1981) Parrinello and Rahman, J Appl Phys, 52, 7182 (1981).

## 17.28 fix brownian command

### 17.29 fix brownian/sphere command

### 17.30 fix brownian/asphere command

#### 17.30.1 Syntax

```
fix ID group-ID style_name temp seed keyword args
```

- ID, group-ID are documented in *fix* command
- style\_name = *brownian* or *brownian/sphere* or *brownian/asphere*
- temp = temperature
- seed = random number generator seed
- one or more keyword/value pairs may be appended
- keyword = *rng* or *dipole* or *gamma\_r\_eigen* or *gamma\_t\_eigen* or *gamma\_r* or *gamma\_t* or *rotation\_temp* or *planar\_rotation*

*rng* value = *uniform* or *gaussian* or *none*

*uniform* = use uniform random number generator

*gaussian* = use gaussian random number generator

*none* = turn off noise

*dipole* value = *mux* and *muy* and *muz* for *brownian/asphere*

*mux*, *muy*, and *muz* = update orientation of dipole having direction (*mux*,\**muy*\*,  
↪\**muz*\*) in body frame of rigid body

*gamma\_r\_eigen* values = *gr1* and *gr2* and *gr3* for *brownian/asphere*

*gr1*, *gr2*, and *gr3* = diagonal entries of body frame rotational friction tensor

*gamma\_r* values = *gr* for *brownian/sphere*

*gr* = magnitude of the (isotropic) rotational friction tensor

*gamma\_t\_eigen* values = *gt1* and *gt2* and *gt3* for *brownian/asphere*

*gt1*, *gt2*, and *gt3* = diagonal entries of body frame translational friction tensor

*gamma\_t* values = *gt* for *brownian* and *brownian/sphere*

*gt* = magnitude of the (isotropic) translational friction tensor

*rotation\_temp* values = *T* for *brownian/sphere* and *brownian/asphere*

*T* = rotation temperature, which can be different then *temp* when out of equilibrium

*planar\_rotation* values = None (constrains rotational diffusion to be in xy plane if,  
↪in 3D)

## 17.30.2 Examples

```

fix 1 all brownian 1.0 12908410 gamma_t 1.0
fix 1 all brownian 1.0 12908410 gamma_t 3.0 rng gaussian
fix 1 all brownian/sphere 1.0 1294019 gamma_t 3.0 gamma_r 1.0
fix 1 all brownian/sphere 1.0 19581092 gamma_t 1.0 gamma_r 0.3 rng none
fix 1 all brownian/asphere 1.0 1294019 gamma_t_eigen 1.0 2.0 3.0 gamma_r_eigen 4.0 7.0 8.
↪0 rng gaussian
fix 1 all brownian/asphere 1.0 1294019 gamma_t_eigen 1.0 2.0 3.0 gamma_r_eigen 4.0 7.0 8.
↪0 dipole 1.0 0.0 0.0

```

## 17.30.3 Description

Perform Brownian Dynamics time integration to update position, velocity, dipole orientation (for spheres) and quaternion orientation (for ellipsoids, with optional dipole update as well) of all particles in the fix group in each timestep. Brownian Dynamics uses Newton's laws of motion in the limit that inertial forces are negligible compared to viscous forces. The stochastic equation of motion for the center of mass positions is

$$d\mathbf{r} = \gamma_t^{-1} \mathbf{F} dt + \sqrt{2k_B T \gamma_t^{-1/2}} d\mathbf{W}_t,$$

in the lab-frame (i.e.  $\gamma_t$  is not diagonal, but only depends on orientation and so the noise is still additive).

The rotational motion for the spherical and ellipsoidal particles is not as simple an expression, but is chosen to replicate the Boltzmann distribution for the case of conservative torques (see *(Ilie)* or *(DeLong)*).

For the style *brownian*, only the positions of the particles are updated. This is therefore suitable for point particle simulations.

For the style *brownian/sphere*, the positions of the particles are updated, and a dipole slaved to the spherical orientation is also updated. This style therefore requires the hybrid atom style *atom\_style dipole* and *atom\_style sphere*. The equation of motion for the dipole is

$$\mu(t + dt) = \frac{\mu(t) + \omega \times \mu dt}{|\mu(t) + \omega \times \mu|}$$

which correctly reproduces a Boltzmann distribution of orientations and rotational diffusion moments (see *(Ilie)*) when

$$\omega = \frac{\mathbf{T}}{\gamma_r} + \sqrt{\frac{2k_B T_{rot}}{\gamma_r}} \frac{d\mathbf{W}}{dt},$$

with  $d\mathbf{W}$  being a random number with zero mean and variance  $dt$  and  $T_{rot}$  is *rotation\_temp*.

For the style *brownian/asphere*, the center of mass positions and the quaternions of ellipsoidal particles are updated. This fix style is suitable for equations of motion where the rotational and translational friction tensors can be diagonalized in a certain (body) reference frame. In this case, the rotational equation of motion is updated via the quaternion

$$\mathbf{q}(t + dt) = \frac{\mathbf{q}(t) + d\mathbf{q}}{|\mathbf{q}(t) + d\mathbf{q}|}$$

which correctly reproduces a Boltzmann distribution of orientations and rotational diffusion moments (see *(Ilie)*) when the quaternion step given by

$$d\mathbf{q} = \Psi \omega dt$$

where  $\Psi$  has rows  $(-q_1, -q_2, -q_3)$ ,  $(q_0, -q_3, q_2)$ ,  $(q_3, q_0, -q_1)$ , and  $(-q_2, q_1, q_0)$ .  $\omega$  is evaluated in the body frame of reference where the friction tensor is diagonal. See *(DeLong)* for more details of a similar algorithm.



---

**Note:** This integrator does not by default assume a relationship between the rotational and translational friction tensors, though such a relationship should exist in the case of no-slip boundary conditions between the particles and the surrounding (implicit) solvent. E.g. in the case of spherical particles, the condition  $\gamma_t = 3\gamma_r/\sigma^2$  must be explicitly accounted for by setting *gamma\_t* to 3x and *gamma\_r* to x (where  $\sigma$  is the spherical diameter). A similar (though more complex) relationship holds for ellipsoids and rod-like particles. The translational diffusion and rotational diffusion are given by *temp/gamma\_t* and *rotation\_temp/gamma\_r*.

---

---

**Note:** Temperature computation using the *compute temp* will not correctly compute temperature of these overdamped dynamics since we are explicitly neglecting inertial effects. Furthermore, this time integrator does not add the stochastic terms or viscous terms to the force and/or torques. Rather, they are just added in to the equations of motion to update the degrees of freedom.

---

---

If the *rng* keyword is used with the *uniform* value, then the noise is generated from a uniform distribution (see (*Dunweg*) for why this works). This is the same method of noise generation as used in *fix langevin*.

If the *rng* keyword is used with the *gaussian* value, then the noise is generated from a gaussian distribution. Typically this added complexity is unnecessary, and one should be fine using the *uniform* value for reasons argued in (*Dunweg*).

If the *rng* keyword is used with the *none* value, then the noise terms are set to zero.

The *gamma\_t* keyword sets the (isotropic) translational viscous damping. Required for (and only compatible with) *brownian* and *brownian/sphere*. The units of *gamma\_t* are mass/time.

The *gamma\_r* keyword sets the (isotropic) rotational viscous damping. Required for (and only compatible with) *brownian/sphere*. The units of *gamma\_r* are mass\*length\*\*2/time.

The *gamma\_r\_eigen*, and *gamma\_t\_eigen* keywords are the eigenvalues of the rotational and viscous damping tensors (having the same units as their isotropic counterparts). Required for (and only compatible with) *brownian/asphere*. For a 2D system, the first two values of *gamma\_r\_eigen* must be inf (only rotation in xy plane), and the third value of *gamma\_t\_eigen* must be inf (only diffusion in xy plane).

If the *dipole* keyword is used, then the dipole moments of the particles are updated as described above. Only compatible with *brownian/asphere* (as *brownian/sphere* updates dipoles automatically).

If the *rotation\_temp* keyword is used, then the rotational diffusion will be occur at this prescribed temperature instead of *temp*. Only compatible with *brownian/sphere* and *brownian/asphere*.

If the *planar\_rotation* keyword is used, then rotation is constrained to the xy plane in a 3D simulation. Only compatible with *brownian/sphere* and *brownian/asphere* in 3D.

---

---

**Note:** For style *brownian/asphere*, the components *gamma\_t\_eigen* =(x,x,x) and *gamma\_r\_eigen* = (y,y,y), the dynamics will replicate those of the *brownian/sphere* style with *gamma\_t* = x and *gamma\_r* = y.

---

### 17.30.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. No global or per-atom quantities are stored by this fix for access by various *output commands*.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.30.5 Restrictions

The style *brownian/sphere* fix requires that atoms store torque and angular velocity (*omega*) as defined by the *atom\_style sphere* command. The style *brownian/asphere* fix requires that atoms store torque and quaternions as defined by the *atom\_style ellipsoid* command. If the *dipole* keyword is used, they must also store a dipole moment as defined by the *atom\_style dipole* command.

This fix is part of the BROWNIAN package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

### 17.30.6 Related commands

*fix propel/self*, *fix langevin*, *fix nve/sphere*,

### 17.30.7 Default

The default for *rng* is *uniform*. The default for the rotational and translational friction tensors are the identity tensor.

---

**(Ilie)** Ilie, Briels, den Otter, Journal of Chemical Physics, 142, 114103 (2015).

**(Delong)** Delong, Usabiaga, Donev, Journal of Chemical Physics. 143, 144107 (2015)

**(Dunweg)** Dunweg and Paul, Int J of Modern Physics C, 2, 817-27 (1991).

## 17.31 fix charge/regulation command

### 17.31.1 Syntax

```
fix ID group-ID charge/regulation cation_type anion_type keyword value(s)
```

- ID, group-ID are documented in fix command
- charge/regulation = style name of this fix command
- cation\_type = atom type of free cations
- anion\_type = atom type of free anions
- zero or more keyword/value pairs may be appended

keyword = *pH*, *pKa*, *pKb*, *pIp*, *pIm*, *pKs*, *acid\_type*, *base\_type*, *lunit\_nm*, *temp*, ↵

↵ *tempfixid*, *nevery*, *nmc*, *rxd*, *seed*, *tag*, *group*, *onllysalt*, *pmcmoves*

*pH* value = pH of the solution (can be specified as an equal-style variable)

*pKa* value = acid dissociation constant (in the -log<sub>10</sub> representation)

`pKb` value = base dissociation constant (in the  $-\log_{10}$  representation)  
`pIp` value = activity (effective concentration) of free cations (in the  $-\log_{10}$  representation)  
`pIm` value = activity (effective concentration) of free anions (in the  $-\log_{10}$  representation)  
`pKs` value = solvent self-dissociation constant (in the  $-\log_{10}$  representation)  
`acid_type` = atom type of acid groups  
`base_type` = atom type of base groups  
`lunit_nm` value = unit length used by LAMMPS (# in the units of nanometers)  
`temp` value = temperature  
`tempfixid` value = fix ID of temperature thermostat  
`nevery` value = invoke this fix every nevery steps  
`nmc` value = number of charge regulation MC moves to attempt every nevery steps  
`rxd` value = cutoff distance for acid/base reaction  
`seed` value = random # seed (positive integer)  
`tag` value = yes or no (yes: The code assign unique tags to inserted ions; no: The tag of all inserted ions is "0")  
`group` value = group-ID, inserted ions are assigned to group group-ID. Can be used multiple times to assign inserted ions to multiple groups.  
`onltsalt` values = flag `charge_cation` `charge_anion`.  
    flag = yes or no (yes: the fix performs only ion insertion/deletion, no: perform acid/base dissociation and ion insertion/deletion)  
    `charge_cation`, `charge_anion` = value of cation/anion charge, must be an integer (only specify if flag = yes)  
`pmcmoves` values = `pmcA` `pmcB` `pmcI` - MC move fractions for acid ionization (`pmcA`), base ionization (`pmcB`) and free ion exchange (`pmcI`)

## 17.31.2 Examples

```

fix chareg all charge/regulation 1 2 acid_type 3 base_type 4 pKa 5.0 pKb 6.0 pH 7.0 pIp
↪3.0 pIm 3.0 nevery 200 nmc 200 seed 123 tempfixid fT

fix chareg all charge/regulation 1 2 pIp 3 pIm 3 onltsalt yes 2 -1 seed 123 tag yes temp
↪1.0

```

## 17.31.3 Description

This fix performs Monte Carlo (MC) sampling of charge regulation and exchange of ions with a reservoir as discussed in (*Curk1*) and (*Curk2*). The implemented method is largely analogous to the grand-reaction ensemble method in (*Landsgesell*). The implementation is parallelized, compatible with existing LAMMPS functionalities, and applicable to any system utilizing discrete, ionizable groups or surface sites.

Specifically, the fix implements the following three types of MC moves, which discretely change the charge state of individual particles and insert ions into the systems:  $A \rightleftharpoons A^- + X^+$ ,  $B \rightleftharpoons B^+ + X^-$ , and  $\emptyset \rightleftharpoons Z^- X^{Z^+} + Z^+ X^{-Z^-}$ . In the former two types of reactions, Monte Carlo moves alter the charge value of specific atoms (A, B) and simultaneously insert a counterion to preserve the charge neutrality of the system, modeling the dissociation/association process. The last type of reaction performs grand canonical MC exchange of ion pairs with a (fictitious) reservoir.

In our implementation “acid” refers to particles that can attain charge  $q = \{0, -1\}$  and “base” to particles with  $q = \{0, 1\}$ , whereas the MC exchange of free ions allows any integer charge values of  $Z^+$  and  $Z^-$ .

Here we provide several practical examples for modeling charge regulation effects in solvated systems. An acid ionization reaction ( $A \rightleftharpoons A^- + H^+$ ) can be defined via a single line in the input file

```
fix acid_reaction all charge/regulation 2 3 acid_type 1 pH 7.0 pKa 5.0 pIp 7.0 pIm 7.0
```

where the fix attempts to charge A (discharge  $A^-$ ) to  $A^-$  (A) and insert (delete) a proton (atom type 2). Besides, the fix implements self-ionization reaction of water  $\emptyset \rightleftharpoons H^+ + OH^-$ .

However, this approach is highly inefficient at  $pH \approx 7$  when the concentration of both protons and hydroxyl ions is low, resulting in a relatively low acceptance rate of MC moves.

A more efficient way is to allow salt ions to participate in ionization reactions, which can be easily achieved via

```
fix acid_reaction2 all charge/regulation 4 5 acid_type 1 pH 7.0 pKa 5.0 pIp 2.0 pIm 2.0
```

where particles of atom type 4 and 5 are the salt cations and anions, both at activity (effective concentration) of  $10^{-2}$  mol/l, see (*Curkl*) and (*Landsgesell*) for more details.

We could have simultaneously added a base ionization reaction ( $B \rightleftharpoons B^+ + OH^-$ )

```
fix acid_base_reaction all charge/regulation 2 3 acid_type 1 base_type 6 pH 7.0 pKa 5.0 ↵
↵pKb 6.0 pIp 7.0 pIm 7.0
```

where the fix will attempt to charge B (discharge  $B^+$ ) to  $B^+$  (B) and insert (delete) a hydroxyl ion  $OH^-$  of atom type 3.

Dissociated ions and salt ions can be combined into a single particle type, which reduces the number of necessary MC moves and increases sampling performance, see (*Curkl*). The  $H^+$  and monovalent salt cation ( $S^+$ ) are combined into a single particle type,  $X^+ = \{H^+, S^+\}$ . In this case “pIp” refers to the effective concentration of the combined cation type  $X^+$  and its value is determined by  $10^{-pIp} = 10^{-pH} + 10^{-pSp}$ , where  $10^{-pSp}$  is the effective concentration of salt cations. For example, at  $pH=7$  and  $pSp=6$  we would find  $pIp \sim 5.958$  and the command that performs reactions with combined ions could read,

```
fix acid_reaction_combined all charge/regulation 2 3 acid_type 1 pH 7.0 pKa 5.0 pIp 5.
↵958 pIm 5.958
```

If neither the acid or the base type is specified, for example,

```
fix salt_reaction all charge/regulation 4 5 pIp 2.0 pIm 2.0
```

the fix simply inserts or deletes an ion pair of a free cation (atom type 4) and a free anion (atom type 5) as done in a conventional grand-canonical MC simulation. Multivalent ions can be inserted (deleted) by using the *onlysalt* keyword.

The fix is compatible with LAMMPS sub-packages such as *molecule* or *rigid*. The acid and base particles can be part of larger molecules or rigid bodies. Free ions that are inserted to or deleted from the system must be defined as single particles (no bonded interactions allowed) and cannot be part of larger molecules or rigid bodies. If *molecule* package is used, all inserted ions have a molecule ID equal to zero.

Note that LAMMPS implicitly assumes a constant number of particles (degrees of freedom). Since using this fix alters the total number of particles during the simulation, any thermostat used by LAMMPS, such as NVT or Langevin, must use a dynamic calculation of system temperature. This can be achieved by specifying a dynamic temperature compute (e.g. *dtemp*) and using it with the desired thermostat, e.g. a Langevin thermostat:

```
compute dtemp all temp
compute_modify dtemp dynamic yes
fix fT all langevin 1.0 1.0 1.0 123
fix_modify fT temp dtemp
```

The units of pH, pKa, pKb, pIp, pIm are considered to be in the standard  $-\log_{10}$  representation assuming reference concentration  $\rho_0 = \text{mol/l}$ . For example, in the dilute ideal solution limit, the concentration of free cations will be

$c_T = 10^{-pI_p} \text{mol/l}$ . To perform the internal unit conversion, the value of the LAMMPS unit length must be specified in nanometers via *lunit\_nm*. The default value is set to the Bjerrum length in water at room temperature (0.71 nm), *lunit\_nm* = 0.71.

The temperature used in MC acceptance probability is set by *temp*. This temperature should be the same as the temperature set by the molecular dynamics thermostat. For most purposes, it is probably best to use *tempfixid* keyword which dynamically sets the temperature equal to the chosen MD thermostat temperature, in the example above we assumed the thermostat fix-ID is *fT*. The inserted particles attain a random velocity corresponding to the specified temperature. Using *tempfixid* overrides any fixed temperature set by *temp*.

The *rxd* keyword can be used to restrict the inserted/deleted counterions to a specific radial distance from an acid or base particle that is currently participating in a reaction. This can be used to simulate more realistic reaction dynamics. If *rxd* = 0 or *rxd* >  $L / 2$ , where  $L$  is the smallest box dimension, the radial restriction is automatically turned off and free ion can be inserted or deleted anywhere in the simulation box.

If the *tag yes* is used, every inserted atom gets a unique tag ID, otherwise, the tag of every inserted atom is set to 0. *tag yes* might cause an integer overflow in very long simulations since the tags are unique to every particle and thus increase with every successful particle insertion.

The *pmcmoves* keyword sets the relative probability of attempting the three types of MC moves (reactions): acid charging, base charging, and ion pair exchange. The fix only attempts to perform particle charging MC moves if *acid\_type* or *base\_type* is defined. Otherwise fix only performs free ion insertion/deletion. For example, if *acid\_type* is not defined, *pmcA* is automatically set to 0. The vector *pmcmoves* is automatically normalized, for example, if set to *pmcmoves* 0 0.33 0.33, the vector would be normalized to [0,0.5,0.5].

The *only\_salt* option can be used to perform multivalent grand-canonical ion-exchange moves. If *only\_salt yes* is used, no charge exchange is performed, only ion insertion/deletion (*pmcmoves* is set to [0,0,1]), but ions can be multivalent. In the example above, an MC move would consist of three ion insertion/deletion to preserve the charge neutrality of the system.

The *group* keyword can be used to add inserted particles to a specific group-ID. All inserted particles are automatically added to group *all*.

## 17.31.4 Output

This fix computes a global vector of length 8, which can be accessed by various output commands. The vector values are the following global quantities:

- 1 = cumulative MC attempts
- 2 = cumulative MC successes
- 3 = current # of neutral acid atoms
- 4 = current # of -1 charged acid atoms
- 5 = current # of neutral base atoms
- 6 = current # of +1 charged base atoms
- 7 = current # of free cations
- 8 = current # of free anions

### 17.31.5 Restrictions

This fix is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The *atom\_style*, used must contain the charge property, for example, the style could be *charge* or *full*. Only usable for 3D simulations. Atoms specified as free ions cannot be part of rigid bodies or molecules and cannot have bonding interactions. The scheme is limited to integer charges, any atoms with non-integer charges will not be considered by the fix.

All interaction potentials used must be continuous, otherwise the MD integration and the particle exchange MC moves do not correspond to the same equilibrium ensemble. For example, if an *lj/cut* pair style is used, the LJ potential must be shifted so that it vanishes at the cutoff. This can be easily achieved using the *pair\_modify* command, i.e., by using: *pair\_modify shift yes*.

---

**Note:** Region restrictions are not yet implemented.

---

### 17.31.6 Related commands

*fix gcmc*, *fix atom/swap*

### 17.31.7 Default

pH = 7.0; pKa = 100.0; pKb = 100.0; pIp = 5.0; pIm = 5.0; pKs = 14.0; acid\_type = -1; base\_type = -1; lunit\_nm = 0.71; temp = 1.0; nevery = 100; nmc = 100; rxd = 0; seed = 0; tag = no; onlysalt = no; pmcmoves = [1/3, 1/3, 1/3], group-ID = all

---

**(Curk1)** T. Curk, J. Yuan, and E. Luijten, “Accelerated simulation method for charge regulation effects”, JCP 156 (2022).

**(Curk2)** T. Curk and E. Luijten, “Charge-regulation effects in nanoparticle self-assembly”, PRL 126 (2021)

**(Landsgesell)** J. Landsgesell, P. Hebbeker, O. Rud, R. Lunkad, P. Kosovan, and C. Holm, “Grand-reaction method for simulations of ionization equilibria coupled to ion partitioning”, Macromolecules 53, 3007-3020 (2020).

## 17.32 fix cmap command

### 17.32.1 Syntax

```
fix ID group-ID cmap filename
```

- ID, group-ID are documented in *fix* command
- cmap = style name of this fix command
- filename = force-field file with CMAP coefficients

## 17.32.2 Examples

```
fix          myCMAP all cmap ../potentials/cmap36.data
read_data    proteinX.data fix myCMAP crossterm CMAP
fix_modify   myCMAP energy yes
```

## 17.32.3 Description

This command enables CMAP 5-body interactions to be added to simulations which use the CHARMM force field. These are relevant for any CHARMM model of a peptide or protein sequences that is 3 or more amino-acid residues long; see (*Buck*) and (*Brooks*) for details, including the analytic energy expressions for CMAP interactions. The CMAP 5-body terms add additional potential energy contributions to pairs of overlapping phi-psi dihedrals of amino-acids, which are important to properly represent their conformational behavior.

The examples/cmap directory has a sample input script and data file for a small peptide, that illustrates use of the fix cmap command.

As in the example above, this fix should be used before reading a data file that contains a listing of CMAP interactions. The *filename* specified should contain the CMAP parameters for a particular version of the CHARMM force field. Two such files are including in the lammmps/potentials directory: charmm22.cmap and charmm36.cmap.

The data file read by the “read\_data” must contain the topology of all the CMAP interactions, similar to the topology data for bonds, angles, dihedrals, etc. Specially it should have a line like this in its header section:

```
N crossterms
```

where N is the number of CMAP 5-body interactions. It should also have a section in the body of the data file like this with N lines:

```
CMAP
      1      1      8      10      12      18      20
      2      5      18      20      22      25      27
      [...]
      N      3     314     315     317     318     330
```

The first column is an index from 1 to N to enumerate the CMAP 5-atom tuples; it is ignored by LAMMPS. The second column is the “type” of the interaction; it is an index into the CMAP force field file. The remaining 5 columns are the atom IDs of the atoms in the two 4-atom dihedrals that overlap to create the CMAP interaction. Note that the “crossterm” and “CMAP” keywords for the header and body sections match those specified in the read\_data command following the data file name; see the *read\_data* page for more details.

A data file containing CMAP 5-body interactions can be generated from a PDB file using the charmm2lammmps.pl script in the tools/ch2lmp directory of the LAMMPS distribution. The script must be invoked with the optional “-cmap” flag to do this; see the tools/ch2lmp/README file for more information. The same conversion script also creates the file of CMAP coefficient data which is read by this command.

The potential energy associated with CMAP interactions can be output as described below. It can also be included in the total potential energy of the system, as output by the *thermo\_style* command, if the *fix\_modify energy* command is used, as in the example above. See the note below about how to include the CMAP energy when performing an *energy minimization*.

### 17.32.4 Restart, `fix_modify`, `output`, `run start/stop`, `minimize info`

This fix writes the list of CMAP cross-terms to *binary restart files*. See the `read_restart` command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The `fix_modify energy` option is supported by this fix to add the potential energy of the CMAP interactions to both the global potential energy and peratom potential energies of the system as part of *thermodynamic output* or output by the `compute pe/atom` command. The default setting for this fix is `fix_modify energy yes`.

The `fix_modify virial` option is supported by this fix to add the contribution due to the CMAP interactions to both the global pressure and per-atom stress of the system via the `compute pressure` and `compute stress/atom` commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is `fix_modify virial yes`.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the potential energy discussed above. The scalar value calculated by this fix is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the `run` command.

The forces due to this fix are imposed during an energy minimization, invoked by the `minimize` command.

The `fix_modify respa` option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

---

**Note:** For energy minimization, if you want the potential energy associated with the CMAP terms forces to be included in the total potential energy of the system (the quantity being minimized), you **MUST** not disable the `fix_modify energy` option for this fix.

---

### 17.32.5 Restrictions

To function as expected this fix command must be issued *before* a `read_data` command but *after* a `read_restart` command.

This fix can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* page for more info.

### 17.32.6 Related commands

`fix_modify`, `read_data`

### 17.32.7 Default

none

---

**(Buck)** Buck, Bouguet-Bonnet, Pastor, MacKerell Jr., *Biophys J*, 90, L36 (2006).

**(Brooks)** Brooks, Brooks, MacKerell Jr., *J Comput Chem*, 30, 1545 (2009).



## 17.33 fix colvars command

### 17.33.1 Syntax

```
fix ID group-ID colvars configfile keyword values ...
```

- ID, group-ID are documented in *fix* command
- colvars = style name of this fix command
- configfile = the configuration file for the colvars module
- keyword = *input* or *output* or *seed* or *tstat*

*input* arg = colvars.state file name or prefix or NULL (default: NULL)

*output* arg = output filename prefix (default: out)

*seed* arg = seed for random number generator (default: 1966)

*unwrap* arg = *yes* or *no*

use unwrapped coordinates in collective variables (default: yes)

*tstat* arg = fix id of a thermostat or NULL (default: NULL)

### 17.33.2 Examples

```
fix mtd all colvars peptide.colvars.inp seed 2122 input peptide.colvars.state output_
↳peptide
fix abf all colvars colvars.inp tstat 1
```

### 17.33.3 Description

This fix interfaces LAMMPS to the collective variables (Colvars) library, which allows to calculate potentials of mean force (PMFs) for any set of colvars, using different sampling methods: currently implemented are the Adaptive Biasing Force (ABF) method, metadynamics, Steered Molecular Dynamics (SMD) and Umbrella Sampling (US) via a flexible harmonic restraint bias.

This documentation describes only the fix colvars command itself and LAMMPS specific parts of the code. The full documentation of the colvars library is available as [this supplementary PDF document](#)

The Colvars library is developed at <https://github.com/colvars/colvars> A detailed discussion of its implementation is in (*Fiorin*).

There are some example scripts for using this package with LAMMPS in the examples/PACKAGES/colvars directory.

The only mandatory argument to the fix is the filename to the colvars input file that contains the input that is independent from the MD program in which the colvars library has been integrated.

The *group-ID* entry is ignored. The collective variable module will always apply to the entire system and there can only be one instance of the colvars fix at a time. The colvars fix will only communicate the minimum information necessary and the colvars library supports multiple, completely independent collective variables, so there is no restriction to functionality by limiting the number of colvars fixes.

The *input* keyword allows to specify a state file that would contain the restart information required in order to continue a calculation from a prerecorded state. Fix colvars records its state in *binary restart* files, so when using the *read\_restart* command, this is usually not needed.

The *output* keyword allows to specify the output prefix. All output files generated will use this prefix followed by the “.colvars.” and a word like “state” or “traj”.

The *seed* keyword contains the seed for the random number generator that will be used in the colvars module.

The *unwrap* keyword controls whether wrapped or unwrapped coordinates are passed to the colvars library for calculation of the collective variables and the resulting forces. The default is *yes*, i.e. to use the image flags to reconstruct the absolute atom positions. Setting this to *no* will use the current local coordinates that are wrapped back into the simulation cell at each re-neighboring instead.

The *tstat* keyword can be either NULL or the label of a thermostating fix that thermostats all atoms in the fix colvars group. This will be used to provide the colvars module with the current thermostat target temperature.

### 17.33.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the current status of the colvars module into *binary restart files*. This is in addition to the text mode status file that is written by the colvars module itself and the kind of information in both files is identical.

The *fix\_modify energy* option is supported by this fix to add the energy change from the biasing force added by Colvars to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify configfile <config file>* option allows to add settings from an additional config file to the colvars module. This option can only be used, after the system has been initialized with a *run* command.

The *fix\_modify config <quoted string>* option allows to add settings from inline strings. Those have to fit on a single line when enclosed in a pair of double quotes (“”), or can span multiple lines when bracketed by a pair of triple double quotes (“””), like python embedded documentation).

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the Colvars energy mentioned above. The scalar value calculated by this fix is “extensive”.

### 17.33.5 Restrictions

This fix is part of the COLVARS package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

There can only be one colvars fix active at a time. Since the interface communicates only the minimum amount of information and colvars module itself can handle an arbitrary number of collective variables, this is not a limitation of functionality.

### 17.33.6 Related commands

*fix smd, fix spring, fix plumed*

### 17.33.7 Default

The default options are input = NULL, output = out, seed = 1966, unwrap yes, and tstat = NULL.

(Fiorin) Fiorin, Klein, Henin, Mol. Phys., DOI:10.1080/00268976.2013.813594

## 17.34 fix controller command

### 17.34.1 Syntax

```
fix ID group-ID controller Nevery alpha Kp Ki Kd pvar setpoint cvar
```

- ID, group-ID are documented in *fix* command
- controller = style name of this fix command
- Nevery = invoke controller every this many timesteps
- alpha = coupling constant for PID equation (see units discussion below)
- Kp = proportional gain in PID equation (unitless)
- Ki = integral gain in PID equation (unitless)
- Kd = derivative gain in PID equation (unitless)
- pvar = process variable of form c\_ID, c\_ID[I], f\_ID, f\_ID[I], or v\_name

```
c_ID = global scalar calculated by a compute with ID
c_ID[I] = Ith component of global vector calculated by a compute with ID
f_ID = global scalar calculated by a fix with ID
f_ID[I] = Ith component of global vector calculated by a fix with ID
v_name = value calculated by an equal-style variable with name
```

- setpoint = desired value of process variable (same units as process variable)
- cvar = name of control variable

### 17.34.2 Examples

```
fix 1 all controller 100 1.0 0.5 0.0 0.0 c_thermo_temp 1.5 tcontrol
fix 1 all controller 100 0.2 0.5 0 100.0 v_pxxwall 1.01325 xwall
fix 1 all controller 10000 0.2 0.5 0 2000 v_avpe -3.785 tcontrol
```

### 17.34.3 Description

This fix enables control of a LAMMPS simulation using a control loop feedback mechanism known as a proportional-integral-derivative (PID) controller. The basic idea is to define a “process variable” which is a quantity that can be monitored during a running simulation. A desired target value is chosen for the process variable. A “control variable” is also defined which is an adjustable attribute of the running simulation, which the process variable will respond to. The PID controller continuously adjusts the control variable based on the difference between the process variable and the target.

Here are examples of ways in which this fix can be used. The `examples/pid` directory contains a script that implements the simple thermostat.

Goal	process variable	control variable
Simple thermostat	instantaneous T	thermostat target T
Find melting temperature	average PE per atom	thermostat target T
Control pressure in non-periodic system	force on wall	position of wall

---

**Note:** For this fix to work, the control variable must actually induce a change in a running LAMMPS simulation. Typically this will only occur if there is some other command (e.g. a thermostat fix) which uses the control variable as an input parameter. This could be done directly or indirectly, e.g. the other command uses a variable as input whose formula uses the control variable. The other command should alter its behavior dynamically as the variable changes.

---



---

**Note:** If there is a command you think could be used in this fashion, but does not currently allow a variable as an input parameter, please notify the LAMMPS developers. It is often not difficult to enable a command to use a variable as an input parameter.

---

The group specified with this command is ignored. However, note that the process variable may be defined by calculations performed by computes and fixes which store their own “group” definitions.

The PID controller is invoked once each *Nevery* timesteps.

The PID controller is implemented as a discretized version of the following dynamic equation:

$$\frac{dc}{dt} = \hat{E} - \alpha \left( K_p e + K_i \int_0^t e dt + K_d \frac{de}{dt} \right)$$

where  $c$  is the continuous time analog of the control variable,  $e = pvar - setpoint$  is the error in the process variable, and  $\alpha$ ,  $K_p$ ,  $K_i$ , and  $K_d$  are constants set by the corresponding keywords described above. The discretized version of this equation is:

$$c_n = \hat{E}c_{n-1} - \alpha \left( K_p \tau e_n + K_i \tau^2 \sum_{i=1}^n e_i + K_d (e_n - e_{n-1}) \right)$$

where  $\tau = Nevery \cdot timestep$  is the time interval between updates, and the subscripted variables indicate the values of  $c$  and  $e$  at successive updates.

From the first equation, it is clear that if the three gain values  $K_p$ ,  $K_i$ ,  $K_d$  are dimensionless constants, then  $\alpha$  must have units of  $[unit\ cvar]/[unit\ pvar]/[unit\ time]$  e.g.  $[eV/K/ps]$ . The advantage of this unit scheme is that the value of the constants should be invariant under a change of either the MD timestep size or the value of *Nevery*. Similarly, if the LAMMPS *unit style* is changed, it should only be necessary to change the value of  $\alpha$  to reflect this, while leaving  $K_p$ ,  $K_i$ , and  $K_d$  unaltered.

When choosing the values of the four constants, it is best to first pick a value and sign for  $\alpha$  that is consistent with the magnitudes and signs of  $pvar$  and  $cvar$ . The magnitude of  $K_p$  should then be tested over a large positive range keeping  $K_i = K_d = 0$ . A good value for  $K_p$  will produce a fast response in  $pvar$ , without overshooting the *setpoint*. For many applications, proportional feedback is sufficient, and so  $K_i = K_d = 0$  can be used. In cases where there is a substantial lag time in the response of  $pvar$  to a change in  $cvar$ , this can be counteracted by increasing  $K_d$ . In situations where  $pvar$  plateaus without reaching *setpoint*, this can be counteracted by increasing  $K_i$ . In the language of Charles Dickens,  $K_p$  represents the error of the present,  $K_i$  the error of the past, and  $K_d$  the error yet to come.

Because this fix updates  $cvar$ , but does not initialize its value, the initial value is that assigned by the user in the input script via the *internal-style variable* command. This value is used (by the other LAMMPS command that used the variable) until this fix performs its first update of  $cvar$  after *Nevery* timesteps. On the first update, the value of the derivative term is set to zero, because the value of  $e_n - 1$  is not yet defined.

The process variable  $pvar$  can be specified as the output of a *compute* or *fix* or the evaluation of a *variable*. In each case, the compute, fix, or variable must produce a global quantity, not a per-atom or local quantity.

If  $pvar$  begins with “c\_”, a compute ID must follow which has been previously defined in the input script and which generates a global scalar or vector. See the individual *compute* doc page for details. If no bracketed integer is appended, the scalar calculated by the compute is used. If a bracketed integer is appended, the *I*th value of the vector calculated by the compute is used. Users can also write code for their own compute styles and *add them to LAMMPS*.

If  $pvar$  begins with “f\_”, a fix ID must follow which has been previously defined in the input script and which generates a global scalar or vector. See the individual *fix* page for details. Note that some fixes only produce their values on certain timesteps, which must be compatible with when fix controller references the values, or else an error results. If no bracketed integer is appended, the scalar calculated by the fix is used. If a bracketed integer is appended, the *I*th value of the vector calculated by the fix is used. Users can also write code for their own fix style and *add them to LAMMPS*.

If  $pvar$  begins with “v\_”, a variable name must follow which has been previously defined in the input script. Only equal-style variables can be referenced. See the *variable* command for details. Note that variables of style *equal* define a formula which can reference individual atom properties or thermodynamic keywords, or they can invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of specifying the process variable.

The target value *setpoint* for the process variable must be a numeric value, in whatever units  $pvar$  is defined for.

The control variable  $cvar$  must be the name of an *internal-style variable* previously defined in the input script. Note that it is not specified with a “v\_” prefix, just the name of the variable. It must be an internal-style variable, because this fix updates its value directly. Note that other commands can use an equal-style versus internal-style variable interchangeably.

#### 17.34.4 Restart, fix\_modify, output, run start/stop, minimize info

Currently, no information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix produces a global vector with 3 values which can be accessed by various *output commands*. The values can be accessed on any timestep, though they are only updated on timesteps that are a multiple of *Nevery*.

The three values are the most recent updates made to the control variable by each of the 3 terms in the PID equation above. The first value is the proportional term, the second is the integral term, the third is the derivative term.

The units of the vector values will be whatever units the control variable is in. The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.34.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.34.6 Related commands

*fix adapt*

### 17.34.7 Default

none

## 17.35 fix damping/cundall command

### 17.35.1 Syntax

```
fix ID group-ID damping/cundall gamma_l gamma_a keyword values ...
```

- ID, group-ID are documented in *fix* command
- damping/cundall = style name of this fix command
- gamma\_l = linear damping coefficient (dimensionless)
- gamma\_a = angular damping coefficient (dimensionless)
- zero or more keyword/value pairs may be appended

keyword = *scale*

scale values = *type ratio* or *v\_name*

type = atom type (1-N)

ratio = factor to scale the damping coefficients by

v\_name = reference to atom style variable *name*

### 17.35.2 Examples

```
fix 1 all damping/cundall 0.8 0.8  
fix 1 all damping/cundall 0.8 0.5 scale 3 2.5  
fix a all damping/cundall 0.8 0.5 scale v_radscale
```

### 17.35.3 Description

Add damping force and torque to finite-size spherical particles in the group following the model of *Cundall, 1987*, as implemented in other granular physics code (e.g., *Yade-DEM*, *PFC*).

The damping is constructed to always have negative mechanical power with respect to the current velocity/angular velocity to ensure dissipation of kinetic energy. If used without additional thermostating (to add kinetic energy to the system), it has the effect of slowly (or rapidly) freezing the system; hence it can also be used as a simple energy minimization technique.

The magnitude of the damping force/torque  $F_d/T_d$  is a fraction  $\gamma \in [0; 1]$  of the current force/torque  $F/T$  on the particle. Damping is applied component-by-component in each direction  $k \in \{x, y, z\}$ :

$$F_{dk} = -\gamma_l F_k \text{sign}(F_k v_k)$$

$$T_{dk} = -\gamma_a T_k \text{sign}(T_k \omega_k)$$

The larger the coefficients, the faster the kinetic energy is reduced.

If the optional keyword *scale* is used,  $\gamma_l$  and  $\gamma_a$  can be scaled up or down by the specified factor for atoms. This factor can be set for different atom types and thus the *scale* keyword used multiple times followed by the atom type and the associated scale factor. Alternately the scaling factor can be computed for each atom (e.g. based on its radius) by using an *atom-style variable*.

---

**Note:** The damping force/torque is computed based on the force/torque at the moment this fix is invoked. Any force/torque added after this fix, e.g., by *fix addforce* or *fix addtorque* will not be damped. When performing simulations with gravity, invoking *fix gravity* after this fix will maintain the specified gravitational acceleration.

---



---

**Note:** This scheme is dependent on the coordinates system and does not correspond to realistic physical processes. It is constructed for numerical convenience and efficacy.

---

This non-viscous damping presents the following advantages:

1. damping is independent of velocity, equally damping regions with distinct natural frequencies,
  2. damping affects acceleration and vanishes for steady uniform motion of the particles,
  3. damping parameter  $\gamma$  is dimensionless and does not require scaling.
- 

### 17.35.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is modifying forces/torques. Default is the outermost level.

The forces/torques due to this fix are imposed during an energy minimization, invoked by the *minimize* command. This fix should only be used with damped dynamics minimizers that allow for non-conservative forces. See the *min\_style* command for details.

### 17.35.5 Restrictions

This fix is part of the GRANULAR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and a radius as defined by the *atom\_style sphere* command.

### 17.35.6 Related commands

*fix viscous*, *fix viscous/sphere*

### 17.35.7 Default

none

### 17.35.8 References

**(Cundall, 1987)** Cundall, P. A. Distinct Element Models of Rock and Soil Structure, in Analytical and Computational Methods in Engineering Rock Mechanics, Ch. 4, pp. 129-163. E. T. Brown, ed. London: Allen & Unwin., 1987.

**(PFC)** PFC Particle Flow Code 6.0 Documentation. Itasca Consulting Group.

**(Yade-DEM)** V. Smilauer et al. (2021), Yade Documentation 3rd ed. The Yade Project. DOI:10.5281/zenodo.5705394 (<https://yade-dem.org/doc/>)

## 17.36 fix deform command

Accelerator Variants: *deform/kk*

### 17.36.1 Syntax

```
fix ID group-ID deform N parameter args ... keyword value ...
```

- ID, group-ID are documented in *fix* command
- deform = style name of this fix command
- N = perform box deformation every this many timesteps
- one or more parameter/arg pairs may be appended

parameter = x or y or z or xy or xz or yz

x, y, z args = style value(s)

style = *final* or *delta* or *scale* or *vel* or *erate* or *trate* or *volume* or *wiggle* or *variable*

→variable

*final* values = lo hi

lo hi = box boundaries at end of run (distance units)

*delta* values = dlo dhi

dlo dhi = change in box boundaries at end of run (distance units)

*scale* values = factor

factor = multiplicative factor for change in box length at end of run

*vel* value = V



V = change box length at this velocity (distance/time units),  
 effectively an engineering strain rate  
*erate* value = R  
 R = engineering strain rate (1/time units)  
*trate* value = R  
 R = true strain rate (1/time units)  
*volume* value = none = adjust this dim to preserve volume of system  
*wiggle* values = A Tp  
 A = amplitude of oscillation (distance units)  
 Tp = period of oscillation (time units)  
*variable* values = v\_name1 v\_name2  
 v\_name1 = variable with name1 for box length change as function of time  
 v\_name2 = variable with name2 for change rate as function of time  
*xy*, *xz*, *yz* args = style value  
*style* = *final* or *delta* or *vel* or *erate* or *trate* or *wiggle*  
*final* value = tilt  
 tilt = tilt factor at end of run (distance units)  
*delta* value = dtilt  
 dtilt = change in tilt factor at end of run (distance units)  
*vel* value = V  
 V = change tilt factor at this velocity (distance/time units),  
 effectively an engineering shear strain rate  
*erate* value = R  
 R = engineering shear strain rate (1/time units)  
*trate* value = R  
 R = true shear strain rate (1/time units)  
*wiggle* values = A Tp  
 A = amplitude of oscillation (distance units)  
 Tp = period of oscillation (time units)  
*variable* values = v\_name1 v\_name2  
 v\_name1 = variable with name1 for tilt change as function of time  
 v\_name2 = variable with name2 for change rate as function of time

- zero or more keyword/value pairs may be appended
- keyword = *remap* or *flip* or *units*

*remap* value = x or v or none  
 x = remap coords of atoms in group into deforming box  
 v = remap velocities of atoms in group when they cross periodic boundaries  
 none = no remapping of x or v  
*flip* value = yes or no  
 allow or disallow box flips when it becomes highly skewed  
*units* value = *lattice* or *box*  
 lattice = distances are defined in lattice units  
 box = distances are defined in simulation box units

## 17.36.2 Examples

```
fix 1 all deform 1 x final 0.0 9.0 z final 0.0 5.0 units box
fix 1 all deform 1 x trate 0.1 y volume z volume
fix 1 all deform 1 xy erate 0.001 remap v
fix 1 all deform 10 y delta -0.5 0.5 xz vel 1.0
```

## 17.36.3 Description

Change the volume and/or shape of the simulation box during a dynamics run. Orthogonal simulation boxes have 3 adjustable parameters ( $x,y,z$ ). Triclinic (non-orthogonal) simulation boxes have 6 adjustable parameters ( $x,y,z,xy,xz,yz$ ). Any or all of them can be adjusted independently and simultaneously by this command.

This fix can be used to perform non-equilibrium MD (NEMD) simulations of a continuously strained system. See the *fix nvt/sllod* and *compute temp/deform* commands for more details. Note that simulation of a continuously extended system (extensional flow) can be modeled using the *UEF package* and its *fix commands*.

For the  $x$ ,  $y$ ,  $z$  parameters, the associated dimension cannot be shrink-wrapped. For the  $xy$ ,  $yz$ ,  $xz$  parameters, the associated second dimension cannot be shrink-wrapped. Dimensions not varied by this command can be periodic or non-periodic. Dimensions corresponding to unspecified parameters can also be controlled by a *fix npt* or *fix nph* command.

The size and shape of the simulation box at the beginning of the simulation run were either specified by the *create\_box* or *read\_data* or *read\_restart* command used to setup the simulation initially if it is the first run, or they are the values from the end of the previous run. The *create\_box*, *read\_data*, and *read\_restart* commands specify whether the simulation box is orthogonal or non-orthogonal (triclinic) and explain the meaning of the  $xy,xz,yz$  tilt factors. If fix deform changes the  $xy,xz,yz$  tilt factors, then the simulation box must be triclinic, even if its initial tilt factors are 0.0.

As described below, the desired simulation box size and shape at the end of the run are determined by the parameters of the fix deform command. Every Nth timestep during the run, the simulation box is expanded, contracted, or tilted to ramped values between the initial and final values.

---

For the  $x$ ,  $y$ , and  $z$  parameters, this is the meaning of their styles and values.

The *final*, *delta*, *scale*, *vel*, and *erate* styles all change the specified dimension of the box via “constant displacement” which is effectively a “constant engineering strain rate”. This means the box dimension changes linearly with time from its initial to final value.

For style *final*, the final lo and hi box boundaries of a dimension are specified. The values can be in lattice or box distance units. See the discussion of the units keyword below.

For style *delta*, plus or minus changes in the lo/hi box boundaries of a dimension are specified. The values can be in lattice or box distance units. See the discussion of the units keyword below.

For style *scale*, a multiplicative factor to apply to the box length of a dimension is specified. For example, if the initial box length is 10, and the factor is 1.1, then the final box length will be 11. A factor less than 1.0 means compression.

For style *vel*, a velocity at which the box length changes is specified in units of distance/time. This is effectively a “constant engineering strain rate”, where  $\text{rate} = V/L_0$  and  $L_0$  is the initial box length. The distance can be in lattice or box distance units. See the discussion of the units keyword below. For example, if the initial box length is 100 Angstroms, and  $V$  is 10 Angstroms/ps, then after 10 ps, the box length will have doubled. After 20 ps, it will have tripled.

The *erate* style changes a dimension of the box at a “constant engineering strain rate”. The units of the specified strain rate are 1/time. See the *units* command for the time units associated with different choices of simulation units, e.g. picoseconds for “metal” units). Tensile strain is unitless and is defined as  $\text{delta}/L_0$ , where  $L_0$  is the original box length and delta is the change relative to the original length. The box length  $L$  as a function of time will change as

$$L(t) = L_0 (1 + \text{erate} * dt)$$

where  $dt$  is the elapsed time (in time units). Thus if *erate*  $R$  is specified as 0.1 and time units are picoseconds, this means the box length will increase by 10% of its original length every picosecond. I.e. strain after 1 ps = 0.1, strain after 2 ps = 0.2, etc.  $R = -0.01$  means the box length will shrink by 1% of its original length every picosecond. Note that for an “engineering” rate the change is based on the original box length, so running with  $R = 1$  for 10 picoseconds expands the box length by a factor of 11 (strain of 10), which is different that what the *trate* style would induce.

The *trate* style changes a dimension of the box at a “constant true strain rate”. Note that this is not an “engineering strain rate”, as the other styles are. Rather, for a “true” rate, the rate of change is constant, which means the box dimension changes non-linearly with time from its initial to final value. The units of the specified strain rate are 1/time. See the *units* command for the time units associated with different choices of simulation units, e.g. picoseconds for “metal” units). Tensile strain is unitless and is defined as  $\Delta/L_0$ , where  $L_0$  is the original box length and  $\Delta$  is the change relative to the original length.

The box length  $L$  as a function of time will change as

$$L(t) = L_0 \exp(\text{trate} * dt)$$

where  $dt$  is the elapsed time (in time units). Thus if *trate*  $R$  is specified as  $\ln(1.1)$  and time units are picoseconds, this means the box length will increase by 10% of its current (not original) length every picosecond. I.e. strain after 1 ps = 0.1, strain after 2 ps = 0.21, etc.  $R = \ln(2)$  or  $\ln(3)$  means the box length will double or triple every picosecond.  $R = \ln(0.99)$  means the box length will shrink by 1% of its current length every picosecond. Note that for a “true” rate the change is continuous and based on the current length, so running with  $R = \ln(2)$  for 10 picoseconds does not expand the box length by a factor of 11 as it would with *erate*, but by a factor of 1024 since the box length will double every picosecond.

Note that to change the volume (or cross-sectional area) of the simulation box at a constant rate, you can change multiple dimensions via *erate* or *trate*. E.g. to double the box volume in a picosecond picosecond, you could set “x erate M”, “y erate M”, “z erate M”, with  $M = \text{pow}(2,1/3) - 1 = 0.26$ , since if each box dimension grows by 26%, the box volume doubles. Or you could set “x trate M”, “y trate M”, “z trate M”, with  $M = \ln(1.26) = 0.231$ , and the box volume would double every picosecond.

The *volume* style changes the specified dimension in such a way that the box volume remains constant while other box dimensions are changed explicitly via the styles discussed above. For example, “x scale 1.1 y scale 1.1 z volume” will shrink the z box length as the x,y box lengths increase, to keep the volume constant (product of x,y,z lengths). If “x scale 1.1 z volume” is specified and parameter y is unspecified, then the z box length will shrink as x increases to keep the product of x,z lengths constant. If “x scale 1.1 y volume z volume” is specified, then both the y,z box lengths will shrink as x increases to keep the volume constant (product of x,y,z lengths). In this case, the y,z box lengths shrink so as to keep their relative aspect ratio constant.

For solids or liquids, note that when one dimension of the box is expanded via *fix deform* (i.e. tensile strain), it may be physically undesirable to hold the other 2 box lengths constant (unspecified by *fix deform*) since that implies a density change. Using the *volume* style for those 2 dimensions to keep the box volume constant may make more physical sense, but may also not be correct for materials and potentials whose Poisson ratio is not 0.5. An alternative is to use *fix npt aniso* with zero applied pressure on those 2 dimensions, so that they respond to the tensile strain dynamically.

The *wiggle* style oscillates the specified box length dimension sinusoidally with the specified amplitude and period. I.e. the box length  $L$  as a function of time is given by

$$L(t) = L_0 + A \sin(2 * \pi t / T_p)$$

where  $L_0$  is its initial length. If the amplitude  $A$  is a positive number the box initially expands, then contracts, etc. If  $A$  is negative then the box initially contracts, then expands, etc. The amplitude can be in lattice or box distance units. See the discussion of the *units* keyword below.

The *variable* style changes the specified box length dimension by evaluating a variable, which presumably is a function of time. The variable with *name1* must be an *equal-style variable* and should calculate a change in box length in units of distance. Note that this distance is in box units, not lattice units; see the discussion of the *units* keyword below. The formula associated with variable *name1* can reference the current timestep. Note that it should return the “change” in

box length, not the absolute box length. This means it should evaluate to 0.0 when invoked on the initial timestep of the run following the definition of fix deform. It should evaluate to a value > 0.0 to dilate the box at future times, or a value < 0.0 to compress the box.

The variable *name2* must also be an *equal-style variable* and should calculate the rate of box length change, in units of distance/time, i.e. the time-derivative of the *name1* variable. This quantity is used internally by LAMMPS to reset atom velocities when they cross periodic boundaries. It is computed internally for the other styles, but you must provide it when using an arbitrary variable.

Here is an example of using the *variable* style to perform the same box deformation as the *wiggle* style formula listed above, where we assume that the current timestep = 0.

```
variable A equal 5.0
variable Tp equal 10.0
variable displace equal "v_A * sin(2*PI * step*dt/v_Tp)"
variable rate equal "2*PI*v_A/v_Tp * cos(2*PI * step*dt/v_Tp)"
fix 2 all deform 1 x variable v_displace v_rate remap v
```

For the *scale*, *vel*, *erate*, *trate*, *volume*, *wiggle*, and *variable* styles, the box length is expanded or compressed around its mid point.

---

For the *xy*, *xz*, and *yz* parameters, this is the meaning of their styles and values. Note that changing the tilt factors of a triclinic box does not change its volume.

The *final*, *delta*, *vel*, and *erate* styles all change the shear strain at a “constant engineering shear strain rate”. This means the tilt factor changes linearly with time from its initial to final value.

For style *final*, the final tilt factor is specified. The value can be in lattice or box distance units. See the discussion of the units keyword below.

For style *delta*, a plus or minus change in the tilt factor is specified. The value can be in lattice or box distance units. See the discussion of the units keyword below.

For style *vel*, a velocity at which the tilt factor changes is specified in units of distance/time. This is effectively an “engineering shear strain rate”, where rate = V/L0 and L0 is the initial box length perpendicular to the direction of shear. The distance can be in lattice or box distance units. See the discussion of the units keyword below. For example, if the initial tilt factor is 5 Angstroms, and the V is 10 Angstroms/ps, then after 1 ps, the tilt factor will be 15 Angstroms. After 2 ps, it will be 25 Angstroms.

The *erate* style changes a tilt factor at a “constant engineering shear strain rate”. The units of the specified shear strain rate are 1/time. See the *units* command for the time units associated with different choices of simulation units, e.g. picoseconds for “metal” units). Shear strain is unitless and is defined as offset/length, where length is the box length perpendicular to the shear direction (e.g. y box length for xy deformation) and offset is the displacement distance in the shear direction (e.g. x direction for xy deformation) from the unstrained orientation.

The tilt factor T as a function of time will change as

$$T(t) = T_0 + L_0 \cdot \text{erate} \cdot dt$$

where T0 is the initial tilt factor, L0 is the original length of the box perpendicular to the shear direction (e.g. y box length for xy deformation), and dt is the elapsed time (in time units). Thus if *erate* R is specified as 0.1 and time units are picoseconds, this means the shear strain will increase by 0.1 every picosecond. I.e. if the xy shear strain was initially 0.0, then strain after 1 ps = 0.1, strain after 2 ps = 0.2, etc. Thus the tilt factor would be 0.0 at time 0, 0.1\*ybox at 1 ps, 0.2\*ybox at 2 ps, etc, where ybox is the original y box length. R = 1 or 2 means the tilt factor will increase by 1 or 2 every picosecond. R = -0.01 means a decrease in shear strain by 0.01 every picosecond.

The *trate* style changes a tilt factor at a “constant true shear strain rate”. Note that this is not an “engineering shear strain rate”, as the other styles are. Rather, for a “true” rate, the rate of change is constant, which means the tilt factor

changes non-linearly with time from its initial to final value. The units of the specified shear strain rate are 1/time. See the *units* command for the time units associated with different choices of simulation units, e.g. picoseconds for “metal” units). Shear strain is unitless and is defined as offset/length, where length is the box length perpendicular to the shear direction (e.g. y box length for xy deformation) and offset is the displacement distance in the shear direction (e.g. x direction for xy deformation) from the unstrained orientation.

The tilt factor  $T$  as a function of time will change as

$$T(t) = T_0 \exp(\text{trate} * dt)$$

where  $T_0$  is the initial tilt factor and  $dt$  is the elapsed time (in time units). Thus if *trate*  $R$  is specified as  $\ln(1.1)$  and time units are picoseconds, this means the shear strain or tilt factor will increase by 10% every picosecond. I.e. if the xy shear strain was initially 0.1, then strain after 1 ps = 0.11, strain after 2 ps = 0.121, etc.  $R = \ln(2)$  or  $\ln(3)$  means the tilt factor will double or triple every picosecond.  $R = \ln(0.99)$  means the tilt factor will shrink by 1% every picosecond. Note that the change is continuous, so running with  $R = \ln(2)$  for 10 picoseconds does not change the tilt factor by a factor of 10, but by a factor of 1024 since it doubles every picosecond. Note that the initial tilt factor must be non-zero to use the *trate* option.

Note that shear strain is defined as the tilt factor divided by the perpendicular box length. The *erate* and *trate* styles control the tilt factor, but assume the perpendicular box length remains constant. If this is not the case (e.g. it changes due to another fix deform parameter), then this effect on the shear strain is ignored.

The *wiggle* style oscillates the specified tilt factor sinusoidally with the specified amplitude and period. I.e. the tilt factor  $T$  as a function of time is given by

$$T(t) = T_0 + A \sin(2 * \pi t / T_p)$$

where  $T_0$  is its initial value. If the amplitude  $A$  is a positive number the tilt factor initially becomes more positive, then more negative, etc. If  $A$  is negative then the tilt factor initially becomes more negative, then more positive, etc. The amplitude can be in lattice or box distance units. See the discussion of the units keyword below.

The *variable* style changes the specified tilt factor by evaluating a variable, which presumably is a function of time. The variable with *name1* must be an *equal-style variable* and should calculate a change in tilt in units of distance. Note that this distance is in box units, not lattice units; see the discussion of the *units* keyword below. The formula associated with variable *name1* can reference the current timestep. Note that it should return the “change” in tilt factor, not the absolute tilt factor. This means it should evaluate to 0.0 when invoked on the initial timestep of the run following the definition of fix deform.

The variable *name2* must also be an *equal-style variable* and should calculate the rate of tilt change, in units of distance/time, i.e. the time-derivative of the *name1* variable. This quantity is used internally by LAMMPS to reset atom velocities when they cross periodic boundaries. It is computed internally for the other styles, but you must provide it when using an arbitrary variable.

Here is an example of using the *variable* style to perform the same box deformation as the *wiggle* style formula listed above, where we assume that the current timestep = 0.

```
variable A equal 5.0
variable Tp equal 10.0
variable displace equal "v_A * sin(2*PI * step*dt/v_Tp)"
variable rate equal "2*PI*v_A/v_Tp * cos(2*PI * step*dt/v_Tp)"
fix 2 all deform 1 xy variable v_displace v_rate remap v
```

All of the tilt styles change the xy, xz, yz tilt factors during a simulation. In LAMMPS, tilt factors (xy,xz,yz) for triclinic boxes are normally bounded by half the distance of the parallel box length. See the discussion of the *flip* keyword below, to allow this bound to be exceeded, if desired.

For example, if  $x_{lo} = 2$  and  $x_{hi} = 12$ , then the x box length is 10 and the xy tilt factor must be between -5 and 5. Similarly, both xz and yz must be between  $-(x_{hi}-x_{lo})/2$  and  $+(y_{hi}-y_{lo})/2$ . Note that this is not a limitation, since if the maximum tilt factor is 5 (as in this example), then configurations with tilt = ..., -15, -5, 5, 15, 25, ... are all equivalent.

To obey this constraint and allow for large shear deformations to be applied via the *xy*, *xz*, or *yz* parameters, the following algorithm is used. If *prd* is the associated parallel box length (10 in the example above), then if the tilt factor exceeds the accepted range of -5 to 5 during the simulation, then the box is flipped to the other limit (an equivalent box) and the simulation continues. Thus for this example, if the initial *xy* tilt factor was 0.0 and “*xy final 100.0*” was specified, then during the simulation the *xy* tilt factor would increase from 0.0 to 5.0, the box would be flipped so that the tilt factor becomes -5.0, the tilt factor would increase from -5.0 to 5.0, the box would be flipped again, etc. The flip occurs 10 times and the final tilt factor at the end of the simulation would be 0.0. During each flip event, atoms are remapped into the new box in the appropriate manner.

The one exception to this rule is if the first dimension in the tilt factor (*x* for *xy*) is non-periodic. In that case, the limits on the tilt factor are not enforced, since flipping the box in that dimension does not change the atom positions due to non-periodicity. In this mode, if you tilt the system to extreme angles, the simulation will simply become inefficient due to the highly skewed simulation box.

---

Each time the box size or shape is changed, the *remap* keyword determines whether atom positions are remapped to the new box. If *remap* is set to *x* (the default), atoms in the *fix* group are remapped; otherwise they are not. Note that their velocities are not changed, just their positions are altered. If *remap* is set to *v*, then any atom in the *fix* group that crosses a periodic boundary will have a delta added to its velocity equal to the difference in velocities between the *lo* and *hi* boundaries. Note that this velocity difference can include tilt components, e.g. a delta in the *x* velocity when an atom crosses the *y* periodic boundary. If *remap* is set to *none*, then neither of these remappings take place.

Conceptually, setting *remap* to *x* forces the atoms to deform via an affine transformation that exactly matches the box deformation. This setting is typically appropriate for solids. Note that though the atoms are effectively “moving” with the box over time, it is not due to their having a velocity that tracks the box change, but only due to the remapping. By contrast, setting *remap* to *v* is typically appropriate for fluids, where you want the atoms to respond to the change in box size/shape on their own and acquire a velocity that matches the box change, so that their motion will naturally track the box without explicit remapping of their coordinates.

---

**Note:** When non-equilibrium MD (NEMD) simulations are performed using this *fix*, the option “*remap v*” should normally be used. This is because *fix nvt/sllod* adjusts the atom positions and velocities to induce a velocity profile that matches the changing box size/shape. Thus atom coordinates should NOT be remapped by *fix deform*, but velocities SHOULD be when atoms cross periodic boundaries, since that is consistent with maintaining the velocity profile already created by *fix nvt/sllod*. LAMMPS will warn you if the *remap* setting is not consistent with *fix nvt/sllod*.

---

**Note:** For non-equilibrium MD (NEMD) simulations using “*remap v*” it is usually desirable that the fluid (or flowing material, e.g. granular particles) stream with a velocity profile consistent with the deforming box. As mentioned above, using a thermostat such as *fix nvt/sllod* or *fix langevin* (with a bias provided by *compute temp/deform*), will typically accomplish that. If you do not use a thermostat, then there is no driving force pushing the atoms to flow in a manner consistent with the deforming box. E.g. for a shearing system the box deformation velocity may vary from 0 at the bottom to 10 at the top of the box. But the stream velocity profile of the atoms may vary from -5 at the bottom to +5 at the top. You can monitor these effects using the *fix ave/chunk*, *compute temp/deform*, and *compute temp/profile* commands. One way to induce atoms to stream consistent with the box deformation is to give them an initial velocity profile, via the *velocity ramp* command, that matches the box deformation rate. This also typically helps the system come to equilibrium more quickly, even if a thermostat is used.

---

**Note:** If a *fix rigid* is defined for rigid bodies, and *remap* is set to *x*, then the center-of-mass coordinates of rigid bodies will be remapped to the changing simulation box. This will be done regardless of whether atoms in the rigid bodies are in the *fix deform* group or not. The velocity of the centers of mass are not remapped even if *remap* is set to *v*, since *fix nvt/sllod* does not currently do anything special for rigid particles. If you wish to perform a NEMD simulation of rigid

---

particles, you can either thermostat them independently or include a background fluid and thermostat the fluid via *fix nvt/sllod*.

---

The *flip* keyword allows the tilt factors for a triclinic box to exceed half the distance of the parallel box length, as discussed above. If the *flip* value is set to *yes*, the bound is enforced by flipping the box when it is exceeded. If the *flip* value is set to *no*, the tilt will continue to change without flipping. Note that if you apply large deformations, this means the box shape can tilt dramatically LAMMPS will run less efficiently, due to the large volume of communication needed to acquire ghost atoms around a processor's irregular-shaped sub-domain. For extreme values of tilt, LAMMPS may also lose atoms and generate an error.

The *units* keyword determines the meaning of the distance units used to define various arguments. A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing. Note that the units choice also affects the *vel* style parameters since it is defined in terms of distance/time. Also note that the units keyword does not affect the *variable* style. You should use the *xlat*, *ylat*, *zlat* keywords of the *thermo\_style* command if you want to include lattice spacings in a variable formula.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 17.36.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix will restore the initial box settings from *binary restart files*, which allows the fix to be properly continue deformation, when using the start/stop options of the *run* command. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*.

This fix can perform deformation over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

#### 17.36.5 Restrictions

You cannot apply x, y, or z deformations to a dimension that is shrink-wrapped via the *boundary* command.

You cannot apply xy, yz, or xz deformations to a second dimension (y in xy) that is shrink-wrapped via the *boundary* command.

## 17.36.6 Related commands

*change\_box*

## 17.36.7 Default

The option defaults are `remap = x`, `flip = yes`, and `units = lattice`.

## 17.37 fix deposit command

### 17.37.1 Syntax

```
fix ID group-ID deposit N type M seed keyword values ...
```

- ID, group-ID are documented in *fix* command
- deposit = style name of this fix command
- N = # of atoms or molecules to insert
- type = atom type to assign to inserted atoms (offset for molecule insertion)
- M = insert a single atom or molecule every M steps
- seed = random # seed (positive integer)
- one or more keyword/value pairs may be appended to args
- keyword = *region* or *id* or *global* or *local* or *near* or *gaussian* or *attempt* or *rate* or *vx* or *vy* or *vz* or *mol* or *rigid* or *shake* or *units*

*region* value = region-ID

region-ID = ID of region to use as insertion volume

*id* value = *max* or *next*

*max* = atom ID for new atom(s) is max ID of all current atoms plus one

*next* = atom ID for new atom(s) increments by one for every deposition

*global* values = lo hi

lo,hi = put new atom/molecule a distance lo-hi above all other atoms (distance ↵  
↵units)

*local* values = lo hi delta

lo,hi = put new atom/molecule a distance lo-hi above any nearby atom beneath it ↵  
↵(distance units)

delta = lateral distance within which a neighbor is considered "nearby" (distance ↵  
↵units)

*near* value = R

R = only insert atom/molecule if further than R from existing particles (distance ↵  
↵units)

*gaussian* values = xmid ymid zmid sigma

xmid,ymid,zmid = center of the gaussian distribution (distance units)

sigma = width of gaussian distribution (distance units)

*attempt* value = Q

Q = attempt a single insertion up to Q times

*rate* value = V

V = z velocity (y in 2d) at which insertion volume moves (velocity units)

*vx* values = vxlo vxhi



```

    vxlo,vxhi = range of x velocities for inserted atom/molecule (velocity units)
    vy values = vylo vyhi
    vylo,vyhi = range of y velocities for inserted atom/molecule (velocity units)
    vz values = vzlo vzhi
    vzlo,vzhi = range of z velocities for inserted atom/molecule (velocity units)
    target values = tx ty tz
    tx,ty,tz = location of target point (distance units)
    mol value = template-ID
    template-ID = ID of molecule template specified in a separate molecule command
    molfrac values = f1 f2 ... fN
    f1 to fN = relative probability of creating each of N molecules in template-ID
    rigid value = fix-ID
    fix-ID = ID of fix rigid/small command
    shake value = fix-ID
    fix-ID = ID of fix shake command
    orient values = rx ry rz
    rx,ry,rz = vector to randomly rotate an inserted molecule around
    units value = lattice or box
    lattice = the geometry is defined in lattice units
    box = the geometry is defined in simulation box units

```

## 17.37.2 Examples

```

fix 3 all deposit 1000 2 100 29494 region myblock local 1.0 1.0 1.0 units box
fix 2 newatoms deposit 10000 1 500 12345 region disk near 2.0 vz -1.0 -0.8
fix 4 sputter deposit 1000 2 500 12235 region sphere vz -1.0 -1.0 target 5.0 5.0 0.0
↳units lattice
fix 5 insert deposit 200 2 100 777 region disk gaussian 5.0 5.0 9.0 1.0 units box

```

## 17.37.3 Description

Insert a single atom or molecule into the simulation domain every *M* timesteps until *N* atoms or molecules have been inserted. This is useful for simulating deposition onto a surface. For the remainder of this doc page, a single inserted atom or molecule is referred to as a “particle”.

If inserted particles are individual atoms, they are assigned the specified atom type. If they are molecules, the type of each atom in the inserted molecule is specified in the file read by the *molecule* command, and those values are added to the specified atom type. E.g. if the file specifies atom types 1,2,3, and those are the atom types you want for inserted molecules, then specify *type* = 0. If you specify *type* = 2, the in the inserted molecule will have atom types 3,4,5.

All atoms in the inserted particle are assigned to two groups: the default group “all” and the group specified in the *fix deposit* command (which can also be “all”).

If you are computing temperature values which include inserted particles, you will want to use the *compute\_modify* dynamic option, which insures the current number of atoms is used as a normalizing factor each time the temperature is computed.

Care must be taken that inserted particles are not too near existing atoms, using the options described below. When inserting particles above a surface in a non-periodic box (see the *boundary* command), the possibility of a particle escaping the surface and flying upward should be considered, since the particle may be lost or the box size may grow infinitely large. A *fix wall/reflect* command can be used to prevent this behavior. Note that if a shrink-wrap boundary is used, it is OK to insert the new particle outside the box, however the box will immediately be expanded to include the new particle. When simulating a sputtering experiment it is probably more realistic to ignore those atoms using the *thermo\_modify* command with the *lost ignore* option and a fixed *boundary*.

The fix deposit command must use the *region* keyword to define an insertion volume. The specified region must have been previously defined with a *region* command. It must be defined with *side = in*.

---

**Note:** LAMMPS checks that the specified region is wholly inside the simulation box. It can do this correctly for orthonormal simulation boxes. However for *triclinic boxes*, it only tests against the larger orthonormal box that bounds the tilted simulation box. If the specified region includes volume outside the tilted box, then an insertion will likely fail, leading to a “lost atoms” error. Thus for triclinic boxes you should insure the specified region is wholly inside the simulation box.

---

The locations of inserted particles are taken from uniform distributed random numbers, unless the *gaussian* keyword is used. Then the individual coordinates are taken from a gaussian distribution of width *sigma* centered on *xmid,ymid,zmid*.

Individual atoms are inserted, unless the *mol* keyword is used. It specifies a *template-ID* previously defined using the *molecule* command, which reads files that define one or more molecules. The coordinates, atom types, charges, etc, as well as any bond/angle/etc and special neighbor information for the molecule can be specified in the molecule file. See the *molecule* command for details. The only settings required to be in each file are the coordinates and types of atoms in the molecule.

If the molecule template contains more than one molecule, the relative probability of depositing each molecule can be specified by the *molfrac* keyword. N relative probabilities, each from 0.0 to 1.0, are specified, where N is the number of molecules in the template. Each time a molecule is deposited, a random number is used to sample from the list of relative probabilities. The N values must sum to 1.0.

If you wish to insert molecules via the *mol* keyword, that will be treated as rigid bodies, use the *rigid* keyword, specifying as its value the ID of a separate *fix rigid/small* command which also appears in your input script.

---

**Note:** If you wish the new rigid molecules (and other rigid molecules) to be thermostatted correctly via *fix rigid/small/nvt* or *fix rigid/small/npt*, then you need to use the “fix\_modify dynamic/dof yes” command for the rigid fix. This is to inform that fix that the molecule count will vary dynamically.

---

If you wish to insert molecules via the *mol* keyword, that will have their bonds or angles constrained via SHAKE, use the *shake* keyword, specifying as its value the ID of a separate *fix shake* command which also appears in your input script.

Each timestep a particle is inserted, the coordinates for its atoms are chosen as follows. For insertion of individual atoms, the “position” referred to in the following description is the coordinate of the atom. For insertion of molecule, the “position” is the geometric center of the molecule; see the *molecule* doc page for details. A random rotation of the molecule around its center point is performed, which determines the coordinates all the individual atoms.

A random position within the region insertion volume is generated. If neither the *global* or *local* keyword is used, the random position is the trial position. If the *global* keyword is used, the random x,y values are used, but the z position of the new particle is set above the highest current atom in the simulation by a distance randomly chosen between lo/hi. (For a 2d simulation, this is done for the y position.) If the *local* keyword is used, the z position is set a distance between lo/hi above the highest current atom in the simulation that is “nearby” the chosen x,y position. In this context, “nearby” means the lateral distance (in x,y) between the new and old particles is less than the *delta* setting.

Once a trial x,y,z position has been selected, the insertion is only performed if no current atom in the simulation is within a distance R of any atom in the new particle, including the effect of periodic boundary conditions if applicable. R is defined by the *near* keyword. Note that the default value for R is 0.0, which will allow atoms to strongly overlap if you are inserting where other atoms are present. This distance test is performed independently for each atom in an inserted molecule, based on the randomly rotated configuration of the molecule. If this test fails, a new random position within the insertion volume is chosen and another trial is made. Up to Q attempts are made. If the particle is not successfully inserted, LAMMPS prints a warning message.

---

**Note:** If you are inserting finite size particles or a molecule or rigid body consisting of finite-size particles, then you should typically set *R* larger than the distance at which any inserted particle may overlap with either a previously inserted particle or an existing particle. LAMMPS will issue a warning if *R* is smaller than this value, based on the radii of existing and inserted particles.

---

The *rate* option moves the insertion volume in the *z* direction (3d) or *y* direction (2d). This enables particles to be inserted from a successively higher height over time. Note that this parameter is ignored if the *global* or *local* keywords are used, since those options choose a *z*-coordinate for insertion independently.

The *vx*, *vy*, and *vz* components of velocity for the inserted particle are set using the values specified for the *vx*, *vy*, and *vz* keywords. Note that normally, new particles should be assigned a negative vertical velocity so that they move towards the surface. For molecules, the same velocity is given to every particle (no rotation or bond vibration).

If the *target* option is used, the velocity vector of the inserted particle is changed so that it points from the insertion position towards the specified target point. The magnitude of the velocity is unchanged. This can be useful, for example, for simulating a sputtering process. E.g. the target point can be far away, so that all incident particles strike the surface as if they are in an incident beam of particles at a prescribed angle.

The *orient* keyword is only used when molecules are deposited. By default, each molecule is inserted at a random orientation. If this keyword is specified, then (*rx,ry,rz*) is used as an orientation vector, and each inserted molecule is rotated around that vector with a random value from zero to  $2\pi$ . For a 2d simulation, *rx = ry = 0.0* is required, since rotations can only be performed around the *z* axis.

The *id* keyword determines how atom IDs and molecule IDs are assigned to newly deposited particles. Molecule IDs are only assigned if molecules are being inserted. For the *max* setting, the atom and molecule IDs of all current atoms are checked. Atoms in the new particle are assigned IDs starting with the current maximum plus one. If a molecule is inserted it is assigned an ID = current maximum plus one. This means that if particles leave the system, the new IDs may replace the lost ones. For the *next* setting, the maximum ID of any atom and molecule is stored at the time the fix is defined. Each time a new particle is added, this value is incremented to assign IDs to the new atom(s) or molecule. Thus atom and molecule IDs for deposited particles will be consecutive even if particles leave the system over time.

The *units* keyword determines the meaning of the distance units used for the other deposition parameters. A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing. Note that the units choice affects all the keyword values that have units of distance or velocity.

---

**Note:** If you are monitoring the temperature of a system where the atom count is changing due to adding particles, you typically should use the *compute\_modify dynamic yes* command for the temperature compute you are using.

---

#### 17.37.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the deposition to *binary restart files*. This includes information about how many particles have been deposited, the random number generator seed, the next timestep for deposition, etc. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

---

**Note:** For this to work correctly, the timestep must **not** be changed after reading the restart with *reset\_timestep*. The fix will try to detect it and stop with an error.

---

None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.37.5 Restrictions

The specified insertion region cannot be a “dynamic” region, as defined by the *region* command.

### 17.37.6 Related commands

*fix pour, region*

### 17.37.7 Default

Insertions are performed for individual atoms, i.e. no *mol* setting is defined. If the *mol* keyword is used, the default for *molfrac* is an equal probabilities for all molecules in the template. Additional option defaults are *id* = max, *delta* = 0.0, *near* = 0.0, *attempt* = 10, *rate* = 0.0, *vx* = 0.0 0.0, *vy* = 0.0 0.0, *vz* = 0.0 0.0, and *units* = lattice.

## 17.38 fix dpd/energy command

Accelerator Variants: *dpd/energy/kk*

### 17.38.1 Syntax

```
fix ID group-ID dpd/energy
```

- ID, group-ID are documented in *fix* command
- dpd/energy = style name of this fix command

### 17.38.2 Examples

```
fix 1 all dpd/energy
```

### 17.38.3 Description

Perform constant energy dissipative particle dynamics (DPD-E) integration. This fix updates the internal energies for particles in the group at each timestep. It must be used in conjunction with a deterministic integrator (e.g. *fix nve*) that updates the particle positions and velocities.

For fix *dpd/energy*, the particle internal temperature is related to the particle internal energy through a mesoparticle equation of state. An additional fix must be specified that defines the equation of state for each particle, e.g. *fix eos/cv*.

This fix must be used with the *pair\_style dpd/fdt/energy* command.

Note that numerous variants of DPD can be specified by choosing an appropriate combination of the integrator and *pair\_style dpd/fdt/energy* command. DPD under isoenergetic conditions can be specified by using *fix dpd/energy*, *fix nve* and *pair\_style dpd/fdt/energy*. DPD under isoenthalpic conditions can be specified by using *fix dpd/energy*, *fix npf* and *pair\_style dpd/fdt/energy*. Examples of each DPD variant are provided in the examples/PACKAGES/dpd-react directory.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 17.38.4 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This fix must be used with an additional fix that specifies time integration, e.g. *fix nve*.

The fix *dpd/energy* requires the *dpd atom\_style* to be used in order to properly account for the particle internal energies and temperature.

The fix *dpd/energy* must be used with an additional fix that specifies the mesoparticle equation of state for each particle.

## 17.38.5 Related commands

*fix nve* *fix eos/cv*

## 17.38.6 Default

none

---

**(Lisal)** M. Lisal, J.K. Brennan, J. Bonet Avalos, “Dissipative particle dynamics at isothermal, isobaric, isoenergetic, and isoenthalpic conditions using Shardlow-like splitting algorithms.”, *J. Chem. Phys.*, 135, 204105 (2011).

**(Larentzos)** J.P. Larentzos, J.K. Brennan, J.D. Moore, and W.D. Mattson, “LAMMPS Implementation of Constant Energy Dissipative Particle Dynamics (DPD-E)”, ARL-TR-6863, U.S. Army Research Laboratory, Aberdeen Proving Ground, MD (2014).

## 17.39 fix edpd/source command

## 17.40 fix tdpd/source command

### 17.40.1 Syntax

```
fix ID group-ID edpd/source keyword values ...
fix ID group-ID tdpd/source cc_index keyword values ...
```

- ID, group-ID are documented in *fix* command
- *edpd/source* or *tdpd/source* = style name of this fix command
- index (only specified for *tdpd/source*) = index of chemical species (1 to Nspecies)
- keyword = *sphere* or *cuboid*

*sphere* values = *cx,cy,cz,radius,source*

*cx,cy,cz* = *x,y,z* center of spherical domain (distance units)

*radius* = radius of a spherical domain (distance units)

*source* = heat source or concentration source (flux units, see below)

*cuboid* values = *cx,cy,cz,dLx,dLy,dLz,source*

*cx,cy,cz* = *x,y,z* lower left corner of a cuboid domain (distance units)

*dLx,dLy,dLz* = *x,y,z* side length of a cuboid domain (distance units)

*source* = heat source or concentration source (flux units, see below)

## 17.40.2 Examples

```
fix 1 all edpd/source sphere 0.0 0.0 0.0 5.0 0.01
fix 1 all edpd/source cuboid 0.0 0.0 0.0 20.0 10.0 10.0 -0.01
fix 1 all tdpd/source 1 sphere 5.0 0.0 0.0 5.0 0.01
fix 1 all tdpd/source 2 cuboid 0.0 0.0 0.0 20.0 10.0 10.0 0.01
```

## 17.40.3 Description

Fix *edpd/source* adds a heat source as an external heat flux to each atom in a spherical or cuboid domain, where the *source* is in units of energy/time. Fix *tdpd/source* adds an external concentration source of the chemical species specified by *index* as an external concentration flux for each atom in a spherical or cuboid domain, where the *source* is in units of mole/volume/time.

This command can be used to give an additional heat/concentration source term to atoms in a simulation, such as for a simulation of a heat conduction with a source term (see Fig.12 in (Li2014)) or diffusion with a source term (see Fig.1 in (Li2015)), as an analog of a periodic Poiseuille flow problem.

If the *sphere* keyword is used, the *cx,cy,cz,radius* defines a spherical domain to apply the source flux to.

If the *cuboid* keyword is used, the *cx,cy,cz,dLx,dLy,dLz* defines a cuboid domain to apply the source flux to.

---

## 17.40.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.40.5 Restrictions

This fix is part of the DPD-MESO package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Fix *edpd/source* must be used with the *pair\_style edpd* command. Fix *tdpd/source* must be used with the *pair\_style tdpd* command.

## 17.40.6 Related commands

*pair\_style edpd*, *pair\_style tdpd*, *compute edpd/temp/atom*, *compute tdpd/cc/atom*

## 17.40.7 Default

none

**(Li2014)** Z. Li, Y.-H. Tang, H. Lei, B. Caswell and G.E. Karniadakis, “Energy-conserving dissipative particle dynamics with temperature-dependent properties”, J. Comput. Phys., 265: 113-127 (2014). DOI: 10.1016/j.jcp.2014.02.003

**(Li2015)** Z. Li, A. Yazdani, A. Tartakovsky and G.E. Karniadakis, “Transport dissipative particle dynamics model for mesoscopic advection-diffusion-reaction problems”, J. Chem. Phys., 143: 014101 (2015). DOI: 10.1063/1.4923254

## 17.41 fix drag command

### 17.41.1 Syntax

```
fix ID group-ID drag x y z fmag delta
```

- ID, group-ID are documented in *fix* command
- drag = style name of this fix command
- x,y,z = coord to drag atoms towards
- fmag = magnitude of force to apply to each atom (force units)
- delta = cutoff distance inside of which force is not applied (distance units)

### 17.41.2 Examples

```
fix center small-molecule drag 0.0 10.0 0.0 5.0 2.0
```

### 17.41.3 Description

Apply a force to each atom in a group to drag it towards the point (x,y,z). The magnitude of the force is specified by `fmag`. If an atom is closer than a distance `delta` to the point, then the force is not applied.

Any of the x,y,z values can be specified as NULL which means do not include that dimension in the distance calculation or force application.

This command can be used to steer one or more atoms to a new location in the simulation.

### 17.41.4 Restart, `fix_modify`, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The `fix_modify respa` option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global 3-vector of forces, which can be accessed by various *output commands*. This is the total force on the group of atoms by the drag force. The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the `run` command. This fix is not invoked during *energy minimization*.

### 17.41.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.41.6 Related commands

*fix spring*, *fix spring/self*, *fix spring/rg*, *fix smd*

### 17.41.7 Default

none

## 17.42 `fix drude` command

### 17.42.1 Syntax

```
fix ID group-ID drude flag1 flag2 ... flagN
```

- ID, group-ID are documented in *fix* command
- drude = style name of this fix command
- flag1 flag2 ... flagN = Drude flag for each atom type (1 to N) in the system



## 17.42.2 Examples

```
fix 1 all drude 1 1 0 1 0 2 2 2
fix 1 all drude C C N C N D D D
```

Example input scripts available: examples/PACKAGES/drude

## 17.42.3 Description

Assign each atom type in the system to be one of 3 kinds of atoms within the Drude polarization model. This fix is designed to be used with the *thermalized Drude oscillator model*. Polarizable models in LAMMPS are described on the *Howto polarizable* doc page.

The three possible types can be designated with an integer (0,1,2) or capital letter (N,C,D):

- 0 or N = non-polarizable atom (not part of Drude model)
- 1 or C = Drude core
- 2 or D = Drude electron

## 17.42.4 Restrictions

This fix should be invoked before any other commands that implement the Drude oscillator model, such as *fix langevin/drude*, *fix tgnvt/drude*, *fix drude/transform*, *compute temp/drude*, *pair\_style thole*.

## 17.42.5 Related commands

*fix langevin/drude*, *fix tgnvt/drude*, *fix drude/transform*, *compute temp/drude*, *pair\_style thole*

## 17.42.6 Default

none

## 17.43 fix drude/transform/direct command

## 17.44 fix drude/transform/inverse command

### 17.44.1 Syntax

```
fix ID group-ID style keyword value ...
```

- ID, group-ID are documented in *fix* command
- style = *drude/transform/direct* or *drude/transform/inverse*

## 17.44.2 Examples

```
fix 3 all drude/transform/direct
fix 1 all drude/transform/inverse
```

Example input scripts available: examples/PACKAGES/drude

## 17.44.3 Description

Transform the coordinates of Drude oscillators from real to reduced and back for thermalizing the Drude oscillators as described in (*Lamoureux*) using a Nose-Hoover thermostat. This fix is designed to be used with the *thermalized Drude oscillator model*. Polarizable models in LAMMPS are described on the *Howto polarizable* doc page.

Drude oscillators are a pair of atoms representing a single polarizable atom. Ideally, the mass of Drude particles would vanish and their positions would be determined self-consistently by iterative minimization of the energy, the cores' positions being fixed. It is however more efficient and it yields comparable results, if the Drude oscillators (the motion of the Drude particle relative to the core) are thermalized at a low temperature. In that case, the Drude particles need a small mass.

The thermostats act on the reduced degrees of freedom, which are defined by the following equations. Note that in these equations upper case denotes atomic or center of mass values and lower case denotes Drude particle or dipole values. Primes denote the transformed (reduced) values, while bare letters denote the original values.

Masses:

$$M' = M + m$$

$$m' = \frac{M m}{M'}$$

Positions:

$$X' = \frac{M X + m x}{M'}$$

$$x' = x - X$$

Velocities:

$$V' = \frac{M V + m v}{M'}$$

$$v' = v - V$$

Forces:

$$F' = F + f$$

$$f' = \frac{M f - m F}{M'}$$

This transform conserves the total kinetic energy

$$\frac{1}{2} (M V^2 + m v^2) = \frac{1}{2} (M' V'^2 + m' v'^2)$$

and the virial defined with absolute positions

$$X F + x f = X' F' + x' f'$$

This fix requires each atom know whether it is a Drude particle or not. You must therefore use the *fix drude* command to specify the Drude status of each atom type.

**Note:** only the Drude core atoms need to be in the group specified for this fix. A Drude electron will be transformed together with its core even if it is not itself in the group. It is safe to include Drude electrons or non-polarizable atoms in the group. The non-polarizable atoms will simply not be transformed.

This fix does NOT perform time integration. It only transform masses, coordinates, velocities and forces. Thus you must use separate time integration fixes, like *fix nve* or *fix npt* to actually update the velocities and positions of atoms. In order to thermalize the reduced degrees of freedom at different temperatures, two Nose-Hoover thermostats must be defined, acting on two distinct groups.

**Note:** The *fix drude/transform/direct* command must appear before any Nose-Hoover thermostating fixes. The *fix drude/transform/inverse* command must appear after any Nose-Hoover thermostating fixes.

Example:

```
fix fDIRECT all drude/transform/direct
fix fNVT gCORES nvt temp 300.0 300.0 100
fix fNVT gDRUDES nvt temp 1.0 1.0 100
fix fINVERSE all drude/transform/inverse
compute TDRUDE all temp/drude
thermo_style custom step cpu etotal ke pe ebond ecoul elong press vol temp c_TDRUDE[1] c_
↪TDRUDE[2]
```

In this example, *gCORES* is the group of the atom cores and *gDRUDES* is the group of the Drude particles (electrons). The centers of mass of the Drude oscillators will be thermostatted at 300.0 and the internal degrees of freedom will be thermostatted at 1.0. The temperatures of cores and Drude particles, in center-of-mass and relative coordinates, are calculated using *compute temp/drude*

In addition, if you want to use a barostat to simulate a system at constant pressure, only one of the Nose-Hoover fixes must be *npt*, the other one should be *nvt*. You must add a *compute temp/com* and a *fix\_modify* command so that the temperature of the *npt* fix be just that of its group (the Drude cores) but the pressure be the overall pressure *thermo\_press*.

Example:

```
compute cTEMP_CORE gCORES temp/com
fix fDIRECT all drude/transform/direct
fix fNPT gCORES npt temp 298.0 298.0 100 iso 1.0 1.0 500
fix_modify fNPT temp cTEMP_CORE press thermo_press
fix fNVT gDRUDES nvt temp 5.0 5.0 100
fix fINVERSE all drude/transform/inverse
```

In this example, *gCORES* is the group of the atom cores and *gDRUDES* is the group of the Drude particles. The centers of mass of the Drude oscillators will be thermostatted at 298.0 and the internal degrees of freedom will be thermostatted at 5.0. The whole system will be barostatted at 1.0.

In order to avoid the flying ice cube problem (irreversible transfer of linear momentum to the center of mass of the system), you may need to add a *fix momentum* command:

```
fix fMOMENTUM all momentum 100 linear 1 1 1
```

---

#### 17.44.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

#### 17.44.5 Restrictions

none

#### 17.44.6 Related commands

*fix drude, fix langevin/drude, compute temp/drude, pair\_style thole*

#### 17.44.7 Default

none

---

(Lamoureux) Lamoureux and Roux, J Chem Phys, 119, 3025-3039 (2003).

### 17.45 fix dt/reset command

#### 17.45.1 Syntax

```
fix ID group-ID dt/reset N Tmin Tmax Xmax keyword values ...
```

- ID, group-ID are documented in *fix* command
- dt/reset = style name of this fix command
- N = re-compute dt every N timesteps
- Tmin = minimum dt allowed which can be NULL (time units)
- Tmax = maximum dt allowed which can be NULL (time units)
- Xmax = maximum distance for an atom to move in one timestep (distance units)
- zero or more keyword/value pairs may be appended
- keyword = *emax* or *units*

*emax* value = Emax

Emax = maximum kinetic energy change for an atom in one timestep (energy units)

*units* value = *lattice* or *box*

lattice = Xmax is defined in lattice units

box = Xmax is defined in simulation box units

## 17.45.2 Examples

```
fix 5 all dt/reset 10 1.0e-5 0.01 0.1
fix 5 all dt/reset 10 0.01 2.0 0.2 units box
fix 5 all dt/reset 5 NULL 0.001 0.5 emax 30 units box
```

## 17.45.3 Description

Reset the timestep size every  $N$  steps during a run, so that no atom moves further than the specified  $X_{max}$  distance, based on current atom velocities and forces. Optionally an additional criterion is imposed by the *emax* keyword, so that no atom's kinetic energy changes by more than the specified  $E_{max}$ .

This can be useful when starting from a configuration with overlapping atoms, where forces will be large. Or it can be useful when running an impact simulation where one or more high-energy atoms collide with a solid, causing a damage cascade.

This fix overrides the timestep size setting made by the *timestep* command. The new timestep size  $dt$  is computed in the following manner.

For each atom, the timestep is computed that would cause it to displace  $X_{max}$  on the next integration step, as a function of its current velocity and force. Since performing this calculation exactly would require the solution to a quartic equation, a cheaper estimate is generated. The estimate is conservative in that the atom's displacement is guaranteed not to exceed  $X_{max}$ , though it may be smaller.

In addition if the *emax* keyword is used, the specified  $E_{max}$  value is enforced as a limit on how much an atom's kinetic energy can change. If the timestep required is even smaller than for the  $X_{max}$  displacement, then the smaller timestep is used.

Given this putative timestep for each atom, the minimum timestep value across all atoms is computed. Then the  $T_{min}$  and  $T_{max}$  bounds are applied, if specified. If one (or both) is specified as NULL, it is not applied.

When the *run style* is *respa*, this fix resets the outer loop (largest) timestep, which is the same timestep that the *timestep* command sets.

Note that the cumulative simulation time (in time units), which accounts for changes in the timestep size as a simulation proceeds, can be accessed by the *thermo\_style time* keyword.

Also note that the *dump\_modify every/time* option allows dump files to be written at intervals specified by simulation time, rather than by timesteps. Simulation time is in time units; see the *units* doc page for details.

## 17.45.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar stores the last timestep on which the timestep was reset to a new value.

The scalar value calculated by this fix is “intensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.45.5 Restrictions

none

## 17.45.6 Related commands

*timestep*, *dump\_modify every/time*

## 17.45.7 Default

The option defaults are units = lattice, and no emax kinetic energy limit.

# 17.46 fix efield command

## 17.46.1 Syntax

```
fix ID group-ID efield ex ey ez keyword value ...
```

- ID, group-ID are documented in *fix* command
- efield = style name of this fix command
- ex,ey,ez = E-field component values (electric field units)
- any of ex,ey,ez can be a variable (see below)
- zero or more keyword/value pairs may be appended to args
- keyword = *region* or *energy*

```
region value = region-ID
```

```
region-ID = ID of region atoms must be in to have added force
```

```
energy value = v_name
```

```
v_name = variable with name that calculates the potential energy of each atom in  
→the added E-field
```

## 17.46.2 Examples

```
fix kick external-field efield 1.0 0.0 0.0  
fix kick external-field efield 0.0 0.0 v_oscillate
```

## 17.46.3 Description

Add a force  $F = qE$  to each charged atom in the group due to an external electric field being applied to the system. If the system contains point-dipoles, also add a torque on the dipoles due to the external electric field.

For charges, any of the 3 quantities defining the E-field components can be specified as an equal-style or atom-style *variable*, namely *ex*, *ey*, *ez*. If the value is a variable, it should be specified as *v\_name*, where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the E-field component.

For point-dipoles, equal-style variables can be used, but atom-style variables are not currently supported, since they imply a spatial gradient in the electric field which means additional terms with gradients of the field are required for the force and torque on dipoles.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent E-field.

Atom-style variables can specify the same formulas as equal-style variables but can also include per-atom values, such as atom coordinates. Thus it is easy to specify a spatially-dependent E-field with optional time-dependence as well.

If the *region* keyword is used, the atom must also be in the specified geometric *region* in order to have force added to it.

---

Adding a force or torque to atoms implies a change in their potential energy as they move or rotate due to the applied E-field.

For dynamics via the “run” command, this energy can be optionally added to the system’s potential energy for thermo-dynamic output (see below). For energy minimization via the “minimize” command, this energy must be added to the system’s potential energy to formulate a self-consistent minimization problem (see below).

The *energy* keyword is not allowed if the added field is a constant vector (*ex,ey,ez*), with all components defined as numeric constants and not as variables. This is because LAMMPS can compute the energy for each charged particle directly as  $E = -x \cdot qE = -q(x*ex + y*ey + z*ez)$ , so that  $-\text{Grad}(E) = F$ . Similarly for point-dipole particles the energy can be computed as  $E = -\mu \cdot E = -(\mu_x*ex + \mu_y*ey + \mu_z*ez)$ .

The *energy* keyword is optional if the added force is defined with one or more variables, and if you are performing dynamics via the *run* command. If the keyword is not used, LAMMPS will set the energy to 0.0, which is typically fine for dynamics.

The *energy* keyword is required if the added force is defined with one or more variables, and you are performing energy minimization via the “minimize” command for charged particles. It is not required for point-dipoles, but a warning is issued since the minimizer in LAMMPS does not rotate dipoles, so you should not expect to be able to minimize the orientation of dipoles in an applied electric field.

The *energy* keyword specifies the name of an atom-style *variable* which is used to compute the energy of each atom as function of its position. Like variables used for *ex, ey, ez*, the energy variable is specified as *v\_name*, where name is the variable name.

Note that when the *energy* keyword is used during an energy minimization, you must insure that the formula defined for the atom-style *variable* is consistent with the force variable formulas, i.e. that  $-\text{Grad}(E) = F$ . For example, if the force due to the electric field were a spring-like  $F = kx$ , then the energy formula should be  $E = -0.5kx^2$ . If you don’t do this correctly, the minimization will not converge properly.

---

#### 17.46.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy inferred by the added force due to the electric field to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*. Note that this energy is a fictitious quantity but is needed so that the *minimize* command can include the forces added by this fix in a consistent manner. I.e. there is a decrease in potential energy when atoms move in the direction of the added force due to the electric field.

The *fix\_modify virial* option is supported by this fix to add the contribution due to the added forces on atoms to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix adding its forces. Default is the outermost level.

This fix computes a global scalar and a global 3-vector of forces, which can be accessed by various *output commands*. The scalar is the potential energy discussed above. The vector is the total force added to the group of atoms. The scalar and vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command. You should not specify force components with a variable that has time-dependence for use with a minimizer, since the minimizer increments the timestep as the iteration count during the minimization.

---

**Note:** If you want the fictitious potential energy associated with the added forces to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

## 17.46.5 Restrictions

None

## 17.46.6 Related commands

*fix addforce*

## 17.46.7 Default

none

## 17.47 fix ehex command

### 17.47.1 Syntax

<code>fix ID group-ID ehex nevery F keyword value</code>
--

- ID, group-ID are documented in *fix* command
- ehex = style name of this fix command
- nevery = add/subtract heat every this many timesteps
- F = energy flux into the reservoir (energy/time units)
- zero or more keyword/value pairs may be appended to args
- keyword = *region* or *constrain* or *com* or *hex*



```

region value = region-ID
  region-ID = ID of region (reservoir) atoms must be in for added thermostating
↔force
constrain value = none
  apply the constraint algorithm (SHAKE or RATTLE) again at the end of the timestep
com value = none
  rescale all sites of a constrained cluster of atom if its COM is in the reservoir
hex value = none
  omit the coordinate correction to recover the HEX algorithm

```

## 17.47.2 Examples

```

# Lennard-Jones, from examples/in.ehex.lj
fix fnve all nve
# specify regions rhot and rcold
...
fix fhot all ehex 1 0.15 region rhot
fix fcold all ehex 1 -0.15 region rcold

# SPC/E water, from examples/in.ehex.spce
fix fnve all nve
# specify regions rhot and rcold
...
fix fhot all ehex 1 0.075 region rhot constrain com
fix fcold all ehex 1 -0.075 region rcold constrain com
fix frattle all rattle 1e-10 400 0 b 1 a 1

```

## 17.47.3 Description

This fix implements the asymmetric version of the enhanced heat exchange algorithm (*Wirnsberger*). The eHEX algorithm is an extension of the heat exchange algorithm (*Ikeshoji*) and adds an additional coordinate integration to account for higher-order truncation terms in the operator splitting. The original HEX algorithm (implemented as *fix heat*) is known to exhibit a slight energy drift limiting the accessible simulation times to a few nanoseconds. This issue is greatly improved by the new algorithm decreasing the energy drift by at least a factor of a hundred (LJ and SPC/E water) with little computational overhead.

In both algorithms (non-translational) kinetic energy is constantly swapped between regions (reservoirs) to impose a heat flux onto the system. The equations of motion are therefore modified if a particle  $i$  is located inside a reservoir  $\Gamma_k$  where  $k > 0$ . We use  $\Gamma_0$  to label those parts of the simulation box which are not thermostatted.) The input parameter *region-ID* of this fix corresponds to  $k$ . The energy swap is modelled by introducing an additional thermostating force to the equations of motion, such that the time evolution of coordinates and momenta of particle  $i$  becomes (*Wirnsberger*)

$$\begin{aligned}\dot{\mathbf{r}}_i &= \mathbf{v}_i, \\ \dot{\mathbf{v}}_i &= \frac{\mathbf{f}_i}{m_i} + \frac{\mathbf{g}_i}{m_i}.\end{aligned}$$

The thermostating force is given by

$$\mathbf{g}_i = \begin{cases} \frac{m_i}{2} \frac{F_{\Gamma_k(\mathbf{r}_i)}}{K_{\Gamma_k(\mathbf{r}_i)}} \left( \mathbf{v}_i - \mathbf{v}_{\Gamma_k(\mathbf{r}_i)} \right) & k(\mathbf{r}_i) > 0 \text{ (inside a reservoir),} \\ 0 & \text{otherwise,} \end{cases}$$

where  $m_i$  is the mass and  $k(\mathbf{r}_i)$  maps the particle position to the respective reservoir. The quantity  $F_{\Gamma_{k(\mathbf{r}_i)}}$  corresponds to the input parameter  $F$ , which is the energy flux into the reservoir. Furthermore,  $K_{\Gamma_{k(\mathbf{r}_i)}}$  and  $v_{\Gamma_{k(\mathbf{r}_i)}}$  denote the non-translational kinetic energy and the center of mass velocity of that reservoir. The thermostating force does not affect the center of mass velocities of the individual reservoirs and the entire simulation box. A derivation of the equations and details on the numerical implementation with velocity Verlet in LAMMPS can be found in reference “(Wirnsberger)”#\_Wirnsberger.

---

**Note:** This fix only integrates the thermostating force and must be combined with another integrator, such as *fix nve*, to solve the full equations of motion.

---

This fix is different from a thermostat such as *fix nvt* or *fix temp/rescale* in that energy is added/subtracted continuously. Thus if there is not another mechanism in place to counterbalance this effect, the entire system will heat or cool continuously.

---

**Note:** If heat is subtracted from the system too aggressively so that the group’s kinetic energy would go to zero, then LAMMPS will halt with an error message. Increasing the value of *nevery* means that heat is added/subtracted less frequently but in larger portions. The resulting temperature profile will therefore be the same.

---

This fix will default to *fix\_heat* (HEX algorithm) if the keyword *hex* is specified.

---

#### Compatibility with SHAKE and RATTLE (rigid molecules):

This fix is compatible with *fix shake* and *fix rattle*. If either of these constraining algorithms is specified in the input script and the keyword *constrain* is set, the bond distances will be corrected a second time at the end of the integration step. It is recommended to specify the keyword *com* in addition to the keyword *constrain*. With this option all sites of a constrained cluster are rescaled, if its center of mass is located inside the region. Rescaling all sites of a cluster by the same factor does not introduce any velocity components along fixed bonds. No rescaling takes place if the center of mass lies outside the region.

---

**Note:** You can only use the keyword *com* along with *constrain*.

---

To achieve the highest accuracy it is recommended to use *fix rattle* with the keywords *constrain* and *com* as shown in the second example. Only if RATTLE is employed, the velocity constraints will be satisfied.

---

**Note:** Even if RATTLE is used and the keywords *com* and *constrain* are both set, the coordinate constraints will not necessarily be satisfied up to the target precision. The velocity constraints are satisfied as long as all sites of a cluster are rescaled (keyword *com*) and the cluster does not span adjacent reservoirs. The current implementation of the eHEX algorithm introduces a small error in the bond distances, which goes to zero with order three in the timestep. For example, in a simulation of SPC/E water with a timestep of 2 fs the maximum relative error in the bond distances was found to be on the order of  $10^{-7}$  for relatively large temperature gradients. A higher precision can be achieved by decreasing the timestep.

---

---

### 17.47.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.47.5 Restrictions

This fix is part of the RIGID package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.47.6 Related commands

*fix heat, fix thermal/conductivity, compute temp, compute temp/region*

### 17.47.7 Default

none

---

(Ikeshoji) Ikeshoji and Hafskjold, Molecular Physics, 81, 251-261 (1994).

(Wirnsberger) Wirnsberger, Frenkel, and Dellago, J Chem Phys, 143, 124104 (2015).

## 17.48 fix electrode/conp command

Accelerator Variant: *electrode/conp/intel*

## 17.49 fix electrode/conq command

Accelerator Variant: *electrode/conq/intel*

## 17.50 fix electrode/thermo command

Accelerator Variant: *electrode/thermo/intel*

### 17.50.1 Syntax

```
fix ID group-ID style args keyword value ...
```

- ID, group-ID are documented in *fix* command
- style = *electrode/conp* or *electrode/conq* or *electrode/thermo*
- args = arguments used by a particular style

```

electrode/conp args = potential eta
electrode/conq args = charge eta
electrode/thermo args = potential eta temp values
  potential = electrode potential
  charge = electrode charge
  eta = reciprocal width of electrode charge smearing
  temp values = T_v tau_v rng_v
    T_v = temperature of thermo-potentiostat
    tau_v = time constant of thermo-potentiostat
    rng_v = integer used to initialize random number generator

```

- zero or more keyword/value pairs may be appended
- keyword = *symm* or *couple* or *etypes* or *ffield* or *write\_mat* or *write\_inv* or *read\_mat* or *read\_inv*

```

symm value = on or off
  turn on/off charge neutrality constraint for the electrodes
couple values = group-ID val
  group-ID = group of atoms treated as additional electrode
  val = electric potential or charge on this electrode
etypes values = type
  type = atom type (can be a range) exclusive to the electrode for optimized neighbor_
↪lists
ffield value = on or off
  turn on/off finite-field implementation
write_mat value = filename
  filename = file to which to write elastance matrix
write_inv value = filename
  filename = file to which to write inverted matrix
read_mat value = filename
  filename = file from which to read elastance matrix
read_inv value = filename
  filename = file from which to read inverted matrix

```

## 17.50.2 Examples

```

fix fxconp bot electrode/conp -1.0 1.805 couple top 1.0 couple ref 0.0 write_inv inv.csv_
↪symm on
fix fxconp electrodes electrode/conq 0.0 1.805
fix fxconp bot electrode/thermo -1.0 1.805 temp 298 100 couple top 1.0

```

## 17.50.3 Description

fix *electrode/conp* mode implements a constant potential method (CPM) (*Siepmann, Reed*). Charges of groups specified via group-ID and optionally with the *couple* keyword are adapted to meet their respective potential at every time step. An arbitrary number of electrodes can be set but the respective groups may not overlap. Electrode charges have a Gaussian charge distribution with reciprocal width *eta*. The energy minimization is achieved via matrix inversion (*Wang*).

fix *electrode/conq* enforces a total charge specified in the input on each electrode. The energy is minimized w.r.t. the charge distribution within the electrode.

fix *electrode/thermo* implements a thermo-potentiostat (*Deissenbeck*). Temperature and time constant of the thermo-potentiostat need to be specified using the *temp* keyword. Currently, only two electrodes are possible with this style.

This fix necessitates the use of a long range solver that calculates and provides the matrix of electrode-electrode interactions and a vector of electrode-electrolyte interactions. The Kspace styles *ewald/electrode*, *pppm/electrode* and *pppm/electrode/intel* are created specifically for this task (*Ahrens-Iwers*).

For systems with non-periodic boundaries in one or two directions dipole corrections are available with the *kpace\_modify*. For *ewald/electrode* a two-dimensional Ewald summation (*Hu*) can be used by setting “slab ew2d”:

```
kspace_modify slab <slab_factor>
kspace_modify wire <wire_factor>
kspace_modify slab ew2d
```

Two implementations for the calculation of the elastance matrix are available with *pppm* and can be selected using the *amat onestep/twostep* keyword. *onestep* is the default; *twostep* can be faster for large electrodes and a moderate mesh size but requires more memory.

```
kspace_modify amat onestep/twostep
```

The *fix\_modify tf* option enables the Thomas-Fermi metallicity model (*Scalfi*) and allows parameters to be set for each atom type.

```
fix_modify ID tf type length voronoi
```

If this option is used parameters must be set for all atom types of the electrode.

The *fix\_modify timer* option turns on (off) additional timer outputs in the log file, for code developers to track optimization.

```
fix_modify ID timer on/off
```

The *fix\_modify set* options allow calculated quantities to be accessed via internal variables. Currently four types of quantities can be accessed:

```
fix_modify ID set v group-ID variablename
fix_modify ID set qsb group-ID variablename
fix_modify ID set mc group-ID1 group-ID2 variablename
fix_modify ID set me group-ID1 group-ID2 variablename
```

One use case is to output the potential that is internally calculated and applied to each electrode group by *fix electrode/conq* or *fix electrode/thermo*. For that case the *v* option makes *fix electrode* update the variable *variablename* with the potential applied to group *group-ID*, where *group-ID* must be a group whose charges are updated by *fix electrode* and *variablename* must be an internal-style variable:

```
fix conq bot electrode/conq -1.0 1.979 couple top 1.0
variable vbot internal 0.0
fix_modify conq set v bot vbot
```

The *qsb* option similarly outputs the total updated charge of the group if its potential were 0.0V. The *mc* option requires two *group-IDs*, and outputs the entry {*group-ID1*, *group-ID2*} of the (symmetric) *macro-capacitance* matrix (MC) which relates the electrodes’ applied potentials (V), total charges (Q), and total charges at 0.0 V (Qsb):

$$\mathbf{Q} = \mathbf{Q}_{SB} + \mathbf{MC} \cdot \mathbf{V}$$

Lastly, the *me* option also requires two *group-IDs* and outputs the entry {*group-ID1*, *group-ID2*} of the *macro-elastance* matrix, which is the inverse of the macro-capacitance matrix. (As the names denote, the macro-capacitance matrix gives electrode charges from potentials, and the macro-elastance matrix gives electrode potentials from charges).

**Warning:** Positions of electrode particles have to be immobilized at all times.

The parallelization for the `fix` works best if electrode atoms are evenly distributed across processors. For a system with two electrodes at the bottom and top of the cell this can be achieved with `processors * * 2`, or with the line

```
if "$(extract_setting(world_size) % 2) == 0" then "processors * * 2"
```

which avoids an error if the script is run on an odd number of processors (such as on just one processor for testing).

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix` *command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

**(Siepmann)** Siepmann and Sprik, J. Chem. Phys. 102, 511 (1995).

**(Reed)** Reed *et al.*, J. Chem. Phys. 126, 084704 (2007).

**(Wang)** Wang *et al.*, J. Chem. Phys. 141, 184102 (2014).

**(Deissenbeck)** Deissenbeck *et al.*, Phys. Rev. Letters 126, 136803 (2021).

**(Ahrens-Iwers)** Ahrens-Iwers and Meissner, J. Chem. Phys. 155, 104104 (2021).

**(Hu)** Hu, J. Chem. Theory Comput. 10, 5254 (2014).

**(Scalfi)** Scalfi *et al.*, J. Chem. Phys., 153, 174704 (2020).

## 17.51 fix electron/stopping command

## 17.52 fix electron/stopping/fit command

### 17.52.1 Syntax

```
fix ID group-ID style args
```

- ID, group-ID are documented in `fix` command
- style = `electron/stopping` or `electron/stopping/fit`

`electron/stopping` args = Ecut file keyword value ...

Ecut = minimum kinetic energy for electronic stopping (energy units)

file = name of the file containing the electronic stopping power table

`electron/stopping/fit` args = Ecut c1 c2 ...

`Ecut` = minimum kinetic energy for electronic stopping (energy units)  
`c1 c2` = linear and quadratic coefficients for the fitted quadratic polynomial

- zero or more keyword/value pairs may be appended to args for style = *electron/stopping*

keyword = *region* or *minneigh*  
*region* value = region-ID  
 region-ID = region whose atoms will be affected by this fix  
*minneigh* value = minneigh  
 minneigh = minimum number of neighbors an atom to have stopping applied

## 17.52.2 Examples

```
fix el all electron/stopping 10.0 elstop-table.txt
fix el all electron/stopping 10.0 elstop-table.txt minneigh 3
fix el mygroup electron/stopping 1.0 elstop-table.txt region bulk
fix 1 all electron/stopping/fit 4.63 3.3e-3 4.0e-8
fix 1 all electron/stopping/fit 3.49 1.8e-3 9.0e-8 7.57 4.2e-3 5.0e-8
```

## 17.52.3 Description

This fix implements inelastic energy loss for fast projectiles in solids. It applies a friction force to fast moving atoms to slow them down due to *electronic stopping* (energy lost via electronic collisions per unit of distance). This fix should be used for simulation of irradiation damage or ion implantation, where the ions can lose noticeable amounts of energy from electron excitations. If the electronic stopping power is not considered, the simulated range of the ions can be severely overestimated (*Nordlund98, Nordlund95*).

The electronic stopping is implemented by applying a friction force to each atom as:

$$\vec{F}_i = \vec{F}_i^0 - \frac{\vec{v}_i}{\|\vec{v}_i\|} \cdot S_e$$

where  $\vec{F}_i$  is the resulting total force on the atom.  $\vec{F}_i^0$  is the original force applied to the atom,  $\vec{v}_i$  is its velocity and  $S_e$  is the stopping power of the ion.

---

**Note:** In addition to electronic stopping, atomic cascades and irradiation simulations require the use of an adaptive timestep (see *fix dt/reset*) and the repulsive ZBL potential (see *ZBL* potential) or similar. Without these settings the interaction between the ion and the target atoms will be faulty. It is also common to use in such simulations a thermostat (*fix\_nvt*) in the borders of the simulation cell.

---

**Note:** This fix removes energy from fast projectiles without depositing it as a heat to the simulation cell. Such implementation might lead to the unphysical results when the amount of energy deposited to the electronic system is large, e.g. simulations of Swift Heavy Ions (energy per nucleon of 100 keV/amu or higher) or multiple projectiles. You could compensate energy loss by coupling bulk atoms with some thermostat or control heat transfer between electronic and atomic subsystems with the two-temperature model (*fix\_tm*).

---

At low velocities the electronic stopping is negligible. The electronic friction is not applied to atoms whose kinetic energy is smaller than *Ecut*, or smaller than the lowest energy value given in the table in *file*. Electronic stopping should be applied only when a projectile reaches bulk material. This fix scans neighbor list and excludes atoms with fewer than *minneigh* neighbors (by default one). If the pair potential cutoff is large, minneigh should be increased, though not

above the number of nearest neighbors in bulk material. An alternative is to disable the check for neighbors by setting *minneigh* to zero and using the *region* keyword. This is necessary when running simulations of cluster bombardment.

If the *region* keyword is used, the atom must also be in the specified geometric *region* in order to have electronic stopping applied to it. This is useful if the position of the bulk material is fixed. By default the electronic stopping is applied everywhere in the simulation cell.

---

The energy ranges and stopping powers are read from the file *file*. Lines starting with # and empty lines are ignored. Otherwise each line must contain exactly **N+1** numbers, where **N** is the number of atom types in the simulation.

The first column is the energy for which the stopping powers on that line apply. The energies must be sorted from the smallest to the largest. The other columns are the stopping powers  $S_e$  for each atom type, in ascending order, in force *units*. The stopping powers for intermediate energy values are calculated with linear interpolation between 2 nearest points.

For example:

```
# This is a comment
#      atom-1   atom-2
# eV   eV/Ang   eV/Ang # units metal
10      0        0
250     60       80
750    100      150
```

If an atom which would have electronic stopping applied to it has a kinetic energy higher than the largest energy given in *file*, LAMMPS will exit with an error message.

The stopping power depends on the energy of the ion and the target material. The electronic stopping table can be obtained from scientific publications, experimental databases or by using *SRIM* software. Other programs such as *CasP* or *PASS* can calculate the energy deposited as a function of the impact parameter of the ion; these results can be used to derive the stopping power.

---

Style *electron/stopping/fit* calculates the electronic stopping power and cumulative energy lost to the electron gas via a quadratic functional and applies a drag force to the classical equations-of-motion for all atoms moving above some minimum cutoff velocity (i.e., kinetic energy). These coefficients can be determined by fitting a quadratic polynomial to electronic stopping data predicted by, for example, SRIM or TD-DFT. Multiple ‘Ecut c1 c2’ values can be provided for multi-species simulations in the order of the atom types. There is an *examples/PACKAGES/electron\_stopping/* directory, which illustrates uses of this command. Details of this implementation are further described in *Stewart2018* and *Lee2020*.

#### 17.52.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify* options are not supported.

This fix computes a global scalar, which can be accessed by various *output commands*. The scalar is the total energy loss from electronic stopping applied by this fix since the start of the latest run. It is considered “intensive”.

The *start/stop* keywords of the *run* command have no effect on this fix.



## 17.52.5 Restrictions

This pair style is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

## 17.52.6 Default

The default is no limitation by region, and minneigh = 1.

**(electronic stopping)** Wikipedia - Electronic Stopping Power: [https://en.wikipedia.org/wiki/Stopping\\_power\\_%28particle\\_radiation%29](https://en.wikipedia.org/wiki/Stopping_power_%28particle_radiation%29)

**(Nordlund98)** Nordlund, Kai, et al. Physical Review B 57.13 (1998): 7556.

**(Nordlund95)** Nordlund, Kai. Computational materials science 3.4 (1995): 448-456.

**(SRIM)** SRIM webpage: <http://www.srim.org/>

**(CasP)** CasP webpage: [https://www.helmholtz-berlin.de/people/gregor-schiwietz/casp\\_en.html](https://www.helmholtz-berlin.de/people/gregor-schiwietz/casp_en.html)

**(PASS)** PASS webpage: <https://www.sdu.dk/en/DPASS>

**(Stewart2018)** J.A. Stewart, et al. (2018) Journal of Applied Physics, 123(16), 165902.

**(Lee2020)** C.W. Lee, et al. (2020) Physical Review B, 102(2), 024107.

## 17.53 fix enforce2d command

Accelerator Variants: *enforce2d/kk*

### 17.53.1 Syntax

```
fix ID group-ID enforce2d
```

- ID, group-ID are documented in *fix* command
- enforce2d = style name of this fix command

### 17.53.2 Examples

```
fix 5 all enforce2d
```

### 17.53.3 Description

Zero out the z-dimension velocity and force on each atom in the group. This is useful when running a 2d simulation to insure that atoms do not move from their initial z coordinate.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 17.53.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

### 17.53.5 Restrictions

none

### 17.53.6 Related commands

none

### 17.53.7 Default

none

## 17.54 fix eos/cv command

### 17.54.1 Syntax

```
fix ID group-ID eos/cv cv
```

- ID, group-ID are documented in *fix* command
- eos/cv = style name of this fix command
- cv = constant-volume heat capacity (energy/temperature units)

## 17.54.2 Examples

```
fix 1 all eos/cv 0.01
```

## 17.54.3 Description

Fix *eos/cv* applies a mesoparticle equation of state to relate the particle internal energy ( $u_i$ ) to the particle internal temperature ( $\theta_i$ ). The *eos/cv* mesoparticle equation of state requires the constant-volume heat capacity, and is defined as follows:

$$u_i = u_i^{mech} + u_i^{cond} = C_V \theta_i$$

where  $C_V$  is the constant-volume heat capacity,  $u^{cond}$  is the internal conductive energy, and  $u^{mech}$  is the internal mechanical energy. Note that alternative definitions of the mesoparticle equation of state are possible.

## 17.54.4 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This command also requires use of the *atom\_style dpd* command.

## 17.54.5 Related commands

*fix shardlow*, *pair dpd/fdt*

## 17.54.6 Default

none

**(Larentzos)** J.P. Larentzos, J.K. Brennan, J.D. Moore, and W.D. Mattson, “LAMMPS Implementation of Constant Energy Dissipative Particle Dynamics (DPD-E)”, ARL-TR-6863, U.S. Army Research Laboratory, Aberdeen Proving Ground, MD (2014).

## 17.55 fix eos/table command

### 17.55.1 Syntax

```
fix ID group-ID eos/table style file N keyword
```

- ID, group-ID are documented in *fix* command
- eos/table = style name of this fix command
- style = *linear* = method of interpolation
- file = filename containing the tabulated equation of state

- N = use N values in *linear* tables
- keyword = name of table keyword corresponding to table file

## 17.55.2 Examples

```
fix 1 all eos/table linear eos.table 100000 KEYWORD
```

## 17.55.3 Description

Fix *eos/table* applies a tabulated mesoparticle equation of state to relate the particle internal energy ( $u_i$ ) to the particle internal temperature ( $dpdTheta_i$ ).

Fix *eos/table* creates interpolation tables of length  $N$  from internal energy values listed in a file as a function of internal temperature.

The interpolation tables are created by fitting cubic splines to the file values and interpolating energy values at each of  $N$  internal temperatures, and vice versa. During a simulation, these tables are used to interpolate internal energy or temperature values as needed. The interpolation is done with the *linear* style.

For the *linear* style, the internal temperature is used to find 2 surrounding table values from which an internal energy is computed by linear interpolation, and vice versa.

The filename specifies a file containing tabulated internal temperature and internal energy values. The keyword specifies a section of the file. The format of this file is described below.

---

The format of a tabulated file is as follows (without the parenthesized comments):

```
# EOS TABLE          (one or more comment or blank lines)

KEYWORD              (keyword is first text on line)
N 500                (N parameter)
                    (blank)
1  1.00 0.000        (index, internal temperature, internal energy)
2  1.02 0.001
...
500 10.0 0.500
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The line can contain additional text, but the initial text must match the argument specified in the fix command.

The next line lists the number of table entries. The parameter “N” is required and its value is the number of table entries that follow. Note that this may be different than the  $N$  specified in the *fix eos/table* command. Let  $N_{table} = N$  in the fix command, and  $N_{file} = “N”$  in the tabulated file. What LAMMPS does is a preliminary interpolation by creating splines using the  $N_{file}$  tabulated values as nodal points. It uses these to interpolate as needed to generate energy and temperature values at  $N_{table}$  different points. The resulting tables of length  $N_{table}$  are then used as described above, when computing energy and temperature relationships. This means that if you want the interpolation tables of length  $N_{table}$  to match exactly what is in the tabulated file (with effectively no preliminary interpolation), you should set  $N_{table} = N_{file}$ .

Following a blank line, the next  $N$  lines list the tabulated values. On each line, the first value is the index from 1 to  $N$ , the second value is the internal temperature (in temperature units), the third value is the internal energy (in energy units).

Note that the internal temperature and internal energy values must increase from one line to the next.

Note that one file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified keyword.

### 17.55.4 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This command also requires use of the *atom\_style dpd* command.

The equation of state must be a monotonically increasing function.

An error will occur if the internal temperature or internal energies are not within the table cutoffs.

### 17.55.5 Related commands

*fix shardlow, pair dpd/fdt*

### 17.55.6 Default

none

## 17.56 fix eos/table/rx command

Accelerator Variants: *eos/table/rx/kk*

### 17.56.1 Syntax

```
fix ID group-ID eos/table/rx style file1 N keyword ...
```

- ID, group-ID are documented in *fix* command
- eos/table/rx = style name of this fix command
- style = *linear* = method of interpolation
- file1 = filename containing the tabulated equation of state
- N = use N values in *linear* tables
- keyword = name of table keyword corresponding to table file
- file2 = filename containing the heats of formation of each species (optional)
- deltaHf = heat of formation for a single species in energy units (optional)
- energyCorr = energy correction in energy units (optional)
- tempCorrCoeff = temperature correction coefficient (optional)

## 17.56.2 Examples

```
fix 1 all eos/table/rx linear eos.table 10000 KEYWORD thermo.table
fix 1 all eos/table/rx linear eos.table 10000 KEYWORD 1.5
fix 1 all eos/table/rx linear eos.table 10000 KEYWORD 1.5 0.025 0.0
```

## 17.56.3 Description

Fix *eos/table/rx* applies a tabulated mesoparticle equation of state to relate the concentration-dependent particle internal energy ( $u_i$ ) to the particle internal temperature ( $\theta_i$ ).

The concentration-dependent particle internal energy ( $u_i$ ) is computed according to the following relation:

$$U_i = \sum_{j=1}^m c_{i,j} (u_j + \Delta H_{f,j}) + \frac{3k_B T}{2} + N k_B T$$

where  $m$  is the number of species,  $c_{i,j}$  is the concentration of species  $j$  in particle  $i$ ,  $u_j$  is the internal energy of species  $j$ ,  $\Delta H_{f,j}$  is the heat of formation of species  $j$ ,  $N$  is the number of molecules represented by the coarse-grained particle,  $k_B$  is the Boltzmann constant, and  $T$  is the temperature of the system. Additionally, it is possible to modify the concentration-dependent particle internal energy relation by adding an energy correction, temperature-dependent correction, and/or a molecule-dependent correction. An energy correction can be specified as a constant (in energy units). A temperature correction can be specified by multiplying a temperature correction coefficient by the internal temperature. A molecular correction can be specified by multiplying a molecule correction coefficient by the average number of product gas particles in the coarse-grain particle.

Fix *eos/table/rx* creates interpolation tables of length  $N$  from  $m$  internal energy values of each species  $u_j$  listed in a file as a function of internal temperature. During a simulation, these tables are used to interpolate internal energy or temperature values as needed. The interpolation is done with the *linear* style. For the *linear* style, the internal temperature is used to find 2 surrounding table values from which an internal energy is computed by linear interpolation. A secant solver is used to determine the internal temperature from the internal energy.

The first filename specifies a file containing tabulated internal temperature and  $m$  internal energy values for each species  $u_j$ . The keyword specifies a section of the file. The format of this file is described below.

The second filename specifies a file containing heat of formation  $\Delta H_{f,j}$  for each species.

In cases where the coarse-grain particle represents a single molecular species (i.e., no reactions occur and fix *rx* is not present in the input file), fix *eos/table/rx* can be applied in a similar manner to fix *eos/table* within a non-reactive DPD simulation. In this case, the heat of formation filename is replaced with the heat of formation value for the single species. Additionally, the energy correction and temperature correction coefficients may also be specified as fix arguments.

The format of a tabulated file is as follows (without the parenthesized comments):

```
# EOS TABLE          (one or more comment or blank lines)

KEYWORD              (keyword is first text on line)
N 500 h2 no2 n2 ... no  (N parameter species1 species2 ... speciesN)
                       (blank)
1  1.00 0.000 ... 0.0000 (index, internal temperature, internal energy of species 1, ..
→ .., internal energy of species m)
2  1.02 0.001 ... 0.0002
...
500 10.0 0.500 ... 1.0000
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The line can contain additional text, but the initial text must match the argument specified in the `fix` command.

The next line lists the number of table entries and the species names that correspond with all the species listed in the reaction equations through the `fix rx` command. The parameter “N” is required and its value is the number of table entries that follow. Let  $N_{file} = “N”$  in the tabulated file. What LAMMPS does is a preliminary interpolation by creating splines using the  $N_{file}$  tabulated values as nodal points.

Following a blank line, the next N lines list the tabulated values. On each line, the first value is the index from 1 to N, the second value is the internal temperature (in temperature units), the third value until the  $m+3$  value are the internal energies of the m species (in energy units).

Note that all internal temperature and internal energy values must increase from one line to the next.

Note that one file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified keyword.

The format of a heat of formation file is as follows (without the parenthesized comments):

```
# HEAT OF FORMATION TABLE (one or more comment or blank lines)
                                (blank)
h2      0.00                    (species name, heat of formation)
no2     0.34
n2      0.00
...
no      0.93
```

Note that the species can be listed in any order. The tag that is used as the species name must correspond with the tags used to define the reactions with the `fix rx` command.

Alternatively, corrections to the EOS can be included by specifying three additional columns that correspond to the energy correction, the temperature correction coefficient and molecule correction coefficient. In this case, the format of the file is as follows:

```
# HEAT OF FORMATION TABLE (one or more comment or blank lines)
                                (blank)
h2      0.00 1.23 0.025 0.0 (species name, heat of formation, energy correction,
↪temperature correction coefficient, molecule correction coefficient)
no2     0.34 0.00 0.000 -1.76
n2      0.00 0.00 0.000 -1.76
...
no      0.93 0.00 0.000 -1.76
```

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix` *command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 17.56.4 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This command also requires use of the *atom\_style dpd* command.

The equation of state must be a monotonically increasing function.

An error will occur if the internal temperature or internal energies are not within the table cutoffs.

## 17.56.5 Related commands

*fix rx, pair dpd/fdt*

## 17.56.6 Default

none

# 17.57 fix evaporate command

## 17.57.1 Syntax

```
fix ID group-ID evaporate N M region-ID seed
```

- ID, group-ID are documented in *fix* command
- evaporate = style name of this fix command
- N = delete atoms every this many timesteps
- M = number of atoms to delete each time
- region-ID = ID of region within which to perform deletions
- seed = random number seed to use for choosing atoms to delete
- zero or more keyword/value pairs may be appended

keyword = *molecule*

*molecule* value = *no* or *yes*



## 17.57.2 Examples

```
fix 1 solvent evaporate 1000 10 surface 49892
fix 1 solvent evaporate 1000 10 surface 38277 molecule yes
```

## 17.57.3 Description

Remove  $M$  atoms from the simulation every  $N$  steps. This can be used, for example, to model evaporation of solvent particles or molecules (i.e. drying) of a system. Every  $N$  steps, the number of atoms in the fix group and within the specified region are counted.  $M$  of these are chosen at random and deleted. If there are less than  $M$  eligible particles, then all of them are deleted.

If the setting for the *molecule* keyword is *no*, then only single atoms are deleted. In this case, you should insure you do not delete only a portion of a molecule (only some of its atoms), or LAMMPS will soon generate an error when it tries to find those atoms. LAMMPS will warn you if any of the atoms eligible for deletion have a non-zero molecule ID, but does not check for this at the time of deletion.

If the setting for the *molecule* keyword is *yes*, then when an atom is chosen for deletion, the entire molecule it is part of is deleted. The count of deleted atoms is incremented by the number of atoms in the molecule, which may make it exceed  $M$ . If the molecule ID of the chosen atom is 0, then it is assumed to not be part of a molecule, and just the single atom is deleted.

As an example, if you wish to delete 10 water molecules every  $N$  steps, you should set  $M$  to 30. If only the water's oxygen atoms were in the fix group, then two hydrogen atoms would be deleted when an oxygen atom is selected for deletion, whether the hydrogen atoms are inside the evaporation region or not.

Note that neighbor lists are re-built on timesteps that atoms are removed. Thus you should not remove atoms too frequently or you will incur overhead due to the cost of building neighbor lists.

---

**Note:** If you are monitoring the temperature of a system where the atom count is changing due to evaporation, you typically should use the *compute\_modify dynamic yes* command for the temperature compute you are using.

---

## 17.57.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global scalar, which can be accessed by various *output commands*. The scalar is the cumulative number of deleted atoms. The scalar value calculated by this fix is “intensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.57.5 Restrictions

None

## 17.57.6 Related commands

*fix deposit*

## 17.57.7 Default

The option defaults are molecule = no.

## 17.58 fix external command

### 17.58.1 Syntax

```
fix ID group-ID external mode args
```

- ID, group-ID are documented in *fix* command
- external = style name of this fix command
- mode = *pf/callback* or *pf/array*

*pf/callback* args = Ncall Napply

Ncall = make callback every Ncall steps

Napply = apply callback forces every Napply steps

*pf/array* args = Napply

Napply = apply array forces every Napply steps

### 17.58.2 Examples

```
fix 1 all external pf/callback 1 1
fix 1 all external pf/callback 100 1
fix 1 all external pf/array 10
```

### 17.58.3 Description

This fix allows external programs that are running LAMMPS through its *library interface* to modify certain LAMMPS properties on specific timesteps, similar to the way other fixes do. The external driver can be a *C/C++ or Fortran program* or a *Python script*.

---

If mode is *pf/callback* then the fix will make a callback every *Ncall* timesteps or minimization iterations to the external program. The external program computes forces on atoms by setting values in an array owned by the fix. The fix then adds these forces to each atom in the group, once every *Napply* steps, similar to the way the *fix addforce* command works. Note that if *Ncall* > *Napply*, the force values produced by one callback will persist, and be used multiple times to update atom forces.

The callback function “foo” is invoked by the fix as:

```
foo(void *ptr, bigint timestep, int nlocal, tagint *ids, double **x, double **fexternal);
```

The arguments are as follows:

- *ptr* = pointer provided by and simply passed back to external driver
- *timestep* = current LAMMPS timestep
- *nlocal* = # of atoms on this processor
- *ids* = list of atom IDs on this processor
- *x* = coordinates of atoms on this processor
- *fexternal* = forces to add to atoms on this processor

Note that *timestep* is a “bigint” which is defined in `src/lmptype.h`, typically as a 64-bit integer. And *ids* is a pointer to type “tagint” which is typically a 32-bit integer unless LAMMPS is compiled with `-DLAMMPS_BIGBIG`. For more info please see the *build settings* section of the manual. Finally, *fexternal* are the forces returned by the driver program.

The fix has a `set_callback()` method which the external driver can call to pass a pointer to its `foo()` function. See the `couple/lammps_quest/lmpqst.cpp` file in the LAMMPS distribution for an example of how this is done. This sample application performs classical MD using quantum forces computed by a density functional code [Quest](#).

If mode is *pf/array* then the fix simply stores force values in an array. The fix adds these forces to each atom in the group, once every *Napply* steps, similar to the way the *fix addforce* command works.

The name of the public force array provided by the `FixExternal` class is

```
double **fexternal;
```

It is allocated by the `FixExternal` class as an (N,3) array where N is the number of atoms owned by a processor. The 3 corresponds to the *fx*, *fy*, *fz* components of force.

It is up to the external program to set the values in this array to the desired quantities, as often as desired. For example, the driver program might perform an MD run in stages of 1000 timesteps each. In between calls to the LAMMPS *run* command, it could retrieve atom coordinates from LAMMPS, compute forces, set values in *fexternal*, etc.

To use this fix during energy minimization, the energy corresponding to the added forces must also be set so as to be consistent with the added forces. Otherwise the minimization will not converge correctly. Correspondingly, the global virial needs to be updated to be use this fix with variable cell calculations (e.g. *fix box/relax* or *fix npt*).

This can be done from the external driver by calling these public methods of the `FixExternal` class:

```
void set_energy_global(double eng);
void set_virial_global(double *virial);
```

where *eng* is the potential energy, and *virial* an array of the 6 stress tensor components. *Eng* is an extensive quantity, meaning it should be the sum over per-atom energies of all affected atoms. It should also be provided in *energy units* consistent with the simulation. See the details below for how to insure this energy setting is used appropriately in a minimization.

Additional public methods that the caller can use to update system properties are:

```
void set_energy_peratom(double *eng);
void set_virial_peratom(double **virial);
void set_vector_length(int n);
void set_vector(int idx, double val);
```

These allow to set per-atom energy contributions, per-atom stress contributions, the length and individual values of a global vector of properties that the caller code may want to communicate to LAMMPS (e.g. for use in *fix ave/time* or in *equal-style variables* or for *custom thermo output*).

## 17.58.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy set by the external driver to both the global potential energy and peratom potential energies of the system as part of *thermodynamic output* or output by the *compute pe/atom* command. The default setting for this fix is *fix\_modify energy yes*. Note that this energy may be a fictitious quantity but it is needed so that the *minimize* command can include the forces added by this fix in a consistent manner. I.e. there is a decrease in potential energy when atoms move in the direction of the added force.

The *fix\_modify virial* option is supported by this fix to add the contribution computed by the external program to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial yes*.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the potential energy discussed above. The scalar stored by this fix is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the fictitious potential energy associated with the added forces to be included in the total potential energy of the system (the quantity being minimized), you **MUST** not disable the *fix\_modify energy* option for this fix.

---

## 17.58.5 Restrictions

none

## 17.58.6 Related commands

none

## 17.58.7 Default

none

## 17.59 fix ffl command

### 17.59.1 Syntax

<code>fix ID id-group ffl tau Tstart Tstop seed [flip-type]</code>
--

- ID, group-ID are documented in *fix* command
- ffl = style name of this fix command
- tau = thermostat parameter (positive real)

- Tstart, Tstop = temperature ramp during the run
- seed = random number seed to use for generating noise (positive integer)
- one more value may be appended

flip-type = determines the flipping type, can be chosen between rescale - no\_flip -  
 ↪ hard - soft, if no flip type is given, rescale will be chosen by default

## 17.59.2 Examples

```
fix 3 boundary ffl 10 300 300 31415
fix 1 all ffl 100 500 500 9265 soft
```

## 17.59.3 Description

Apply a Fast-Forward Langevin Equation (FFL) thermostat as described in (*Hijazi*). Contrary to *fix langevin*, this fix performs both thermostating and evolution of the Hamiltonian equations of motion, so it should not be used together with *fix nve* – at least not on the same atom groups.

The time-evolution of a single particle undergoing Langevin dynamics is described by the equations

$$\frac{dq}{dt} = \frac{p}{m},$$

$$\frac{dp}{dt} = -\gamma p + W + F,$$

where  $F$  is the physical force,  $\gamma$  is the friction coefficient, and  $W$  is a Gaussian random force.

The friction coefficient is the inverse of the thermostat parameter :  $\gamma = 1/\tau$ , with  $\tau$  the thermostat parameter *tau*. The thermostat parameter is given in the time units,  $\gamma$  is in inverse time units.

Equilibrium sampling a temperature  $T$  is obtained by specifying the target value as the *Tstart* and *Tstop* arguments, so that the internal constants depending on the temperature are computed automatically.

The random number *seed* must be a positive integer. A Marsaglia random number generator is used. Each processor uses the input seed to generate its own unique seed and its own stream of random numbers. Thus the dynamics of the system will not be identical on two runs on different numbers of processors.

The flipping type *flip-type* can be chosen between 4 types described in (*Hijazi*). The flipping operation occurs during the thermostating step and it flips the momenta of the atoms. If *no\_flip* is chosen, no flip will be executed and the integration will be the same as a standard Langevin thermostat (*Bussi*). The other flipping types are : rescale - hard - soft.

## 17.59.4 Restart, fix\_modify, output, run start/stop, minimize info

The instantaneous values of the extended variables are written to *binary restart files*. Because the state of the random number generator is not saved in restart files, this means you cannot do “exact” restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior. Note however that you should use a different seed each time you restart, otherwise the same sequence of random numbers will be used each time, which might lead to stochastic synchronization and subtle artifacts in the sampling.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

## 17.59.5 Restrictions

In order to perform constant-pressure simulations please use *fix press/berendsen*, rather than *fix npt*, to avoid duplicate integration of the equations of motion.

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 17.59.6 Related commands

*fix nvt*, *fix temp/rescale*, *fix viscous*, *fix nvt*, *pair\_style dpd/tstat*, *fix gld*, *fix gle*

---

**(Hijazi)** M. Hijazi, D. M. Wilkins, M. Ceriotti, J. Chem. Phys. 148, 184109 (2018)

**(Bussi)** G. Bussi, M. Parrinello, Phs. Rev. E 75, 056707 (2007)

## 17.60 fix filter/corotate command

### 17.60.1 Syntax

```
fix ID group-ID filter/corotate keyword value ...
```

- ID, group-ID are documented in *fix* command
- one or more constraint/value pairs are appended
- constraint = *b* or *a* or *t* or *m*

*b* values = one or more bond types  
*a* values = one or more angle types  
*t* values = one or more atom types  
*m* value = one or more mass values

### 17.60.2 Examples

```
timestep 8  
run_style respa 3 2 8 bond 1 pair 2 kspace 3  
fix cor all filter/corotate m 1.0  
  
fix cor all filter/corotate b 4 19 a 3 5 2
```

### 17.60.3 Description

This fix implements a corotational filter for a mollified impulse method. In biomolecular simulations, it allows the usage of larger timesteps for long-range electrostatic interactions. For details, see (*Fath*).

When using *run\_style respa* for a biomolecular simulation with high-frequency covalent bonds, the outer time-step is restricted to below  $\sim 4$ fs due to resonance problems. This fix filters the outer stage of the respa and thus a larger (outer) time-step can be used. Since in large biomolecular simulations the computation of the long-range electrostatic contributions poses a major bottleneck, this can significantly accelerate the simulation.

The filter computes a cluster decomposition of the molecular structure following the criteria indicated by the options a, b, t and m. This process is similar to the approach in *fix shake*, however, the clusters are not kept constrained. Instead, the position is slightly modified only for the computation of long-range forces. A good cluster decomposition constitutes in building clusters which contain the fastest covalent bonds inside clusters.

If the clusters are chosen suitably, the *run\_style respa* is stable for outer timesteps of at least 8fs.

### 17.60.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about these fixes is written to *binary restart files*. None of the *fix\_modify* options are relevant to these fixes. No global or per-atom quantities are stored by these fixes for access by various *output commands*. No parameter of these fixes can be used with the *start/stop* keywords of the *run* command. These fixes are not invoked during *energy minimization*.

### 17.60.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Currently, it does not support *molecule templates*.

### 17.60.6 Related commands

### 17.60.7 Default

none

---

(*Fath*) Fath, Hochbruck, Singh, J Comp Phys, 333, 180-198 (2017).

## 17.61 fix flow/gauss command

### 17.61.1 Syntax

```
fix ID group-ID flow/gauss xflag yflag zflag keyword
```

- ID, group-ID are documented in *fix* command
- flow/gauss = style name of this fix command

- xflag,yflag,zflag = 0 or 1

<pre>0 = do not conserve current in this dimension 1 = conserve current in this dimension</pre>
---

- zero or more keyword/value pairs may be appended
- keyword = *energy*  
  *energy* value = *no* or *yes*  
    *no* = do not compute work done by this fix  
    *yes* = compute work done by this fix

## 17.61.2 Examples

<pre>fix GD fluid flow/gauss 1 0 0 fix GD fluid flow/gauss 1 1 1 energy yes</pre>
---

## 17.61.3 Description

This fix implements the Gaussian dynamics (GD) method to simulate a system at constant mass flux (*Strong*). GD is a nonequilibrium molecular dynamics simulation method that can be used to study fluid flows through pores, pipes, and channels. In its original implementation GD was used to compute the pressure required to achieve a fixed mass flux through an opening. The flux can be conserved in any combination of the directions, x, y, or z, using xflag,yflag,zflag. This fix does not initialize a net flux through a system, it only conserves the center-of-mass momentum that is present when the fix is declared in the input script. Use the *velocity* command to generate an initial center-of-mass momentum.

GD applies an external fluctuating gravitational field that acts as a driving force to keep the system away from equilibrium. To maintain steady state, a profile-unbiased thermostat must be implemented to dissipate the heat that is added by the driving force. *Compute temp/profile* can be used to implement a profile-unbiased thermostat.

A common use of this fix is to compute a pressure drop across a pipe, pore, or membrane. The pressure profile can be computed in LAMMPS with *compute stress/atom* and *fix ave/chunk*, or with the hardy method in *fix atc*. Note that the simple *compute stress/atom* method is only accurate away from inhomogeneities in the fluid, such as fixed wall atoms. Further, the computed pressure profile must be corrected for the acceleration applied by GD before computing a pressure drop or comparing it to other methods, such as the pump method (*Zhu*). The pressure correction is discussed and described in (*Strong*).

For a complete example including the considerations discussed above, see the examples/PACKAGES/flow\_gauss directory.

---

**Note:** Only the flux of the atoms in group-ID will be conserved. If the velocities of the group-ID atoms are coupled to the velocities of other atoms in the simulation, the flux will not be conserved. For example, in a simulation with fluid atoms and harmonically constrained wall atoms, if a single thermostat is applied to group *all*, the fluid atom velocities will be coupled to the wall atom velocities, and the flux will not be conserved. This issue can be avoided by thermostatting the fluid and wall groups separately.

---

Adding an acceleration to atoms does work on the system. This added energy can be optionally subtracted from the potential energy for the thermodynamic output (see below) to check that the timestep is small enough to conserve energy. Since the applied acceleration is fluctuating in time, the work cannot be computed from a potential. As a result, computing the work is slightly more computationally expensive than usual, so it is not performed by default. To invoke the work calculation, use the *energy* keyword. The *fix\_modify energy* option also invokes the work calculation,



and overrides an *energy no* setting here. If neither *energy yes* or *fix\_modify energy yes* are set, the global scalar computed by the fix will return zero.

---

**Note:** In order to check energy conservation, any other fixes that do work on the system must have *fix\_modify energy yes* set as well. This includes thermostat fixes and any constraints that hold the positions of wall atoms fixed, such as *fix spring/self*.

---

If this fix is used in a simulation with the *rRESPA* integrator, the applied acceleration must be computed and applied at the same rRESPA level as the interactions between the flowing fluid and the obstacle. The rRESPA level at which the acceleration is applied can be changed using the *fix\_modify respa* option discussed below. If the flowing fluid and the obstacle interact through multiple interactions that are computed at different rRESPA levels, then there must be a separate flow/gauss fix for each level. For example, if the flowing fluid and obstacle interact through pairwise and long-range Coulomb interactions, which are computed at rRESPA levels 3 and 4, respectively, then there must be two separate flow/gauss fixes, one that specifies *fix\_modify respa 3* and one with *fix\_modify respa 4*.

---

#### 17.61.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy added by the fix to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify respa* option is supported by this fix. This allows the user to set at which level of the *rRESPA* integrator the fix computes and adds the external acceleration. Default is the outermost level.

This fix computes a global scalar and a global 3-vector of forces, which can be accessed by various *output commands*. The scalar is the negative of the work done on the system, see the discussion above. It is only calculated if the *energy* keyword is enabled or *fix\_modify energy yes* is set.

The vector is the total force that this fix applied to the group of atoms on the current timestep. The scalar and vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

This fix is not invoked during *energy minimization*.

#### 17.61.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 17.61.6 Related commands

*fix addforce*, *compute temp/profile*, *velocity*

## 17.61.7 Default

The option default for the *energy* keyword is `energy = no`.

---

**(Strong)** Strong and Eaves, J. Phys. Chem. B 121, 189 (2017).

**(Evans)** Evans and Morriss, Phys. Rev. Lett. 56, 2172 (1986).

**(Zhu)** Zhu, Tajkhorshid, and Schulten, Biophys. J. 83, 154 (2002).

## 17.62 fix freeze command

Accelerator Variants: *freeze/kk*

### 17.62.1 Syntax

```
fix ID group-ID freeze
```

- ID, group-ID are documented in *fix* command
- freeze = style name of this fix command

### 17.62.2 Examples

```
fix 2 bottom freeze
```

### 17.62.3 Description

Zero out the force and torque on a granular particle. This is useful for preventing certain particles from moving in a simulation. The *granular pair styles* also detect if this fix has been defined and compute interactions between frozen and non-frozen particles appropriately, as if the frozen particle has infinite mass. A similar functionality for normal (point) particles can be obtained using *fix setforce*.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 17.62.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global 3-vector of forces, which can be accessed by various *output commands*. This is the total force on the group of atoms before the forces on individual atoms are changed by the fix. The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.62.5 Restrictions

This fix is part of the GRANULAR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

There can only be a single freeze fix defined. This is because other the *granular pair styles* treat frozen particles differently and need to be able to reference a single group to which this fix is applied.

### 17.62.6 Related commands

*atom\_style sphere, fix setforce*

### 17.62.7 Default

none

## 17.63 fix gcmc command

### 17.63.1 Syntax

```
fix ID group-ID gcmc N X M type seed T mu displace keyword values ...
```

- ID, group-ID are documented in *fix* command
- gcmc = style name of this fix command
- N = invoke this fix every N steps
- X = average number of GCMC exchanges to attempt every N steps
- M = average number of MC moves to attempt every N steps
- type = atom type for inserted atoms (must be 0 if mol keyword used)
- seed = random # seed (positive integer)
- T = temperature of the ideal gas reservoir (temperature units)
- mu = chemical potential of the ideal gas reservoir (energy units)
- displace = maximum Monte Carlo translation distance (length units)
- zero or more keyword/value pairs may be appended to args

keyword = *mol*, *region*, *maxangle*, *pressure*, *fugacity\_coeff*, *full\_energy*, *charge*,  
↪*group*, *grouptype*, *intra\_energy*, *tfac\_insert*, or *overlap\_cutoff*

*mol* value = template-ID  
template-ID = ID of molecule template specified in a separate *molecule* command

*mcmoves* values = Patomtrans Pmoltrans Pmolrotate  
Patomtrans = proportion of atom translation MC moves  
Pmoltrans = proportion of molecule translation MC moves  
Pmolrotate = proportion of molecule rotation MC moves

*rigid* value = fix-ID  
fix-ID = ID of *fix rigid/small* command

*shake* value = fix-ID  
fix-ID = ID of *fix shake* command

*region* value = region-ID  
region-ID = ID of region where GCMC exchanges and MC moves are allowed

*maxangle* value = maximum molecular rotation angle (degrees)  
*pressure* value = pressure of the gas reservoir (pressure units)  
*fugacity\_coeff* value = fugacity coefficient of the gas reservoir (unitless)  
*full\_energy* = compute the entire system energy when performing GCMC exchanges and  
↪MC moves

*charge* value = charge of inserted atoms (charge units)  
*group* value = group-ID  
group-ID = group-ID for inserted atoms (string)

*grouptype* values = type group-ID  
type = atom type (int)  
group-ID = group-ID for inserted atoms (string)

*intra\_energy* value = intramolecular energy (energy units)  
*tfac\_insert* value = scale up/down temperature of inserted atoms (unitless)  
*overlap\_cutoff* value = maximum pair distance for overlap rejection (distance,  
↪units)

*max* value = Maximum number of molecules allowed in the system  
*min* value = Minimum number of molecules allowed in the system

## 17.63.2 Examples

```
fix 2 gas gcmc 10 1000 1000 2 29494 298.0 -0.5 0.01
fix 3 water gcmc 10 100 100 0 3456543 3.0 -2.5 0.1 mol my_one_water maxangle 180 full_
↪energy
fix 4 my_gas gcmc 1 10 10 1 123456543 300.0 -12.5 1.0 region disk
```

## 17.63.3 Description

This fix performs grand canonical Monte Carlo (GCMC) exchanges of atoms or molecules with an imaginary ideal gas reservoir at the specified T and chemical potential ( $\mu$ ) as discussed in (*Frenkel*). It also attempts Monte Carlo (MC) moves (translations and molecule rotations) within the simulation cell or region. If used with the *fix nvt* command, simulations in the grand canonical ensemble ( $\mu$ VT, constant chemical potential, constant volume, and constant temperature) can be performed. Specific uses include computing isotherms in microporous materials, or computing vapor-liquid coexistence curves.

Every N timesteps the fix attempts both GCMC exchanges (insertions or deletions) and MC moves of gas atoms or molecules. On those timesteps, the average number of attempted GCMC exchanges is X, while the average number of attempted MC moves is M. For GCMC exchanges of either molecular or atomic gasses, these exchanges can be either deletions or insertions, with equal probability.

The possible choices for MC moves are translation of an atom, translation of a molecule, and rotation of a molecule. The relative amounts of each are determined by the optional *mcmoves* keyword (see below). The default behavior is as follows. If the *mol* keyword is used, only molecule translations and molecule rotations are performed with equal probability. Conversely, if the *mol* keyword is not used, only atom translations are performed. *M* should typically be chosen to be approximately equal to the expected number of gas atoms or molecules of the given type within the simulation cell or region, which will result in roughly one MC move per atom or molecule per MC cycle.

All inserted particles are always added to two groups: the default group “all” and the fix group specified in the fix command. In addition, particles are also added to any groups specified by the *group* and *groupstype* keywords. If inserted particles are individual atoms, they are assigned the atom type given by the *type* argument. If they are molecules, the *type* argument has no effect and must be set to zero. Instead, the type of each atom in the inserted molecule is specified in the file read by the *molecule* command.

---

**Note:** Care should be taken to apply fix gcmc only to a group that contains only those atoms and molecules that you wish to manipulate using Monte Carlo. Hence it is generally not a good idea to specify the default group “all” in the fix command, although it is allowed.

---

This fix cannot be used to perform GCMC insertions of gas atoms or molecules other than the exchanged type, but GCMC deletions, and MC translations, and rotations can be performed on any atom/molecule in the fix group. All atoms in the simulation cell can be moved using regular time integration translations, e.g. via *fix nvt*, resulting in a hybrid GCMC+MD simulation. A smaller-than-usual timestep size may be needed when running such a hybrid simulation, especially if the inserted molecules are not well equilibrated.

This command may optionally use the *region* keyword to define an exchange and move volume. The specified region must have been previously defined with a *region* command. It must be defined with *side = in*. Insertion attempts occur only within the specified region. For non-rectangular regions, random trial points are generated within the rectangular bounding box until a point is found that lies inside the region. If no valid point is generated after 1000 trials, no insertion is performed, but it is counted as an attempted insertion. Move and deletion attempt candidates are selected from gas atoms or molecules within the region. If there are no candidates, no move or deletion is performed, but it is counted as an attempt move or deletion. If an attempted move places the atom or molecule center-of-mass outside the specified region, a new attempted move is generated. This process is repeated until the atom or molecule center-of-mass is inside the specified region.

If used with *fix nvt*, the temperature of the imaginary reservoir, *T*, should be set to be equivalent to the target temperature used in *fix nvt*. Otherwise, the imaginary reservoir will not be in thermal equilibrium with the simulation cell. Also, it is important that the temperature used by *fix nvt* be dynamic/dof, which can be achieved as follows:

```
compute mdtemp mdatoms temp
compute_modify mdtemp dynamic/dof yes
fix mdnvt mdatoms nvt temp 300.0 300.0 10.0
fix_modify mdnvt temp mdtemp
```

Note that neighbor lists are re-built every timestep that this fix is invoked, so you should not set *N* to be too small. However, periodic rebuilds are necessary in order to avoid dangerous rebuilds and missed interactions. Specifically, avoid performing so many MC translations per timestep that atoms can move beyond the neighbor list skin distance. See the *neighbor* command for details.

When an atom or molecule is to be inserted, its coordinates are chosen at a random position within the current simulation cell or region, and new atom velocities are randomly chosen from the specified temperature distribution given by *T*. The effective temperature for new atom velocities can be increased or decreased using the optional keyword *tfac\_insert* (see below). Relative coordinates for atoms in a molecule are taken from the template molecule provided by the user. The center of mass of the molecule is placed at the insertion point. The orientation of the molecule is chosen at random by rotating about this point.

Individual atoms are inserted, unless the *mol* keyword is used. It specifies a *template-ID* previously defined using the *molecule* command, which reads a file that defines the molecule. The coordinates, atom types, charges, etc., as well as

any bonding and special neighbor information for the molecule can be specified in the molecule file. See the *molecule* command for details. The only settings required to be in this file are the coordinates and types of atoms in the molecule.

When not using the *mol* keyword, you should ensure you do not delete atoms that are bonded to other atoms, or LAMMPS will soon generate an error when it tries to find bonded neighbors. LAMMPS will warn you if any of the atoms eligible for deletion have a non-zero molecule ID, but does not check for this at the time of deletion.

If you wish to insert molecules using the *mol* keyword that will be treated as rigid bodies, use the *rigid* keyword, specifying as its value the ID of a separate *fix rigid/small* command which also appears in your input script.

---

**Note:** If you wish the new rigid molecules (and other rigid molecules) to be thermostatted correctly via *fix rigid/small/nvt* or *fix rigid/small/npt*, then you need to use the “fix\_modify dynamic/dof yes” command for the rigid fix. This is to inform that fix that the molecule count will vary dynamically.

---

If you wish to insert molecules via the *mol* keyword, that will have their bonds or angles constrained via SHAKE, use the *shake* keyword, specifying as its value the ID of a separate *fix shake* command which also appears in your input script.

Optionally, users may specify the relative amounts of different MC moves using the *mcmoves* keyword. The values *Patomtrans*, *Pmoltrans*, *Pmolrotate* specify the average proportion of atom translations, molecule translations, and molecule rotations, respectively. The values must be non-negative integers or real numbers, with at least one non-zero value. For example, (10,30,0) would result in 25% of the MC moves being atomic translations, 75% molecular translations, and no molecular rotations.

Optionally, users may specify the maximum rotation angle for molecular rotations using the *maxangle* keyword and specifying the angle in degrees. Rotations are performed by generating a random point on the unit sphere and a random rotation angle on the range [0,maxangle). The molecule is then rotated by that angle about an axis passing through the molecule center of mass. The axis is parallel to the unit vector defined by the point on the unit sphere. The same procedure is used for randomly rotating molecules when they are inserted, except that the maximum angle is 360 degrees.

Note that fix gcmc does not use configurational bias MC or any other kind of sampling of intramolecular degrees of freedom. Inserted molecules can have different orientations, but they will all have the same intramolecular configuration, which was specified in the molecule command input.

For atomic gasses, inserted atoms have the specified atom type, but deleted atoms are any atoms that have been inserted or that already belong to the fix group. For molecular gasses, exchanged molecules use the same atom types as in the template molecule supplied by the user. In both cases, exchanged atoms/molecules are assigned to two groups: the default group “all” and the fix group (which can also be “all”).

The chemical potential is a user-specified input parameter defined as:

$$\mu = \mu^{id} + \mu^{ex}$$

The second term  $\mu_{ex}$  is the excess chemical potential due to energetic interactions and is formally zero for the fictitious gas reservoir but is non-zero for interacting systems. So, while the chemical potential of the reservoir and the simulation cell are equal,  $\mu_{ex}$  is not, and as a result, the densities of the two are generally quite different. The first term  $\mu_{id}$  is the ideal gas contribution to the chemical potential.  $\mu_{id}$  can be related to the density or pressure of the fictitious gas reservoir by:

$$\begin{aligned}\mu^{id} &= kT \ln \rho \Lambda^3 \\ &= kT \ln \frac{\phi P \Lambda^3}{k_B T}\end{aligned}$$

where  $k_B$  is the Boltzmann constant,  $T$  is the user-specified temperature,  $\rho$  is the number density,  $P$  is the pressure, and  $\phi$  is the fugacity coefficient. The constant  $\Lambda$  is required for dimensional consistency. For all unit styles except *lj* it

is defined as the thermal de Broglie wavelength

$$\Lambda = \sqrt{\frac{h^2}{2\pi m k_B T}}$$

where  $h$  is Planck's constant, and  $m$  is the mass of the exchanged atom or molecule. For unit style *lj*,  $\Lambda$  is simply set to unity. Note that prior to March 2017,  $\Lambda$  for unit style *lj* was calculated using the above formula with  $h$  set to the rather specific value of 0.18292026. Chemical potential under the old definition can be converted to an equivalent value under the new definition by subtracting  $3kT \ln(\Lambda_{old})$ .

As an alternative to specifying  $\mu$  directly, the ideal gas reservoir can be defined by its pressure  $P$  using the *pressure* keyword, in which case the user-specified chemical potential is ignored. The user may also specify the fugacity coefficient  $\phi$  using the *fugacity\_coeff* keyword, which defaults to unity.

The *full\_energy* option means that the fix calculates the total potential energy of the entire simulated system, instead of just the energy of the part that is changed. The total system energy before and after the proposed GCMC exchange or MC move is then used in the Metropolis criterion to determine whether or not to accept the proposed change. By default, this option is off, in which case only partial energies are computed to determine the energy difference due to the proposed change.

The *full\_energy* option is needed for systems with complicated potential energy calculations, including the following:

- long-range electrostatics (kspace)
- many-body pair styles
- hybrid pair styles
- eam pair styles
- tail corrections
- need to include potential energy contributions from other fixes

In these cases, LAMMPS will automatically apply the *full\_energy* keyword and issue a warning message.

When the *mol* keyword is used, the *full\_energy* option also includes the intramolecular energy of inserted and deleted molecules, whereas this energy is not included when *full\_energy* is not used. If this is not desired, the *intra\_energy* keyword can be used to define an amount of energy that is subtracted from the final energy when a molecule is inserted, and subtracted from the initial energy when a molecule is deleted. For molecules that have a non-zero intramolecular energy, this will ensure roughly the same behavior whether or not the *full\_energy* option is used.

Inserted atoms and molecules are assigned random velocities based on the specified temperature  $T$ . Because the relative velocity of all atoms in the molecule is zero, this may result in inserted molecules that are systematically too cold. In addition, the intramolecular potential energy of the inserted molecule may cause the kinetic energy of the molecule to quickly increase or decrease after insertion. The *tfac\_insert* keyword allows the user to counteract these effects by changing the temperature used to assign velocities to inserted atoms and molecules by a constant factor. For a particular application, some experimentation may be required to find a value of *tfac\_insert* that results in inserted molecules that equilibrate quickly to the correct temperature.

Some fixes have an associated potential energy. Examples of such fixes include: *efield*, *gravity*, *addforce*, *langevin*, *restrain*, *temp/berendsen*, *temp/rescale*, and *wall fixes*. For that energy to be included in the total potential energy of the system (the quantity used when performing GCMC exchange and MC moves), you MUST enable the *fix\_modify energy* option for that fix. The doc pages for individual *fix* commands specify if this should be done.

Use the *charge* option to insert atoms with a user-specified point charge. Note that doing so will cause the system to become non-neutral. LAMMPS issues a warning when using long-range electrostatics (kspace) with non-neutral systems. See the *compute group/group* documentation for more details about simulating non-neutral systems with kspace on.

Use of this fix typically will cause the number of atoms to fluctuate, therefore, you will want to use the `compute_modify dynamic/dof` command to insure that the current number of atoms is used as a normalizing factor each time temperature is computed. A simple example of this is:

```
compute_modify thermo_temp dynamic yes
```

A more complicated example is listed earlier on this page in the context of NVT dynamics.

---

**Note:** If the density of the cell is initially very small or zero, and increases to a much larger density after a period of equilibration, then certain quantities that are only calculated once at the start (kspace parameters) may no longer be accurate. The solution is to start a new simulation after the equilibrium density has been reached.

---

With some pair\_styles, such as *Buckingham*, *Born-Mayer-Huggins* and *ReaxFF*, two atoms placed close to each other may have an arbitrary large, negative potential energy due to the functional form of the potential. While these unphysical configurations are inaccessible to typical dynamical trajectories, they can be generated by Monte Carlo moves. The `overlap_cutoff` keyword suppresses these moves by effectively assigning an infinite positive energy to all new configurations that place any pair of atoms closer than the specified overlap cutoff distance.

The `max` and `min` keywords allow for the restriction of the number of atoms in the simulation. They automatically reject all insertion or deletion moves that would take the system beyond the set boundaries. Should the system already be beyond the boundary, only moves that bring the system closer to the bounds may be accepted.

The `group` keyword adds all inserted atoms to the `group` of the group-ID value. The `group_type` keyword adds all inserted atoms of the specified type to the `group` of the group-ID value.

#### 17.63.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the fix to *binary restart files*. This includes information about the random number generator seed, the next timestep for MC exchanges, the number of MC step attempts and successes etc. See the `read_restart` command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

---

**Note:** For this to work correctly, the timestep must **not** be changed after reading the restart with `reset_timestep`. The fix will try to detect it and stop with an error.

---

None of the `fix_modify` options are relevant to this fix.

This fix computes a global vector of length 8, which can be accessed by various *output commands*. The vector values are the following global cumulative quantities:

- 1 = translation attempts
- 2 = translation successes
- 3 = insertion attempts
- 4 = insertion successes
- 5 = deletion attempts
- 6 = deletion successes
- 7 = rotation attempts
- 8 = rotation successes



The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.63.5 Restrictions

This fix is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

Do not set “neigh\_modify once yes” or else this fix will never be called. RENEIGHBORING is required.

Only usable for 3D simulations.

Can be run in parallel, but aspects of the GCMC part will not scale well in parallel. Currently, molecule translations and rotations are not supported with more than one MPI process. It is still possible to do parallel molecule exchange without translation and rotation moves by setting MC moves to zero and/or by using the *mcmoves* keyword with *Pmoltrans* = *Pmolrotate* = 0 .

When using fix *gcmc* in combination with fix *shake* or fix *rigid*, only GCMC exchange moves are supported, so the argument *M* must be zero.

When using fix *gcmc* in combination with fix *rigid*, deletion of the last remaining molecule is not allowed for technical reasons, and so the molecule count will never drop below 1, regardless of the specified chemical potential.

Note that very lengthy simulations involving insertions/deletions of billions of gas molecules may run out of atom or molecule IDs and trigger an error, so it is better to run multiple shorter-duration simulations. Likewise, very large molecules have not been tested and may turn out to be problematic.

Use of multiple fix *gcmc* commands in the same input script can be problematic if using a template molecule. The issue is that the user-referenced template molecule in the second fix *gcmc* command may no longer exist since it might have been deleted by the first fix *gcmc* command. An existing template molecule will need to be referenced by the user for each subsequent fix *gcmc* command.

### 17.63.6 Related commands

*fix atom/swap*, *fix nvt*, *neighbor*, *fix deposit*, *fix evaporate*, *delete\_atoms*

### 17.63.7 Default

The option defaults are *mol* = no, *maxangle* = 10, *overlap\_cutoff* = 0.0, *fugacity\_coeff* = 1.0, *intra\_energy* = 0.0, *tfac\_insert* = 1.0. (*Patomtrans*, *Pmoltrans*, *Pmolrotate*) = (1, 0, 0) for *mol* = no and (0, 1, 1) for *mol* = yes. *full\_energy* = no, except for the situations where *full\_energy* is required, as listed above.

---

(Frenkel) Frenkel and Smit, Understanding Molecular Simulation, Academic Press, London, 2002.

## 17.64 fix gld command

### 17.64.1 Syntax

```
fix ID group-ID gld Tstart Tstop N_k seed series c_1 tau_1 ... c_N_k tau_N_k keyword_
↪values ...
```

- ID, group-ID are documented in *fix* command
- gld = style name of this fix command
- Tstart,Tstop = desired temperature at start/end of run (temperature units)
- N\_k = number of terms in the Prony series representation of the memory kernel
- seed = random number seed to use for white noise (positive integer)
- series = *pprony* is presently the only available option
- c\_k = the weight of the kth term in the Prony series (mass per time units)
- tau\_k = the time constant of the kth term in the Prony series (time units)
- zero or more keyword/value pairs may be appended

keyword = *frozen* or *zero*

*frozen* value = *no* or *yes*

*no* = initialize extended variables using values drawn from equilibrium\_  
↪distribution at Tstart

*yes* = initialize extended variables to zero (i.e., from equilibrium\_  
↪distribution at zero temperature)

*zero* value = *no* or *yes*

*no* = do not set total random force to zero

*yes* = set total random force to zero

### 17.64.2 Examples

```
fix 1 all gld 1.0 1.0 2 82885 pprony 0.5 1.0 1.0 2.0 frozen yes zero yes
fix 3 rouse gld 7.355 7.355 4 48823 pprony 107.1 0.02415 186.0 0.04294 428.6 0.09661_
↪1714 0.38643
```

### 17.64.3 Description

Applies Generalized Langevin Dynamics to a group of atoms, as described in (*Baczewski*). This is intended to model the effect of an implicit solvent with a temporally non-local dissipative force and a colored Gaussian random force, consistent with the Fluctuation-Dissipation Theorem. The functional form of the memory kernel associated with the temporally non-local force is constrained to be a Prony series.

---

**Note:** While this fix bears many similarities to *fix langevin*, it has one significant difference. Namely, *fix gld* performs time integration, whereas *fix langevin* does NOT. To this end, the specification of another fix to perform time integration, such as *fix nve*, is NOT necessary.

---

With this fix active, the force on the  $j$ th atom is given as

$$\mathbf{F}_j(t) = \mathbf{F}_j^C(t) - \int_0^t \Gamma_j(t-s) \mathbf{v}_j(s) ds + \mathbf{F}_j^R(t)$$

$$\Gamma_j(t-s) = \sum_{k=1}^{N_k} \frac{c_k}{\tau_k} e^{-(t-s)/\tau_k}$$

$$\langle \mathbf{F}_j^R(t), \mathbf{F}_j^R(s) \rangle = k_B T \Gamma_j(t-s)$$

Here, the first term is representative of all conservative (pairwise, bonded, etc) forces external to this fix, the second is the temporally non-local dissipative force given as a Prony series, and the third is the colored Gaussian random force.

The Prony series form of the memory kernel is chosen to enable an extended variable formalism, with a number of exemplary mathematical features discussed in (*Baczewski*). In particular,  $3N_k$  extended variables are added to each atom, which effect the action of the memory kernel without having to explicitly evaluate the integral over time in the second term of the force. This also has the benefit of requiring the generation of uncorrelated random forces, rather than correlated random forces as specified in the third term of the force.

Presently, the Prony series coefficients are limited to being greater than or equal to zero, and the time constants are limited to being greater than zero. To this end, the value of series MUST be set to *pprony*, for now. Future updates will allow for negative coefficients and other representations of the memory kernel. It is with these updates in mind that the series option was included.

The units of the Prony series coefficients are chosen to be mass per time to ensure that the numerical integration scheme stably approaches the Newtonian and Langevin limits. Details of these limits, and the associated numerical concerns are discussed in (*Baczewski*).

The desired temperature at each timestep is ramped from *Tstart* to *Tstop* over the course of the next run.

The random # *seed* must be a positive integer. A Marsaglia random number generator is used. Each processor uses the input seed to generate its own unique seed and its own stream of random numbers. Thus the dynamics of the system will not be identical on two runs on different numbers of processors.

The keyword/value option pairs are used in the following ways.

The keyword *frozen* can be used to specify how the extended variables associated with the GLD memory kernel are initialized. Specifying no (the default), the initial values are drawn at random from an equilibrium distribution at *Tstart*, consistent with the Fluctuation-Dissipation Theorem. Specifying yes, initializes the extended variables to zero.

The keyword *zero* can be used to eliminate drift due to the thermostat. Because the random forces on different atoms are independent, they do not sum exactly to zero. As a result, this fix applies a small random force to the entire system, and the center-of-mass of the system undergoes a slow random walk. If the keyword *zero* is set to *yes*, the total random force is set exactly to zero by subtracting off an equal part of it from each atom in the group. As a result, the center-of-mass of a system with zero initial momentum will not drift over time.

### 17.64.4 Restart, *fix\_modify*, output, run start/stop, minimize info

The instantaneous values of the extended variables are written to *binary restart files*. Because the state of the random number generator is not saved in restart files, this means you cannot do “exact” restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior.

None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

### 17.64.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.64.6 Related commands

*fix langevin*, *fix viscous*, *pair\_style dpd/tstat*

### 17.64.7 Default

The option defaults are frozen = no, zero = no.

---

(Baczewski) A.D. Baczewski and S.D. Bond, J. Chem. Phys. 139, 044107 (2013).

## 17.65 *fix gle* command

### 17.65.1 Syntax

```
fix ID id-group gle Ns Tstart Tstop seed Amatrix [noneq Cmatrix] [every stride]
```

- ID, group-ID are documented in *fix* command
- gle = style name of this fix command
- Ns = number of additional fictitious momenta
- Tstart, Tstop = temperature ramp during the run
- Amatrix = file to read the drift matrix A from
- seed = random number seed to use for generating noise (positive integer)
- zero or more keyword/value pairs may be appended

keyword = *noneq* or *every*  
*noneq* Cmatrix = file to read the non-equilibrium covariance matrix from  
*every* stride = apply the GLE once every time steps. Reduces the accuracy  
of the integration of the GLE, but has \*no effect\* on the accuracy of  $\int_{\text{equilibrium}}$   
sampling. It might change sampling properties when used together with *noneq*.

## 17.65.2 Examples

```
fix 3 boundary gle 6 300 300 31415 smart.A
fix 1 all gle 6 300 300 31415 qt-300k.A noneq qt-300k.C
```

## 17.65.3 Description

Apply a Generalized Langevin Equation (GLE) thermostat as described in (*Cerioti*). The formalism allows one to obtain a number of different effects ranging from efficient sampling of all vibrational modes in the system to inexpensive (approximate) modelling of nuclear quantum effects. Contrary to *fix langevin*, this fix performs both thermostating and evolution of the Hamiltonian equations of motion, so it should not be used together with *fix nve* – at least not on the same atom groups.

Each degree of freedom in the thermostatted group is supplemented with  $N_s$  additional degrees of freedom  $s$ , and the equations of motion become

$$\begin{aligned} dq/dt &= p/m \\ d(p,s)/dt &= (F, \emptyset) - A(p,s) + B dW/dt \end{aligned}$$

where  $F$  is the physical force,  $A$  is the drift matrix (that generalizes the friction in Langevin dynamics),  $B$  is the diffusion term and  $dW/dt$  un-correlated Gaussian random forces. The  $A$  matrix couples the physical  $(q,p)$  dynamics with that of the additional degrees of freedom, and makes it possible to obtain effectively a history-dependent noise and friction kernel.

The drift matrix should be given as an external file *Afile*, as a  $(N_s+1 \times N_s+1)$  matrix in inverse time units. Matrices that are optimal for a given application and the system of choice can be obtained from (*GLE4MD*).

Equilibrium sampling a temperature  $T$  is obtained by specifying the target value as the *Tstart* and *Tstop* arguments, so that the diffusion matrix that gives canonical sampling for a given  $A$  is computed automatically. However, the GLE framework also allow for non-equilibrium sampling, that can be used for instance to model inexpensively zero-point energy effects (*Cerioti2*). This is achieved specifying the *noneq* keyword followed by the name of the file that contains the static covariance matrix for the non-equilibrium dynamics. Please note, that the covariance matrix is expected to be given in **temperature units**.

Since integrating GLE dynamics can be costly when used together with simple potentials, one can use the *every* optional keyword to apply the Langevin terms only once every several MD steps, in a multiple time-step fashion. This should be used with care when doing non-equilibrium sampling, but should have no effect on equilibrium averages when using canonical sampling.

The random number *seed* must be a positive integer. A Marsaglia random number generator is used. Each processor uses the input seed to generate its own unique seed and its own stream of random numbers. Thus the dynamics of the system will not be identical on two runs on different numbers of processors.

Note also that the Generalized Langevin Dynamics scheme that is implemented by the *fix gld* scheme is closely related to the present one. In fact, it should be always possible to cast the Prony series form of the memory kernel used by GLD into an appropriate input matrix for *fix gle*. While the GLE scheme is more general, the form used by *fix gld* can be more directly related to the representation of an implicit solvent environment.

### 17.65.4 Restart, fix\_modify, output, run start/stop, minimize info

The instantaneous values of the extended variables are written to *binary restart files*. Because the state of the random number generator is not saved in restart files, this means you cannot do “exact” restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior. Note however that you should use a different seed each time you restart, otherwise the same sequence of random numbers will be used each time, which might lead to stochastic synchronization and subtle artifacts in the sampling.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

### 17.65.5 Restrictions

The GLE thermostat in its current implementation should not be used with rigid bodies, SHAKE or RATTLE. It is expected that all the thermostatted degrees of freedom are fully flexible, and the sampled ensemble will not be correct otherwise.

In order to perform constant-pressure simulations please use *fix press/berendsen*, rather than *fix npt*, to avoid duplicate integration of the equations of motion.

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.65.6 Related commands

*fix nvt*, *fix temp/rescale*, *fix viscous*, *fix nvt*, *pair\_style dpd/tstat*, *fix gl*

---

**(Ceriotti)** Ceriotti, Bussi and Parrinello, J Chem Theory Comput 6, 1170-80 (2010)

**(GLE4MD)** <http://gle4md.org/>

**(Ceriotti2)** Ceriotti, Bussi and Parrinello, Phys Rev Lett 103, 030603 (2009)

## 17.66 fix gravity command

Accelerator Variants: *gravity/omp*, *gravity/kk*

## 17.66.1 Syntax

```
fix ID group gravity magnitude style args
```

- ID, group are documented in *fix* command
- gravity = style name of this fix command
- magnitude = size of acceleration (force/mass units)
- magnitude can be a variable (see below)
- style = *chute* or *spherical* or *gradient* or *vector*

*chute* args = angle

angle = angle in +x away from -z or -y axis in 3d/2d (in degrees)

angle can be a variable (see below)

*spherical* args = phi theta

phi = azimuthal angle from +x axis (in degrees)

theta = angle from +z or +y axis in 3d/2d (in degrees)

phi or theta can be a variable (see below)

*vector* args = x y z

x y z = vector direction to apply the acceleration

x or y or z can be a variable (see below)

## 17.66.2 Examples

```
fix 1 all gravity 1.0 chute 24.0
fix 1 all gravity v_increase chute 24.0
fix 1 all gravity 1.0 spherical 0.0 -180.0
fix 1 all gravity 10.0 spherical v_phi v_theta
fix 1 all gravity 100.0 vector 1 1 0
```

## 17.66.3 Description

Impose an additional acceleration on each particle in the group. This fix is typically used with granular systems to include a “gravity” term acting on the macroscopic particles. More generally, it can represent any kind of driving field, e.g. a pressure gradient inducing a Poiseuille flow in a fluid. Note that this fix operates differently than the *fix addforce* command. The *addforce* fix adds the same force to each atom, independent of its mass. This command imparts the same acceleration to each atom (force/mass).

The *magnitude* of the acceleration is specified in force/mass units. For granular systems (LJ units) this is typically 1.0. See the *units* command for details.

Style *chute* is typically used for simulations of chute flow where the specified *angle* is the chute angle, with flow occurring in the +x direction. For 3d systems, the tilt is away from the z axis; for 2d systems, the tilt is away from the y axis.

Style *spherical* allows an arbitrary 3d direction to be specified for the acceleration vector. *Phi* and *theta* are defined in the usual spherical coordinates. Thus for acceleration acting in the -z direction, *theta* would be 180.0 (or -180.0). *Theta* = 90.0 and *phi* = -90.0 would mean acceleration acts in the -y direction. For 2d systems, *phi* is ignored and *theta* is an angle in the xy plane where *theta* = 0.0 is the y-axis.

Style *vector* imposes an acceleration in the vector direction given by (x,y,z). Only the direction of the vector is important; it’s length is ignored. For 2d systems, the z component is ignored.

Any of the quantities *magnitude*, *angle*, *phi*, *theta*, *x*, *y*, *z* which define the gravitational magnitude and direction, can be specified as an equal-style *variable*. If the value is a variable, it should be specified as *v\_name*, where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the quantity. You should insure that the variable calculates a result in the appropriate units, e.g. force/mass or degrees.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent gravitational field.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 17.66.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the gravitational potential energy of the system to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global scalar which can be accessed by various *output commands*. This scalar is the gravitational potential energy of the particles in the defined field, namely  $\text{mass} * (\mathbf{g} \cdot \mathbf{x})$  for each particles, where *x* and *mass* are the particles position and mass, and *g* is the gravitational field. The scalar value calculated by this fix is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.66.5 Restrictions

none

#### 17.66.6 Related commands

*atom\_style sphere*, *fix addforce*



## 17.66.7 Default

none

## 17.67 fix grem command

### 17.67.1 Syntax

```
fix ID group-ID grem lambda eta H0 thermostat-ID
```

- ID, group-ID are documented in *fix* command
- grem = style name of this fix command
- lambda = intercept parameter of linear effective temperature function
- eta = slope parameter of linear effective temperature function
- H0 = shift parameter of linear effective temperature function
- thermostat-ID = ID of Nose-Hoover thermostat or barostat used in simulation

### 17.67.2 Examples

```
fix          fxgREM all grem 400 -0.01 -30000 fxnpt
thermo_modify press fxgREM_press

fix          fxgREM all grem 502 -0.15 -80000 fxnvt
```

### 17.67.3 Description

This fix implements the molecular dynamics version of the generalized replica exchange method (gREM) originally developed by (*Kim*), which uses non-Boltzmann ensembles to sample over first order phase transitions. The is done by defining replicas with an enthalpy dependent effective temperature

$$T_{eff} = \lambda + \eta(H - H_0)$$

with  $\eta$  negative and steep enough to only intersect the characteristic microcanonical temperature ( $T_s$ ) of the system once, ensuring a unimodal enthalpy distribution in that replica.  $\lambda$  is the intercept and effects the generalized ensemble similar to how temperature effects a Boltzmann ensemble.  $H_0$  is a reference enthalpy, and is typically set as the lowest desired sampled enthalpy. Further explanation can be found in our recent papers (*Malolepsza*).

This fix requires a Nose-Hoover thermostat fix reference passed to the grem as *thermostat-ID*. Two distinct temperatures exist in this generalized ensemble, the effective temperature defined above, and a kinetic temperature that controls the velocity distribution of particles as usual. Either constant volume or constant pressure algorithms can be used.

The fix enforces a generalized ensemble in a single replica only. Typically, this ideology is combined with replica exchange with replicas differing by  $\lambda$  only for simplicity, but this is not required. A multi-replica simulation can be run within the LAMMPS environment using the *temper/grem* command. This utilizes LAMMPS partition mode and requires the number of available processors be on the order of the number of desired replicas. A 100-replica simulation would require at least 100 processors (1 per world at minimum). If many replicas are needed on a small

number of processors, multi-replica runs can be run outside of LAMMPS. An example of this can be found in examples/PACKAGES/grem and has no limit on the number of replicas per processor. However, this is very inefficient and error prone and should be avoided if possible.

In general, defining the generalized ensembles is unique for every system. When starting a many-replica simulation without any knowledge of the underlying microcanonical temperature, there are several tricks we have utilized to optimize the process. Choosing a less-steep  $\eta$  yields broader distributions, requiring fewer replicas to map the microcanonical temperature. While this likely struggles from the same sampling problems gREM was built to avoid, it provides quick insight to  $T_s$ . Initially using an evenly-spaced  $\lambda$  distribution identifies regions where small changes in enthalpy lead to large temperature changes. Replicas are easily added where needed.

---

## 17.67.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *thermo\_modify press* option is supported by this fix to add the rescaled kinetic pressure as part of *thermodynamic output*.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the effective temperature  $T_{eff}$ . The scalar value calculated by this fix is “intensive”.

## 17.67.5 Restrictions

This fix is part of the REPLICa package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 17.67.6 Related commands

*temper/grem, fix nvt, fix npt, thermo\_modify*

## 17.67.7 Default

none

---

**(Kim)** Kim, Keyes, Straub, J Chem. Phys, 132, 224107 (2010).

**(Malolepsza)** Malolepsza, Secor, Keyes, J Phys Chem B 119 (42), 13379-13384 (2015).

## 17.68 fix halt command

### 17.68.1 Syntax

```
fix ID group-ID halt N attribute operator avalue keyword value ...
```

- ID, group-ID are documented in *fix* command
- halt = style name of this fix command
- N = check halt condition every N steps

- attribute = *bondmax* or *tlimit* or *v\_name*  
*bondmax* = length of longest bond in the system (in length units)  
*tlimit* = elapsed CPU time (in seconds)  
*diskfree* = free disk space (in MBytes)  
*v\_name* = name of *equal-style variable*
- operator = "<" or "<=" or ">" or ">=" or "==" or "!=" or "|^"
- avalue = numeric value to compare attribute to
- zero or more keyword/value pairs may be appended
- keyword = *error* or *message* or *path*  
*error* value = *hard* or *soft* or *continue*  
*message* value = *yes* or *no*  
*path* value = path to check for free space (may be in quotes)

## 17.68.2 Examples

```
fix 10 all halt 1 bondmax > 1.5
fix 10 all halt 10 v_myCheck != 0 error soft
fix 10 all halt 100 diskfree < 1000000.0 path "dump storage/."
```

## 17.68.3 Description

Check a condition every *N* steps during a simulation run. *N* must be  $\geq 1$ . If the condition is met, exit the run. In this context a “run” can be dynamics or minimization iterations, as specified by the *run* or *minimize* command.

The specified group-ID is ignored by this fix.

The specified *attribute* can be one of the options listed above, namely *bondmax*, *tlimit*, *diskfree*, or an *equal-style variable* referenced as *v\_name*, where “name” is the name of a variable that has been defined previously in the input script.

The *bondmax* attribute will loop over all bonds in the system, compute their current lengths, and set *attribute* to the longest bond distance.

The *tlimit* attribute queries the elapsed CPU time (in seconds) since the current run began, and sets *attribute* to that value. This is an alternative way to limit the length of a simulation run, similar to the *timer* timeout command. There are two differences in using this method versus the timer command option. The first is that the clock starts at the beginning of the current run (not when the timer or fix command is specified), so that any setup time for the run is not included in the elapsed time. The second is that the timer invocation and syncing across all processors (via MPI\_Allreduce) is not performed once every *N* steps by this command. Instead it is performed (typically) only a small number of times and the elapsed times are used to predict when the end-of-the-run will be. Both of these attributes can be useful when performing benchmark calculations for a desired length of time with minimal overhead. For example, if a run is performing 1000s of timesteps/sec, the overhead for syncing the timer frequently across a large number of processors may be non-negligible.

The *diskfree* attribute will check for available disk space (in MBytes) on supported operating systems. By default it will check the file system of the current working directory. This can be changed with the optional *path* keyword, which will take the path to a file or folder on the file system to be checked as argument. This path must be given with single or double quotes, if it contains blanks or other special characters (like \$).

Equal-style variables evaluate to a numeric value. See the *variable* command for a description. They calculate formulas which can involve mathematical operations, atom properties, group properties, thermodynamic properties, global

values calculated by a *compute* or *fix*, or references to other *variables*. Thus they are a very general means of computing some attribute of the current system. For example, the following “bondmax” variable will calculate the same quantity as the `hstyle = bondmax` option.

```
compute      bdist all bond/local dist
compute      bmax all reduce max c_bdist
variable     bondmax equal c_bmax
```

Thus these two versions of a `fix halt` command will do the same thing:

```
fix 10 all halt 1 bondmax > 1.5
fix 10 all halt 1 v_bondmax > 1.5
```

The version with “bondmax” will just run somewhat faster, due to less overhead in computing bond lengths and not storing them in a separate compute.

A variable can be used to implement a large variety of conditions, including to stop when a specific file exists. Example:

```
variable exit equal is_file(EXIT)
fix 10 all halt 100 v_exit != 0 error soft
```

Will stop the current run command when a file `EXIT` is created in the current working directory. The condition can be cleared by removing the file through the *shell* command.

The choice of operators listed above are the usual comparison operators. The XOR operation (exclusive or) is also included as “|<sup>^</sup>”. In this context, XOR means that if either the attribute or avalue is 0.0 and the other is non-zero, then the result is “true”. Otherwise it is “false”.

The specified *avalue* must be a numeric value.

---

The optional *error* keyword determines how the current run is halted. If its value is *hard*, then LAMMPS will stop with an error message.

If its value is *soft*, LAMMPS will exit the current run, but continue to execute subsequent commands in the input script. However, additional *run* or *minimize* commands will be skipped. For example, this allows a script to output the current state of the system, e.g. via a *write\_dump* or *write\_restart* command.

If its value is *continue*, the behavior is the same as for *soft*, except subsequent *run* or *minimize* commands are executed. This allows your script to remedy the condition that triggered the halt, if necessary. Note that you may wish use the *unfix* command on the `fix halt` ID, so that the same condition is not immediately triggered in a subsequent run.

The optional *message* keyword determines whether a message is printed to the screen and logfile when the halt condition is triggered. If *message* is set to yes, a one line message with the values that triggered the halt is printed. If *message* is set to no, no message is printed; the run simply exits. The latter may be desirable for post-processing tools that extract thermodynamic information from log files.

#### 17.68.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

## 17.68.5 Restrictions

The *diskfree* attribute is currently only supported on Linux, MacOSX, and BSD.

## 17.68.6 Related commands

*variable*

## 17.68.7 Default

The option defaults are error = hard, message = yes, and path = “.”.

## 17.69 fix heat command

### 17.69.1 Syntax

```
fix ID group-ID heat N eflux
```

- ID, group-ID are documented in *fix* command
- heat = style name of this fix command
- N = add/subtract heat every this many timesteps
- eflux = rate of heat addition or subtraction (energy/time units)
- eflux can be a variable (see below)
- zero or more keyword/value pairs may be appended to args
- keyword = *region*

```
region value = region-ID
```

```
region-ID = ID of region atoms must be in to have added force
```

### 17.69.2 Examples

```
fix 3 qin heat 1 1.0
fix 3 qin heat 10 v_flux
fix 4 qout heat 1 -1.0 region top
```

### 17.69.3 Description

Add non-translational kinetic energy (heat) to a group of atoms in a manner that conserves their aggregate momentum. Two of these fixes can be used to establish a temperature gradient across a simulation domain by adding heat (energy) to one group of atoms (hot reservoir) and subtracting heat from another (cold reservoir). E.g. a simulation sampling from the McDLT ensemble.

If the *region* keyword is used, the atom must be in both the group and the specified geometric *region* in order to have energy added or subtracted to it. If not specified, then the atoms in the group are affected wherever they may move to.

Heat addition/subtraction is performed every N timesteps.

The *eflux* parameter can be specified as a numeric constant or as an equal- or atom-style *variable*. If the value is a variable, it should be specified as *v\_name*, where *name* is the variable name. In this case, the variable will be evaluated each timestep, and its current value(s) used to determine the flux.

If *eflux* is a numeric constant or equal-style variable which evaluates to a scalar value, then *eflux* determines the change in aggregate energy of the entire group of atoms per unit time, e.g. in eV/ps for *metal units*. In this case it is an “extensive” quantity, meaning its magnitude should be scaled with the number of atoms in the group. Note that since *eflux* also has per-time units (i.e. it is a flux), this means that a larger value of *N* will add/subtract a larger amount of energy each time the fix is invoked.

---

**Note:** The heat-exchange (HEX) algorithm implemented by this fix is known to exhibit a pronounced energy drift. An improved algorithm (eHEX) is available as a *fix ehex* command and might be preferable if energy conservation is important.

---

If *eflux* is specified as an atom-style variable (see below), then the variable computes one value per atom. In this case, each value is the energy flux for a single atom, again in units of energy per unit time. In this case, each value is an “intensive” quantity, which need not be scaled with the number of atoms in the group.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent flux.

Atom-style variables can specify the same formulas as equal-style variables but can also include per-atom values, such as atom coordinates. Thus it is easy to specify a spatially-dependent flux with optional time-dependence as well.

---

**Note:** If heat is subtracted from the system too aggressively so that the group’s kinetic energy would go to zero, or any individual atom’s kinetic energy would go to zero for the case where *eflux* is an atom-style variable, then LAMMPS will halt with an error message.

---

Fix heat is different from a thermostat such as *fix nvt* or *fix temp/rescale* in that energy is added/subtracted continually. Thus if there is not another mechanism in place to counterbalance this effect, the entire system will heat or cool continuously. You can use multiple heat fixes so that the net energy change is 0.0 or use *fix viscous* to drain energy from the system.

This fix does not change the coordinates of its atoms; it only scales their velocities. Thus you must still use an integration fix (e.g. *fix nve*) on the affected atoms. This fix should not normally be used on atoms that have their temperature controlled by another fix - e.g. *fix nvt* or *fix langevin* fix.

## 17.69.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global scalar which can be accessed by various *output commands*. This scalar is the most recent value by which velocities were scaled. The scalar value calculated by this fix is “intensive”. If *eflux* is specified as an atom-style variable, this fix computes the average value by which the velocities were scaled for all of the atoms that had their velocities scaled.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.69.5 Restrictions

none

## 17.69.6 Related commands

*fix ehex, compute temp, compute temp/region*

## 17.69.7 Default

none

# 17.70 fix hyper/global command

## 17.70.1 Syntax

```
fix ID group-ID hyper/global cutbond qfactor Vmax Tequil
```

- ID, group-ID are documented in *fix* command
- hyper/global = style name of this fix command
- cutbond = max distance at which a pair of atoms is considered bonded (distance units)
- qfactor = max strain at which bias potential goes to 0.0 (unitless)
- Vmax = height of bias potential (energy units)
- Tequil = equilibration temperature (temperature units)

## 17.70.2 Examples

```
fix 1 all hyper/global 1.0 0.3 0.8 300.0
```

## 17.70.3 Description

This fix is meant to be used with the *hyper* command to perform a bond-boost global hyperdynamics (GHD) simulation. The role of this fix is to select a single pair of atoms in the system at each timestep to add a global bias potential to, which will alter the dynamics of the system in a manner that effectively accelerates time. This is in contrast to the *fix hyper/local* command, which can be used to perform a local hyperdynamics (LHD) simulation, by adding a local bias potential to multiple pairs of atoms at each timestep. GHD can time accelerate a small simulation with up to a few 100 atoms. For larger systems, LHD is needed to achieve good time acceleration.

For a system that undergoes rare transition events, where one or more atoms move over an energy barrier to a new potential energy basin, the effect of the bias potential is to induce more rapid transitions. This can lead to a dramatic speed-up in the rate at which events occurs, without altering their relative frequencies, thus leading to an overall increase in the elapsed real time of the simulation as compared to running for the same number of timesteps with normal MD. See the *hyper* page for a more general discussion of hyperdynamics and citations that explain both GHD and LHD.

The equations and logic used by this fix and described here to perform GHD follow the description given in (Voter2013). The bond-boost form of a bias potential for HD is due to Miron and Fichthorn as described in (Miron). In LAMMPS we use a simplified version of bond-boost GHD where a single bond in the system is biased at any one timestep.

Bonds are defined between each pair of atoms  $ij$ , whose  $R_{ij}^0$  distance is less than  $cutbond$ , when the system is in a quenched state (minimum) energy. Note that these are not “bonds” in a covalent sense. A bond is simply any pair of atoms that meet the distance criterion.  $Cutbond$  is an argument to this fix; it is discussed below. A bond is only formed if one or both of the  $ij$  atoms are in the specified group.

The current strain of bond  $ij$  (when running dynamics) is defined as

$$E_{ij} = \frac{R_{ij} - R_{ij}^0}{R_{ij}^0}$$

where  $R_{ij}$  is the current distance between atoms  $i$  and  $j$ , and  $R_{ij}^0$  is the equilibrium distance in the quenched state.

The bias energy  $V_{ij}$  of any bond between atoms  $i$  and  $j$  is defined as

$$V_{ij} = V^{max} \cdot \left( 1 - \left( \frac{E_{ij}}{q} \right)^2 \right) \text{ for } |E_{ij}| < qfactor \text{ or } 0 \text{ otherwise}$$

where the prefactor  $V^{max}$  and the cutoff  $qfactor$  are arguments to this fix; they are discussed below. This functional form is an inverse parabola centered at 0.0 with height  $V^{max}$  and which goes to 0.0 at +/-  $qfactor$ .

Let  $E^{max}$  be the maximum of  $|E_{ij}|$  for all  $ij$  bonds in the system on a given timestep. On that step,  $V_{ij}$  is added as a bias potential to only the single bond with strain  $E^{max}$ , call it  $V_{ij}^{max}$ . Note that  $V_{ij}^{max}$  will be 0.0 if  $E^{max} \geq qfactor$  on that timestep. Also note that  $V_{ij}^{max}$  is added to the normal interatomic potential that is computed between all atoms in the system at every step.

The derivative of  $V_{ij}^{max}$  with respect to the position of each atom in the  $E^{max}$  bond gives a bias force  $F_{ij}^{max}$  acting on the bond as

$$F_{ij}^{max} = -\frac{dV_{ij}^{max}}{dE_{ij}} = \frac{2V^{max} E - ij}{qfactor^2} \text{ for } |E_{ij}| < qfactor \text{ or } 0 \text{ otherwise}$$

which can be decomposed into an equal and opposite force acting on only the two  $ij$  atoms in the  $E^{max}$  bond.

The time boost factor for the system is given each timestep  $I$  by

$$B_i = e^{\beta V_{ij}^{max}}$$

where  $\beta = \frac{1}{kT_{equil}}$ , and  $T_{equil}$  is the temperature of the system and an argument to this fix. Note that  $B_i \geq 1$  at every step.

---

**Note:** To run a GHD simulation, the input script must also use the *fix langevin* command to thermostat the atoms at the same  $T_{equil}$  as specified by this fix, so that the system is running constant-temperature (NVT) dynamics. LAMMPS does not check that this is done.

---

The elapsed time  $t_{hyper}$  for a GHD simulation running for  $N$  timesteps is simply

$$t_{hyper} = \sum_{i=1, N} B - i \cdot dt$$

where  $dt$  is the timestep size defined by the *timestep* command. The effective time acceleration due to GHD is thus  $t_{hyper} / N \cdot dt$ , where  $N \cdot dt$  is elapsed time for a normal MD run of  $N$  timesteps.

Note that in GHD, the boost factor varies from timestep to timestep. Likewise, which bond has  $E^{max}$  strain and thus which pair of atoms the bias potential is added to, will also vary from timestep to timestep. This is in contrast to local hyperdynamics (LHD) where the boost factor is an input parameter; see the *fix hyper/local* page for details.



Here is additional information on the input parameters for GHD.

The *cutbond* argument is the cutoff distance for defining bonds between pairs of nearby atoms. A pair of *ij* atoms in their equilibrium, minimum-energy configuration, which are separated by a distance  $R_{ij} < \text{cutbond}$ , are flagged as a bonded pair. Setting *cutbond* to be ~25% larger than the nearest-neighbor distance in a crystalline lattice is a typical choice for solids, so that bonds exist only between nearest neighbor pairs.

The *qfactor* argument is the limiting strain at which the bias potential goes to 0.0. It is dimensionless, so a value of 0.3 means a bond distance can be up to 30% larger or 30% smaller than the equilibrium (quenched)  $R_{0ij}$  distance and the two atoms in the bond could still experience a non-zero bias force.

If *qfactor* is set too large, then transitions from one energy basin to another are affected because the bias potential is non-zero at the transition state (e.g. saddle point). If *qfactor* is set too small then little boost is achieved because the  $E_{ij}$  strain of some bond in the system will (nearly) always exceed *qfactor*. A value of 0.3 for *qfactor* is typically reasonable.

The *Vmax* argument is the prefactor on the bias potential. Ideally, it should be set to a value slightly less than the smallest barrier height for an event to occur. Otherwise the applied bias potential may be large enough (when added to the interatomic potential) to produce a local energy basin with a maxima in the center. This can produce artificial energy minima in the same basin that trap an atom. Or if *Vmax* is even larger, it may induce an atom(s) to rapidly transition to another energy basin. Both cases are “bad dynamics” which violate the assumptions of GHD that guarantee an accelerated time-accurate trajectory of the system.

Note that if *Vmax* is set too small, the GHD simulation will run correctly. There will just be fewer events because the hyper time ( $t_{\text{hyper}}$  equation above) will be shorter.

---

**Note:** If you have no physical intuition as to the smallest barrier height in your system, a reasonable strategy to determine the largest *Vmax* you can use for a GHD model, is to run a sequence of simulations with smaller and smaller *Vmax* values, until the event rate does not change (as a function of hyper time).

---

The *Tequil* argument is the temperature at which the system is simulated; see the comment above about the *fix langevin* thermostating. It is also part of the beta term in the exponential factor that determines how much boost is achieved as a function of the bias potential.

In general, the lower the value of *Tequil* and the higher the value of *Vmax*, the more time boost will be achievable by the GHD algorithm.

---

## 17.70.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the energy of the bias potential to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

This fix computes a global scalar and global vector of length 12, which can be accessed by various *output commands*. The scalar is the magnitude of the bias potential (energy units) applied on the current timestep. The vector stores the following quantities:

- 1 = boost factor on this step (unitless)
- 2 = max strain  $E_{ij}$  of any bond on this step (absolute value, unitless)
- 3 = ID of first atom in the max-strain bond
- 4 = ID of second atom in the max-strain bond

- 5 = average # of bonds/atom on this step
- 6 = fraction of timesteps where the biased bond has bias = 0.0 during this run
- 7 = fraction of timesteps where the biased bond has negative strain during this run
- 8 = max drift distance of any atom during this run (distance units)
- 9 = max bond length during this run (distance units)
- 10 = cumulative hyper time since fix was defined (time units)
- 11 = cumulative count of event timesteps since fix was defined
- 12 = cumulative count of atoms in events since fix was defined

The first 5 quantities are for the current timestep. Quantities 6-9 are for the current hyper run. They are reset each time a new hyper run is performed. Quantities 10-12 are cumulative across multiple runs (since the point in the input script the fix was defined).

For value 8, drift is the distance an atom moves between two quenched states when the second quench determines an event has occurred. Atoms involved in an event will typically move the greatest distance since others typically remain near their original quenched position.

For value 11, events are checked for by the *hyper* command once every *Nevent* timesteps. This value is the count of those timesteps on which one (or more) events was detected. It is NOT the number of distinct events, since more than one event may occur in the same *Nevent* time window.

For value 12, each time the *hyper* command checks for an event, it invokes a compute to flag zero or more atoms as participating in one or more events. E.g. atoms that have displaced more than some distance from the previous quench state. Value 12 is the cumulative count of the number of atoms participating in any of the events that were found.

The scalar and vector values calculated by this fix are all “intensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.70.5 Restrictions

This command can only be used if LAMMPS was built with the REPLICA package. See the *Build package* page for more info.

### 17.70.6 Related commands

*hyper*, *fix hyper/local*

### 17.70.7 Default

none

---

(**Voter2013**) S. Y. Kim, D. Perez, A. F. Voter, J Chem Phys, 139, 144110 (2013).

(**Miron**) R. A. Miron and K. A. Fichthorn, J Chem Phys, 119, 6210 (2003).

## 17.71 fix hyper/local command

### 17.71.1 Syntax

```
fix ID group-ID hyper/local cutbond qfactor Vmax Tequil Dcut alpha Btarget
```

- ID, group-ID are documented in *fix* command
- hyper/local = style name of this fix command
- cutbond = max distance at which a pair of atoms is considered bonded (distance units)
- qfactor = max strain at which bias potential goes to 0.0 (unitless)
- Vmax = estimated height of bias potential (energy units)
- Tequil = equilibration temperature (temperature units)
- Dcut = minimum distance between boosted bonds (distance units)
- alpha = boostostat relaxation time (time units)
- Btarget = desired time boost factor (unitless)
- zero or more keyword/value pairs may be appended
- keyword = *bound* or *reset* or *check/ghost* or *check/bias*

*bound* value = Bfrac

Bfrac = -1 or a value  $\geq 0.0$

*reset* value = Rfreq

Rfreq = -1 or 0 or timestep value  $> 0$

*check/ghost* values = none

*check/bias* values = Nevery error/warn/ignore

### 17.71.2 Examples

```
fix 1 all hyper/local 1.0 0.3 0.8 300.0
fix 1 all hyper/local 1.0 0.3 0.8 300.0 bound 0.1 reset 0
```

### 17.71.3 Description

This fix is meant to be used with the *hyper* command to perform a bond-boost local hyperdynamics (LHD) simulation. The role of this fix is to select multiple pairs of atoms in the system at each timestep to add a local bias potential, which will alter the dynamics of the system in a manner that effectively accelerates time. This is in contrast to the *fix hyper/global* command, which can be used to perform a global hyperdynamics (GHD) simulation, by adding a global bias potential to a single pair of atoms at each timestep. GHD can time accelerate a small simulation with up to a few 100 atoms. For larger systems, LHD is needed to achieve good time acceleration.

For a system that undergoes rare transition events, where one or more atoms move over an energy barrier to a new potential energy basin, the effect of the bias potential is to induce more rapid transitions. This can lead to a dramatic speed-up in the rate at which events occurs, without altering their relative frequencies, thus leading to an overall increase in the elapsed real time of the simulation as compared to running for the same number of timesteps with normal MD. See the *hyper* page for a more general discussion of hyperdynamics and citations that explain both GHD and LHD.

The equations and logic used by this fix and described here to perform LHD follow the description given in (Voter2013). The bond-boost form of a bias potential for HD is due to Miron and Fichthorn as described in (Miron).

To understand this description, you should first read the description of the GHD algorithm on the *fix hyper/global* doc page. This description of LHD builds on the GHD description.

The definition of bonds and  $E_{ij}$  are the same for GHD and LHD. The formulas for  $V_{ij}^{max}$  and  $F_{ij}^{max}$  are also the same except for a prefactor  $C_{ij}$ , explained below.

The bias energy  $V_{ij}$  applied to a bond  $ij$  with maximum strain is

$$V_{ij}^{max} = C_{ij} \cdot V^{max} \cdot \left( 1 - \left( \frac{E_{ij}}{q} \right)^2 \right) \text{ for } |E_{ij}| < qfactor \text{ or } 0 \text{ otherwise}$$

The derivative of  $V_{ij}^{max}$  with respect to the position of each atom in the  $ij$  bond gives a bias force  $F_{ij}^{max}$  acting on the bond as

$$F_{ij}^{max} = -\frac{dV_{ij}^{max}}{dE_{ij}} = 2C_{ij}V^{max} \frac{E_{ij}}{qfactor^2} \text{ for } |E_{ij}| < qfactor \text{ or } 0 \text{ otherwise}$$

which can be decomposed into an equal and opposite force acting on only the two atoms  $i$  and  $j$  in the  $ij$  bond.

The key difference is that in GHD a bias energy and force is added (on a particular timestep) to only one bond (pair of atoms) in the system, which is the bond with maximum strain  $E^{max}$ .

In LHD, a bias energy and force can be added to multiple bonds separated by the specified  $Dcut$  distance or more. A bond  $ij$  is biased if it is the maximum strain bond within its local “neighborhood”, which is defined as the bond  $ij$  plus any neighbor bonds within a distance  $Dcut$  from  $ij$ . The “distance” between bond  $ij$  and bond  $kl$  is the minimum distance between any of the  $ik$ ,  $il$ ,  $jk$ , and  $jl$  pairs of atoms.

For a large system, multiple bonds will typically meet this requirement, and thus a bias potential  $V_{ij}^{max}$  will be applied to many bonds on the same timestep.

In LHD, all bonds store a  $C_{ij}$  prefactor which appears in the  $V_{ij}^{max}$  and  $F_{ij}^{max}$  equations above. Note that the  $C_{ij}$  factor scales the strength of the bias energy and forces whenever bond  $ij$  is the maximum strain bond in its neighborhood.

$C_{ij}$  is initialized to 1.0 when a bond between the  $ij$  atoms is first defined. The specified  $Btarget$  factor is then used to adjust the  $C_{ij}$  prefactors for each bond every timestep in the following manner.

An instantaneous boost factor  $B_{ij}$  is computed each timestep for each bond, as

$$B_{ij} = e^{\beta V_{kl}^{max}}$$

where  $V_{kl}^{max}$  is the bias energy of the maxstrain bond  $kl$  within bond  $ij$ 's neighborhood,  $\beta = \frac{1}{kT_{equil}}$ , and  $T_{equil}$  is the temperature of the system and an argument to this fix.

---

**Note:** To run an LHD simulation, the input script must also use the *fix langevin* command to thermostat the atoms at the same  $T_{equil}$  as specified by this fix, so that the system is running constant-temperature (NVT) dynamics. LAMMPS does not check that this is done.

---

Note that if  $ij == kl$ , then bond  $ij$  is a biased bond on that timestep, otherwise it is not. But regardless, the boost factor  $B_{ij}$  can be thought of an estimate of time boost currently being applied within a local region centered on bond  $ij$ . For LHD, we want this to be the specified  $Btarget$  value everywhere in the simulation domain.

To accomplish this, if  $B_{ij} < Btarget$ , the  $C_{ij}$  prefactor for bond  $ij$  is incremented on the current timestep by an amount proportional to the inverse of the specified  $\alpha$  and the difference ( $B_{ij} - Btarget$ ). Conversely if  $B_{ij} > Btarget$ ,  $C_{ij}$  is decremented by the same amount. This procedure is termed “boostostatting” in (Voter2013). It drives all of the individual  $C_{ij}$  to values such that when  $V_{ij}^{max}$  is applied as a bias to bond  $ij$ , the resulting boost factor  $B_{ij}$  will be close to  $Btarget$  on average. Thus the LHD time acceleration factor for the overall system is effectively  $Btarget$ .

Note that in LHD, the boost factor  $B_{target}$  is specified by the user. This is in contrast to global hyperdynamics (GHD) where the boost factor varies each timestep and is computed as a function of  $V_{max}$ ,  $E_{max}$ , and  $T_{equil}$ ; see the [fix hyper/global](#) page for details.

Here is additional information on the input parameters for LHD.

Note that the *cutbond*, *qfactor*, and *Tequil* arguments have the same meaning as for GHD. The *Vmax* argument is slightly different. The *Dcut*, *alpha*, and *Btarget* parameters are unique to LHD.

The *cutbond* argument is the cutoff distance for defining bonds between pairs of nearby atoms. A pair of I,J atoms in their equilibrium, minimum-energy configuration, which are separated by a distance  $R_{ij} < cutbond$ , are flagged as a bonded pair. Setting *cutbond* to be ~25% larger than the nearest-neighbor distance in a crystalline lattice is a typical choice for solids, so that bonds exist only between nearest neighbor pairs.

The *qfactor* argument is the limiting strain at which the bias potential goes to 0.0. It is dimensionless, so a value of 0.3 means a bond distance can be up to 30% larger or 30% smaller than the equilibrium (quenched)  $R_{ij}^0$  distance and the two atoms in the bond could still experience a non-zero bias force.

If *qfactor* is set too large, then transitions from one energy basin to another are affected because the bias potential is non-zero at the transition state (e.g. saddle point). If *qfactor* is set too small than little boost can be achieved because the  $E_{ij}$  strain of some bond in the system will (nearly) always exceed *qfactor*. A value of 0.3 for *qfactor* is typically a reasonable value.

The *Vmax* argument is a fixed prefactor on the bias potential. There is also a dynamic prefactor  $C_{ij}$ , driven by the choice of *Btarget* as discussed above. The product of these should be a value less than the smallest barrier height for an event to occur. Otherwise the applied bias potential may be large enough (when added to the interatomic potential) to produce a local energy basin with a maxima in the center. This can produce artificial energy minima in the same basin that trap an atom. Or if  $C_{ij} \cdot V^{max}$  is even larger, it may induce an atom(s) to rapidly transition to another energy basin. Both cases are “bad dynamics” which violate the assumptions of LHD that guarantee an accelerated time-accurate trajectory of the system.

---

**Note:** It may seem that  $V^{max}$  can be set to any value, and  $C_{ij}$  will compensate to reduce the overall prefactor if necessary. However the  $C_{ij}$  are initialized to 1.0 and the booststating procedure typically operates slowly enough that there can be a time period of bad dynamics if  $V^{max}$  is set too large. A better strategy is to set  $V^{max}$  to the slightly smaller than the lowest barrier height for an event (the same as for GHD), so that the  $C_{ij}$  remain near unity.

---

The *Tequil* argument is the temperature at which the system is simulated; see the comment above about the [fix langevin](#) thermostating. It is also part of the beta term in the exponential factor that determines how much boost is achieved as a function of the bias potential. See the discussion of the *Btarget* argument below.

As discussed above, the *Dcut* argument is the distance required between two locally maxstrain bonds for them to both be selected as biased bonds on the same timestep. Computationally, the larger *Dcut* is, the more work (computation and communication) must be done each timestep within the LHD algorithm. And the fewer bonds can be simultaneously biased, which may mean the specified *Btarget* time acceleration cannot be achieved.

Physically *Dcut* should be a long enough distance that biasing two pairs of atoms that close together will not influence the dynamics of each pair. E.g. something like 2x the cutoff of the interatomic potential. In practice a *Dcut* value of ~10 Angstroms seems to work well for many solid-state systems.

---

**Note:** You should insure that ghost atom communication is performed for a distance of at least  $Dcut + cutedvent =$  the distance one or more atoms move (between quenched states) to be considered an “event”. It is an argument to the “compute event/displace” command used to detect events. By default the ghost communication distance is set by the *pair\_style* cutoff, which will typically be  $< Dcut$ . The *comm\_modify cutoff* command should be used to override the ghost cutoff explicitly, e.g.

---

`comm_modify` cutoff 12.0

Note that this fix does not know the *cutevent* parameter, but uses half the *cutbond* parameter as an estimate to warn if the ghost cutoff is not long enough.

As described above the *alpha* argument is a prefactor in the boostostat update equation for each bond's  $C_{ij}$  prefactor. *Alpha* is specified in time units, similar to other thermostat or barostat damping parameters. It is roughly the physical time it will take the boostostat to adjust a  $C_{ij}$  value from a too high (or too low) value to a correct one. An *alpha* setting of a few ps is typically good for solid-state systems. Note that the *alpha* argument here is the inverse of the alpha parameter discussed in (Voter2013).

The *Btarget* argument is the desired time boost factor (a value > 1) that all the atoms in the system will experience. The elapsed time *t\_hyper* for an LHD simulation running for *N* timesteps is simply

$$t_{hyper} = B_{target} \cdot N \cdot dt$$

where *dt* is the timestep size defined by the *timestep* command. The effective time acceleration due to LHD is thus  $\frac{t_{hyper}}{N \cdot dt} = B_{target}$ , where  $N \cdot dt$  is the elapsed time for a normal MD run of *N* timesteps.

You cannot choose an arbitrarily large setting for *Btarget*. The maximum value you should choose is

$$B_{target} = e^{\beta V_{small}}$$

where  $V_{small}$  is the smallest event barrier height in your system,  $\beta = \frac{1}{kT_{equil}}$ , and  $T_{equil}$  is the specified temperature of the system (both by this fix and the Langevin thermostat).

Note that if *Btarget* is set smaller than this, the LHD simulation will run correctly. There will just be fewer events because the hyper time (*t\_hyper* equation above) will be shorter.

---

**Note:** If you have no physical intuition as to the smallest barrier height in your system, a reasonable strategy to determine the largest *Btarget* you can use for an LHD model, is to run a sequence of simulations with smaller and smaller *Btarget* values, until the event rate does not change (as a function of hyper time).

---

Here is additional information on the optional keywords for this fix.

The *bound* keyword turns on min/max bounds for bias coefficients  $C_{ij}$  for all bonds.  $C_{ij}$  is a prefactor for each bond on the bias potential of maximum strength  $V^{max}$ . Depending on the choice of *alpha* and *Btarget* and *Vmax*, the boostostatting can cause individual  $C_{ij}$  values to fluctuate. If the fluctuations are too large  $C_{ij} \cdot V^{max}$  can exceed low barrier heights and induce bad event dynamics. Bounding the  $C_{ij}$  values is a way to prevent this. If *Bfrac* is set to -1 or any negative value (the default) then no bounds are enforced on  $C_{ij}$  values (except they must always be  $\geq 0.0$ ). A *Bfrac* setting  $\geq 0.0$  sets a lower bound of  $1.0 - Bfrac$  and upper bound of  $1.0 + Bfrac$  on each  $C_{ij}$  value. Note that all  $C_{ij}$  values are initialized to 1.0 when a bond is created for the first time. Thus *Bfrac* limits the bias potential height to  $Vmax \pm Bfrac \cdot Vmax$ .

The *reset* keyword allow *Vmax* to be adjusted dynamically depending on the average value of all  $C_{ij}$  prefactors. This can be useful if you are unsure what value of *Vmax* will match the *Btarget* boost for the system. The  $C_{ij}$  values will then adjust in aggregate (up or down) so that  $C_{ij} \cdot V^{max}$  produces a boost of *Btarget*, but this may conflict with the *bound* keyword settings. By using *bound* and *reset* together,  $V^{max}$  itself can be reset, and desired bounds still applied to the  $C_{ij}$  values.

A setting for *Rfreq* of -1 (the default) means *Vmax* never changes. A setting of 0 means  $V^{max}$  is adjusted every time an event occurs and bond pairs are recalculated. A setting of  $N > 0$  timesteps means  $V^{max}$  is adjusted on the first time an event occurs on a timestep  $\geq N$  steps after the previous adjustment. The adjustment to  $V^{max}$  is computed as follows. The current average of all  $C_{ij} \cdot V^{max}$  values is computed and the  $V^{max}$  is reset to that value. All  $C_{ij}$  values are changed to new prefactors such the new  $C_{ij} \cdot V^{max}$  is the same as it was previously. If the *bound* keyword was

used, those bounds are enforced on the new  $C_{ij}$  values. Henceforth, new bonds are assigned a  $C_{ij} = 1.0$ , which means their bias potential magnitude is the new  $V^{max}$ .

The *check/ghost* keyword turns on extra computation each timestep to compute statistics about ghost atoms used to determine which bonds to bias. The output of these stats are the vector values 14 and 15, described below. If this keyword is not enabled, the output of the stats will be zero.

The *check/bias* keyword turns on extra computation and communication to check if any biased bonds are closer than *Dcut* to each other, which should not be the case if LHD is operating correctly. Thus it is a debugging check. The *Nevery* setting determines how often the check is made. The *error*, *warn*, or *ignore* setting determines what is done if the count of too-close bonds is not zero. Either the code will exit, or issue a warning, or silently tally the count. The count can be output as vector value 17, as described below. If this keyword is not enabled, the output of that statistic will be 0.

Note that both of these computations are costly, hence they are only enabled by these keywords.

---

### 17.71.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the energy of the bias potential to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

This fix computes a global scalar and global vector of length 28, which can be accessed by various *output commands*. The scalar is the magnitude of the bias potential (energy units) applied on the current timestep, summed over all biased bonds. The vector stores the following quantities:

- 1 = average boost for all bonds on this step (unitless)
- 2 = # of biased bonds on this step
- 3 = max strain  $E_{ij}$  of any bond on this step (absolute value, unitless)
- 4 = value of  $V^{max}$  on this step (energy units)
- 5 = average bias coeff for all bonds on this step (unitless)
- 6 = min bias coeff for all bonds on this step (unitless)
- 7 = max bias coeff for all bonds on this step (unitless)
- 8 = average # of bonds/atom on this step
- 9 = average neighbor bonds/bond on this step within *Dcut*
- 10 = average boost for all bonds during this run (unitless)
- 11 = average # of biased bonds/step during this run
- 12 = fraction of biased bonds with no bias during this run
- 13 = fraction of biased bonds with negative strain during this run
- 14 = max bond length during this run (distance units)
- 15 = average bias coeff for all bonds during this run (unitless)
- 16 = min bias coeff for any bond during this run (unitless)
- 17 = max bias coeff for any bond during this run (unitless)
- 18 = max drift distance of any bond atom during this run (distance units)

- 19 = max distance from proc subbox of any ghost atom with  $\text{maxstrain} < qfactor$  during this run (distance units)
- 20 = max distance outside my box of any ghost atom with any  $\text{maxstrain}$  during this run (distance units)
- 21 = count of ghost atoms that could not be found on reneighbor steps during this run
- 22 = count of bias overlaps ( $< Dcut$ ) found during this run
- 23 = cumulative hyper time since fix created (time units)
- 24 = cumulative count of event timesteps since fix created
- 25 = cumulative count of atoms in events since fix created
- 26 = cumulative # of new bonds formed since fix created

27 = average boost for biased bonds on this step (unitless) 28 = # of bonds with absolute strain  $\geq q$  on this step

The first quantities 1-9 are for the current timestep. Quantities 10-22 are for the current hyper run. They are reset each time a new hyper run is performed. Quantities 23-26 are cumulative across multiple runs (since the point in the input script the fix was defined).

For value 10, each bond instantaneous boost factor is given by the equation for  $B_{ij}$  above. The total system boost (average across all bonds) fluctuates, but should average to a value close to the specified  $B_{target}$ .

For value 12, the numerator is a count of all biased bonds on each timestep whose bias energy = 0.0 due to  $E_{ij} \geq qfactor$ . The denominator is the count of all biased bonds on all timesteps.

For value 13, the numerator is a count of all biased bonds on each timestep with negative strain. The denominator is the count of all biased bonds on all timesteps.

Values 18-22 are mostly useful for debugging and diagnostic purposes.

For value 18, drift is the distance an atom moves between two quenched states when the second quench determines an event has occurred. Atoms involved in an event will typically move the greatest distance since others typically remain near their original quenched position.

For values 19-21, neighbor atoms in the full neighbor list with cutoff  $Dcut$  may be ghost atoms outside a processor's sub-box. Before the next event occurs they may move further than  $Dcut$  away from the sub-box boundary. Value 19 is the furthest (from the sub-box) any ghost atom in the neighbor list with  $\text{maxstrain} < qfactor$  was accessed during the run. Value 20 is the same except that the ghost atom's  $\text{maxstrain}$  may be  $\geq qfactor$ , which may mean it is about to participate in an event. Value 21 is a count of how many ghost atoms could not be found on reneighbor steps, presumably because they moved too far away due to their participation in an event (which will likely be detected at the next quench).

Typical values for 19 and 20 should be slightly larger than  $Dcut$ , which accounts for ghost atoms initially at a  $Dcut$  distance moving thermally before the next event takes place.

Note that for values 19 and 20 to be computed, the optional keyword *check/ghost* must be specified. Otherwise these values will be zero. This is because computing them incurs overhead, so the values are only computed if requested.

Value 21 should be zero or small. As explained above a small count likely means some ghost atoms were participating in their own events and moved a longer distance. If the value is large, it likely means the communication cutoff for ghosts is too close to  $Dcut$  leading to many not-found ghost atoms before the next event. This may lead to a reduced number of bonds being selected for biasing, since the code assumes those atoms are part of highly strained bonds. As explained above, the *comm\_modify cutoff* command can be used to set a longer cutoff.

For value 22, no two bonds should be biased if they are within a  $Dcut$  distance of each other. This value should be zero, indicating that no pair of biased bonds are closer than  $Dcut$  from each other.

Note that for value 22 to be computed, the optional keyword *check/bias* must be specified and it determines how often this check is performed. This is because performing the check incurs overhead, so if only computed as often as requested.



The result at the end of the run is the cumulative total from every timestep the check was made. Note that the value is a count of atoms in bonds which found other atoms in bonds too close, so it is almost always an over-count of the number of too-close bonds.

Value 23 is simply the specified *boost* factor times the number of timesteps times the timestep size.

For value 24, events are checked for by the *hyper* command once every *Nevent* timesteps. This value is the count of those timesteps on which one (or more) events was detected. It is NOT the number of distinct events, since more than one event may occur in the same *Nevent* time window.

For value 25, each time the *hyper* command checks for an event, it invokes a compute to flag zero or more atoms as participating in one or more events. E.g. atoms that have displaced more than some distance from the previous quench state. Value 25 is the cumulative count of the number of atoms participating in any of the events that were found.

Value 26 tallies the number of new bonds created by the bond reset operation. Bonds between a specific I,J pair of atoms may persist for the entire hyperdynamics simulation if neither I or J are involved in an event.

Value 27 computes the average boost for biased bonds only on this step.

Value 28 is the count of bonds with an absolute value of strain  $\geq q$  on this step.

The scalar value is an “extensive” quantity since it grows with the system size; the vector values are all “intensive”.

This fix also computes a local vector of length the number of bonds currently in the system. The value for each bond is its  $C_{ij}$  prefactor (bias coefficient). These values can be accessed by various *output commands*. A particularly useful one is the *fix ave/histo* command which can be used to histogram the  $C_{ij}$  values to see if they are distributed reasonably close to 1.0, which indicates a good choice of  $V^{max}$ .

The local values calculated by this fix are unitless.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.71.5 Restrictions

This fix is part of the REPLICa package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

## 17.71.6 Related commands

*hyper*, *fix hyper/global*

## 17.71.7 Default

The default settings for optimal keywords are *bounds* = -1 and *reset* = -1. The *check/ghost* and *check/bias* keywords are not enabled by default.

---

(**Voter2013**) S. Y. Kim, D. Perez, A. F. Voter, J Chem Phys, 139, 144110 (2013).

(**Miron**) R. A. Miron and K. A. Fichthorn, J Chem Phys, 119, 6210 (2003).

## 17.72 fix imd command

### 17.72.1 Syntax

```
fix ID group-ID imd trate port keyword values ...
```

- ID, group-ID are documented in *fix* command
- imd = style name of this fix command
- port = port number on which the fix listens for an IMD client
- keyword = *unwrap* or *fscale* or *trate*

*unwrap* arg = *on* or *off*

*off* = coordinates are wrapped back into the principal unit cell (default)

*on* = "unwrapped" coordinates using the image flags used

*fscale* arg = factor

factor = floating point number to scale IMD forces (default: 1.0)

*trate* arg = transmission rate of coordinate data sets (default: 1)

*nowait* arg = *on* or *off*

*off* = LAMMPS waits to be connected to an IMD client before continuing (default)

*on* = LAMMPS listens for an IMD client, but continues with the run

### 17.72.2 Examples

```
fix vmd all imd 5678
fix comm all imd 8888 trate 5 unwrap on fscale 10.0
```

### 17.72.3 Description

This fix implements the “Interactive MD” (IMD) protocol which allows realtime visualization and manipulation of MD simulations through the IMD protocol, as initially implemented in VMD and NAMD. Specifically it allows LAMMPS to connect an IMD client, for example the [VMD visualization program](#), so that it can monitor the progress of the simulation and interactively apply forces to selected atoms.

If LAMMPS is compiled with the pre-processor flag `-DLAMMPS_ASYNC_IMD` then `fix imd` will use POSIX threads to spawn a IMD communication thread on MPI rank 0 in order to offload data reading and writing from the main execution thread and potentially lower the inferred latencies for slow communication links. This feature has only been tested under linux.

There are example scripts for using this package with LAMMPS in `examples/PACKAGES/imd`. Additional examples and a driver for use with the Novint Falcon game controller as haptic device can be found at: <http://sites.google.com/site/akohlmey/software/vrpn-icms>.

The source code for this fix includes code developed by the Theoretical and Computational Biophysics Group in the Beckman Institute for Advanced Science and Technology at the University of Illinois at Urbana-Champaign. We thank them for providing a software interface that allows codes like LAMMPS to hook to [VMD](#).

Upon initialization of the fix, it will open a communication port on the node with MPI task 0 and wait for an incoming connection. As soon as an IMD client is connected, the simulation will continue and the fix will send the current coordinates of the fix’s group to the IMD client at every `trate` MD step. When using `r-RESPA`, `trate` applies to the steps of the outmost `RESPA` level. During a run with an active IMD connection also the IMD client can request to apply forces to selected atoms of the fix group.

The port number selected must be an available network port number. On many machines, port numbers < 1024 are reserved for accounts with system manager privilege and specific applications. If multiple imd fixes would be active at the same time, each needs to use a different port number.

The *nowait* keyword controls the behavior of the fix when no IMD client is connected. With the default setting of *off*, LAMMPS will wait until a connection is made before continuing with the execution. Setting *nowait* to *on* will have the LAMMPS code be ready to connect to a client, but continue with the simulation. This can for example be used to monitor the progress of an ongoing calculation without the need to be permanently connected or having to download a trajectory file.

The *trate* keyword allows to select how often the coordinate data is sent to the IMD client. It can also be changed on request of the IMD client through an IMD protocol message. The *unwrap* keyword allows to send “unwrapped” coordinates to the IMD client that undo the wrapping back of coordinates into the principle unit cell, as done by default in LAMMPS. The *fscale* keyword allows to apply a scaling factor to forces transmitted by the IMD client. The IMD protocols stipulates that forces are transferred in kcal/mol/angstrom under the assumption that coordinates are given in angstrom. For LAMMPS runs with different units or as a measure to tweak the forces generated by the manipulation of the IMD client, this option allows to make adjustments.

To connect VMD to a listening LAMMPS simulation on the same machine with fix imd enabled, one needs to start VMD and load a coordinate or topology file that matches the fix group. When the VMD command prompts appears, one types the command line:

```
imd connect localhost 5678
```

This assumes that *fix imd* was started with 5678 as a port number for the IMD protocol.

The steps to do interactive manipulation of a running simulation in VMD are the following:

In the Mouse menu of the VMD Main window, select “Mouse -> Force -> Atom”. You may alternately select “Residue”, or “Fragment” to apply forces to whole residues or fragments. Your mouse can now be used to apply forces to your simulation. Click on an atom, residue, or fragment and drag to apply a force. Click quickly without moving the mouse to turn the force off. You can also use a variety of 3D position trackers to apply forces to your simulation. Game controllers or haptic devices with force-feedback such as the Novint Falcon or Sensable PHANTOM allow you to feel the resistance due to inertia or interactions with neighbors that the atoms experience you are trying to move, as if they were real objects. See the [VMD IMD Homepage](#) and the [VRPN-ICMS Homepage](#) for more details.

If IMD control messages are received, a line of text describing the message and its effect will be printed to the LAMMPS output screen, if screen output is active.

#### 17.72.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global scalar or vector or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.72.5 Restrictions

This fix is part of the MISC package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

When used in combination with VMD, a topology or coordinate file has to be loaded, which matches (in number and ordering of atoms) the group the fix is applied to. The fix internally sorts atom IDs by ascending integer value; in VMD (and thus the IMD protocol) those will be assigned 0-based consecutive index numbers.

When using multiple active IMD connections at the same time, each needs to use a different port number.

## 17.72.6 Related commands

none

## 17.72.7 Default

none

# 17.73 fix indent command

## 17.73.1 Syntax

```
fix ID group-ID indent K keyword values ...
```

- ID, group-ID are documented in *fix* command
- indent = style name of this fix command
- K = force constant for indenter surface (force/distance<sup>2</sup> units)
- one or more keyword/value pairs may be appended
- keyword = *sphere* or *cylinder* or *plane* or *side* or *units*

*sphere* args = x y z R

x,y,z = initial position of center of indenter (distance units)

R = sphere radius of indenter (distance units)

any of x,y,z,R can be a variable (see below)

*cylinder* args = dim c1 c2 R

dim = x or y or z = axis of cylinder

c1,c2 = coords of cylinder axis in other 2 dimensions (distance units)

R = cylinder radius of indenter (distance units)

any of c1,c2,R can be a variable (see below)

*plane* args = dim pos side

dim = x or y or z = plane perpendicular to this dimension

pos = position of plane in dimension x, y, or z (distance units)

pos can be a variable (see below)

side = *lo* or *hi*

*side* value = *in* or *out*

*in* = the indenter acts on particles inside the sphere or cylinder

*out* = the indenter acts on particles outside the sphere or cylinder

*units* value = *lattice* or *box*

lattice = the geometry is defined in lattice units  
 box = the geometry is defined in simulation box units

### 17.73.2 Examples

```
fix 1 all indent 10.0 sphere 0.0 0.0 15.0 3.0
fix 1 all indent 10.0 sphere v_x v_y 0.0 v_radius side in
fix 2 flow indent 10.0 cylinder z 0.0 0.0 10.0 units box
```

### 17.73.3 Description

Insert an indenter within a simulation box. The indenter repels all atoms in the group that touch it, so it can be used to push into a material or as an obstacle in a flow. Or it can be used as a constraining wall around a simulation; see the discussion of the *side* keyword below.

The indenter can either be spherical or cylindrical or planar. You must set one of those 3 keywords.

A spherical indenter exerts a force of magnitude

$$F(r) = -K (r - R)^2$$

on each atom where  $K$  is the specified force constant,  $r$  is the distance from the atom to the center of the indenter, and  $R$  is the radius of the indenter. The force is repulsive and  $F(r) = 0$  for  $r > R$ .

A cylindrical indenter exerts the same force, except that  $r$  is the distance from the atom to the center axis of the cylinder. The cylinder extends infinitely along its axis.

Spherical and cylindrical indenters account for periodic boundaries in two ways. First, the center point of a spherical indenter ( $x,y,z$ ) or axis of a cylindrical indenter ( $c1,c2$ ) is remapped back into the simulation box, if the box is periodic in a particular dimension. This occurs every timestep if the indenter geometry is specified with a variable (see below), e.g. it is moving over time. Second, the calculation of distance to the indenter center or axis accounts for periodic boundaries. Both of these mean that an indenter can effectively move through and straddle one or more periodic boundaries.

A planar indenter is really an axis-aligned infinite-extent wall exerting the same force on atoms in the system, where  $R$  is the position of the plane and  $r-R$  is the distance from the plane. If the *side* parameter of the plane is specified as *lo* then it will indent from the lo end of the simulation box, meaning that atoms with a coordinate less than the plane's current position will be pushed towards the hi end of the box and atoms with a coordinate higher than the plane's current position will feel no force. Vice versa if *side* is specified as *hi*.

Any of the 4 quantities defining a spherical indenter's geometry can be specified as an equal-style *variable*, namely  $x$ ,  $y$ ,  $z$ , or  $R$ . Similarly, for a cylindrical indenter, any of  $c1$ ,  $c2$ , or  $R$ , can be a variable. For a planar indenter, *pos* can be a variable. If the value is a variable, it should be specified as  $v\_name$ , where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to define the indenter geometry.

Note that equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify indenter properties that change as a function of time or span consecutive runs in a continuous fashion. For the latter, see the *start* and *stop* keywords of the *run* command and the *elaplong* keyword of *thermo\_style custom* for details.

For example, if a spherical indenter's x-position is specified as  $v\_x$ , then this variable definition will keep it's center at a relative position in the simulation box, 1/4 of the way from the left edge to the right edge, even if the box size changes:

```
variable x equal "xlo + 0.25*lx"
```

Similarly, either of these variable definitions will move the indenter from an initial position at 2.5 at a constant velocity of 5:

```
variable x equal "2.5 + 5*elaplong*dt"
variable x equal vdisplace(2.5,5)
```

If a spherical indenter's radius is specified as `v_r`, then these variable definitions will grow the size of the indenter at a specified rate.

```
variable r0 equal 0.0
variable rate equal 1.0
variable r equal "v_r0 + step*dt*v_rate"
```

If the *side* keyword is specified as *out*, which is the default, then particles outside the indenter are pushed away from its outer surface, as described above. This only applies to spherical or cylindrical indenters. If the *side* keyword is specified as *in*, the action of the indenter is reversed. Particles inside the indenter are pushed away from its inner surface. In other words, the indenter is now a containing wall that traps the particles inside it. If the radius shrinks over time, it will squeeze the particles.

The *units* keyword determines the meaning of the distance units used to define the indenter geometry. A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing. The (x,y,z) coords of the indenter position are scaled by the x,y,z lattice spacings respectively. The radius of a spherical or cylindrical indenter is scaled by the x lattice spacing.

Note that the *units* keyword only affects indenter geometry parameters specified directly with numbers, not those specified as variables. In the latter case, you should use the *xlat*, *ylat*, *zlat* keywords of the *thermo\_style* command if you want to include lattice spacings in a variable formula.

The force constant *K* is not affected by the *units* keyword. It is always in force/distance<sup>2</sup> units where force and distance are defined by the *units* command. If you wish *K* to be scaled by the lattice spacing, you can define *K* with a variable whose formula contains *xlat*, *ylat*, *zlat* keywords of the *thermo\_style* command, e.g.

```
variable k equal 100.0/xlat/xlat
fix 1 all indent $k sphere ...
```

#### 17.73.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the energy of interaction between atoms and the indenter to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*. The energy of each particle interacting with the indenter is  $K/3 (r - R)^3$ .

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global scalar energy and a global 3-vector of forces (on the indenter), which can be accessed by various *output commands*. The scalar and vector values calculated by this fix are “extensive”.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command. Note that if you define the indenter geometry with a variable using a time-dependent formula, LAMMPS uses the iteration count in the minimizer as the timestep. But it is almost certainly a bad idea to have the indenter change its position or size during a minimization. LAMMPS does not check if you have done this.

---

**Note:** If you want the atom/indenter interaction energy to be included in the total potential energy of the system (the quantity being minimized), you must enable the *fix\_modify energy* option for this fix.

---

### 17.73.5 Restrictions

none

### 17.73.6 Related commands

none

### 17.73.7 Default

The option defaults are `side = out` and `units = lattice`.

## 17.74 fix ipi command

### 17.74.1 Syntax

```
fix ID group-ID ipi address port [unix] [reset]
```

- ID, group-ID are documented in *fix* command
- ipi = style name of this fix command
- address = internet address (FQDN or IP), or UNIX socket name
- port = port number (ignored for UNIX sockets)
- zero or more keywords may be appended
- keyword = *unix* or *reset*
  - unix* args = none = use a unix socket
  - reset* args = none = reset electrostatics at each call

### 17.74.2 Examples

```
fix 1 all ipi my.server.com 12345
fix 1 all ipi mysocket 666 unix reset
```

### 17.74.3 Description

This fix enables LAMMPS to be run as a client for the i-PI Python wrapper (*IPI*) for performing a path integral molecular dynamics (PIMD) simulation. The philosophy behind i-PI is described in the following publication (*IPI-CPC*).

A version of the i-PI package, containing only files needed for use with LAMMPS, is provided in the `tools/i-pi` directory. See the `tools/i-pi/manual.pdf` for an introduction to i-PI. The `examples/PACKAGES/i-pi` directory contains example scripts for using i-PI with LAMMPS.

In brief, the path integral molecular dynamics is performed by the Python wrapper, while the client (LAMMPS in this case) simply computes forces and energy for each configuration. The communication between the two components takes place using sockets, and is reduced to the bare minimum. All the parameters of the dynamics are specified in the input of i-PI, and all the parameters of the force field must be specified as LAMMPS inputs, preceding the *fix ipi* command.

The server address must be specified by the *address* argument, and can be either the IP address, the fully-qualified name of the server, or the name of a UNIX socket for local, faster communication. In the case of internet sockets, the *port* argument specifies the port number on which i-PI is listening, while the *unix* optional switch specifies that the socket is a UNIX socket.

Note that there is no check of data integrity, or that the atomic configurations make sense. It is assumed that the species in the i-PI input are listed in the same order as in the data file of LAMMPS. The initial configuration is ignored, as it will be substituted with the coordinates received from i-PI before forces are ever evaluated.

A note of caution when using potentials that contain long-range electrostatics, or that contain parameters that depend on box size: all of these options will be initialized based on the cell size in the LAMMPS-side initial configuration and kept constant during the run. This is required to e.g. obtain reproducible and conserved forces. If the cell varies too wildly, it may be advisable to re-initialize these interactions at each call. This behavior can be requested by setting the *reset* switch.

### 17.74.4 Restart, fix\_modify, output, run start/stop, minimize info

There is no restart information associated with this fix, since all the dynamical parameters are dealt with by i-PI.

### 17.74.5 Restrictions

Using this fix on anything other than all atoms requires particular care, since i-PI will know nothing on atoms that are not those whose coordinates are transferred. However, one could use this strategy to define an external potential acting on the atoms that are moved by i-PI.

This fix is part of the MISC package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info. Because of the use of UNIX domain sockets, this fix will only work in a UNIX environment.

### 17.74.6 Related commands

*fix nve*

---

(*IPI-CPC*) Ceriotti, More and Manolopoulos, *Comp Phys Comm*, 185, 1019-1026 (2014).

(*IPI*) <http://epfl-cosmo.github.io/gle4md/index.html?page=ipi>



## 17.75 fix langevin command

Accelerator Variants: *langevin/kk*

### 17.75.1 Syntax

```
fix ID group-ID langevin Tstart Tstop damp seed keyword values ...
```

- ID, group-ID are documented in *fix* command
- langevin = style name of this fix command
- Tstart,Tstop = desired temperature at start/end of run (temperature units)
- Tstart can be a variable (see below)
- damp = damping parameter (time units)
- seed = random number seed to use for white noise (positive integer)
- zero or more keyword/value pairs may be appended
- keyword = *angmom* or *omega* or *scale* or *tally* or *zero*

*angmom* value = *no* or *factor*

*no* = do not thermostat rotational degrees of freedom via the angular momentum

*factor* = do thermostat rotational degrees of freedom via the angular momentum and

→ apply numeric scale factor as discussed below

*gjf* value = *no* or *vfull* or *vhalf*

*no* = use standard formulation

*vfull* = use Gronbech-Jensen/Farago formulation

*vhalf* = use 2GJ formulation

*omega* value = *no* or *yes*

*no* = do not thermostat rotational degrees of freedom via the angular velocity

*yes* = do thermostat rotational degrees of freedom via the angular velocity

*scale* values = type ratio

type = atom type (1-N)

ratio = factor by which to scale the damping coefficient

*tally* value = *no* or *yes*

*no* = do not tally the energy added/subtracted to atoms

*yes* = do tally the energy added/subtracted to atoms

*zero* value = *no* or *yes*

*no* = do not set total random force to zero

*yes* = set total random force to zero

### 17.75.2 Examples

```
fix 3 boundary langevin 1.0 1.0 1000.0 699483
fix 1 all langevin 1.0 1.1 100.0 48279 scale 3 1.5
fix 1 all langevin 1.0 1.1 100.0 48279 angmom 3.333
```

### 17.75.3 Description

Apply a Langevin thermostat as described in (*Schneider*) to a group of atoms which models an interaction with a background implicit solvent. Used with *fix nve*, this command performs Brownian dynamics (BD), since the total force on each atom will have the form:

$$\begin{aligned}F &= F_c + F_f + F_r \\F_f &= -\frac{m}{\text{damp}}v \\F_r &\propto \sqrt{\frac{k_B T m}{dt \text{ damp}}}\end{aligned}$$

$F_c$  is the conservative force computed via the usual inter-particle interactions (*pair\_style*, *bond\_style*, etc). The  $F_f$  and  $F_r$  terms are added by this fix on a per-particle basis. See the *pair\_style dpd/tstat* command for a thermostating option that adds similar terms on a pairwise basis to pairs of interacting particles.

$F_f$  is a frictional drag or viscous damping term proportional to the particle's velocity. The proportionality constant for each atom is computed as  $\frac{m}{\text{damp}}$ , where  $m$  is the mass of the particle and *damp* is the damping factor specified by the user.

$F_r$  is a force due to solvent atoms at a temperature  $T$  randomly bumping into the particle. As derived from the fluctuation/dissipation theorem, its magnitude as shown above is proportional to  $\sqrt{\frac{k_B T m}{dt \text{ damp}}}$ , where  $k_B$  is the Boltzmann constant,  $T$  is the desired temperature,  $m$  is the mass of the particle,  $dt$  is the timestep size, and *damp* is the damping factor. Random numbers are used to randomize the direction and magnitude of this force as described in (*Dunweg*), where a uniform random number is used (instead of a Gaussian random number) for speed.

Note that unless you use the *omega* or *angmom* keywords, the thermostat effect of this fix is applied to only the translational degrees of freedom for the particles, which is an important consideration for finite-size particles, which have rotational degrees of freedom, are being thermostatted. The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

---

**Note:** Unlike the *fix nvt* command which performs Nose/Hoover thermostating AND time integration, this fix does NOT perform time integration. It only modifies forces to effect thermostating. Thus you must use a separate time integration fix, like *fix nve* to actually update the velocities and positions of atoms using the modified forces. Likewise, this fix should not normally be used on atoms that also have their temperature controlled by another fix - e.g. by *fix nvt* or *fix temp/rescale* commands.

---

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostating.

The desired temperature at each timestep is a ramped value during the run from *Tstart* to *Tstop*.

*Tstart* can be specified as an equal-style or atom-style *variable*. In this case, the *Tstop* setting is ignored. If the value is a variable, it should be specified as *v\_name*, where *name* is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the target temperature.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent temperature.

Atom-style variables can specify the same formulas as equal-style variables but can also include per-atom values, such as atom coordinates. Thus it is easy to specify a spatially-dependent temperature with optional time-dependence as well.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a "bias" from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

The *damp* parameter is specified in time units and determines how rapidly the temperature is relaxed. For example, a value of 100.0 means to relax the temperature in a timespan of (roughly) 100 time units ( $\tau$  or fs or ps - see the *units* command). The damp factor can be thought of as inversely related to the viscosity of the solvent. I.e. a small relaxation time implies a high-viscosity solvent and vice versa. See the discussion about  $\gamma$  and viscosity in the documentation for the *fix viscous* command for more details.

The random # *seed* must be a positive integer. A Marsaglia random number generator is used. Each processor uses the input seed to generate its own unique seed and its own stream of random numbers. Thus the dynamics of the system will not be identical on two runs on different numbers of processors.

---

The keyword/value option pairs are used in the following ways.

The keyword *angmom* and *omega* keywords enable thermostating of rotational degrees of freedom in addition to the usual translational degrees of freedom. This can only be done for finite-size particles.

A simulation using *atom\_style sphere* defines an omega for finite-size spheres. A simulation using *atom\_style ellipsoid* defines a finite size and shape for aspherical particles and an angular momentum. The Langevin formulas for thermostating the rotational degrees of freedom are the same as those above, where force is replaced by torque, *m* is replaced by the moment of inertia *I*, and *v* is replaced by omega (which is derived from the angular momentum in the case of aspherical particles).

The rotational temperature of the particles can be monitored by the *compute temp/sphere* and *compute temp/asphere* commands with their rotate options.

For the *omega* keyword there is also a scale factor of  $\frac{10.0}{3.0}$  that is applied as a multiplier on the  $F_f$  (damping) term in the equation above and of  $\sqrt{\frac{10.0}{3.0}}$  as a multiplier on the  $F_r$  term. This does not affect the thermostating behavior of the Langevin formalism but insures that the randomized rotational diffusivity of spherical particles is correct.

For the *angmom* keyword a similar scale factor is needed which is  $\frac{10.0}{3.0}$  for spherical particles, but is anisotropic for aspherical particles (e.g. ellipsoids). Currently LAMMPS only applies an isotropic scale factor, and you can choose its magnitude as the specified value of the *angmom* keyword. If your aspherical particles are (nearly) spherical than a value of  $\frac{10.0}{3.0} = 3.\bar{3}$  is a good choice. If they are highly aspherical, a value of 1.0 is as good a choice as any, since the effects on rotational diffusivity of the particles will be incorrect regardless. Note that for any reasonable scale factor, the thermostating effect of the *angmom* keyword on the rotational temperature of the aspherical particles should still be valid.

The keyword *scale* allows the damp factor to be scaled up or down by the specified factor for atoms of that type. This can be useful when different atom types have different sizes or masses. It can be used multiple times to adjust damp for several atom types. Note that specifying a ratio of 2 increases the relaxation time which is equivalent to the solvent's viscosity acting on particles with  $\frac{1}{2}$  the diameter. This is the opposite effect of scale factors used by the *fix viscous* command, since the damp factor in *fix langevin* is inversely related to the  $\gamma$  factor in *fix viscous*. Also note that the damping factor in *fix langevin* includes the particle mass in  $F_f$ , unlike *fix viscous*. Thus the mass and size of different atom types should be accounted for in the choice of ratio values.

The keyword *tally* enables the calculation of the cumulative energy added/subtracted to the atoms as they are thermostatted. Effectively it is the energy exchanged between the infinite thermal reservoir and the particles. As described below, this energy can then be printed out or added to the potential energy of the system to monitor energy conservation.

---

**Note:** This accumulated energy does NOT include kinetic energy removed by the *zero* flag. LAMMPS will print a warning when both options are active.

---

The keyword *zero* can be used to eliminate drift due to the thermostat. Because the random forces on different atoms are independent, they do not sum exactly to zero. As a result, this fix applies a small random force to the entire system, and the center-of-mass of the system undergoes a slow random walk. If the keyword *zero* is set to *yes*, the total random force is set exactly to zero by subtracting off an equal part of it from each atom in the group. As a result, the center-of-mass of a system with zero initial momentum will not drift over time.

The keyword *gif* can be used to run the *Gronbech-Jensen/Farago* time-discretization of the Langevin model. As described in the papers cited below, the purpose of this method is to enable longer timesteps to be used (up to the numerical stability limit of the integrator), while still producing the correct Boltzmann distribution of atom positions.

The current implementation provides the user with the option to output the velocity in one of two forms: *vfull* or *vhalf*, which replaces the outdated option *yes*. The *gif* option *vfull* outputs the on-site velocity given in *Gronbech-Jensen/Farago*; this velocity is shown to be systematically lower than the target temperature by a small amount, which grows quadratically with the timestep. The *gif* option *vhalf* outputs the 2GJ half-step velocity given in *Gronbech-Jensen/Gronbech-Jensen*; for linear systems, this velocity is shown to not have any statistical errors for any stable time step. An overview of statistically correct Boltzmann and Maxwell-Boltzmann sampling of true on-site and true half-step velocities is given in *Gronbech-Jensen*. Regardless of the choice of output velocity, the sampling of the configurational distribution of atom positions is the same, and linearly consistent with the target temperature.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 17.75.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. Because the state of the random number generator is not saved in restart files, this means you cannot do “exact” restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a temperature *compute* you have defined to this fix which will be used in its thermostating procedure, as described above. For consistency, the group used by this fix and by the compute should be the same.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*, but only if the *tally* keyword is set to *yes*. See the *thermo\_style* page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”. Note that calculation of this quantity also requires setting the *tally* keyword to *yes*.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

## 17.75.5 Restrictions

For *gjf* do not choose `damp=dt/2`. *gjf* is not compatible with `run_style respa`.

## 17.75.6 Related commands

*fix nvt*, *fix temp/rescale*, *fix viscous*, *fix nvt*, *pair\_style dpd/tstat*

## 17.75.7 Default

The option defaults are `angmom = no`, `omega = no`, `scale = 1.0` for all types, `tally = no`, `zero = no`, `gjf = no`.

**(Dunweg)** Dunweg and Paul, Int J of Modern Physics C, 2, 817-27 (1991).

**(Schneider)** Schneider and Stoll, Phys Rev B, 17, 1302 (1978).

**(Gronbech-Jensen)** Gronbech-Jensen and Farago, Mol Phys, 111, 983 (2013); Gronbech-Jensen, Hayre, and Farago, Comp Phys Comm, 185, 524 (2014)

**(Gronbech-Jensen)** Gronbech Jensen and Gronbech-Jensen, Mol Phys, 117, 2511 (2019)

**(Gronbech-Jensen)** Gronbech-Jensen, Mol Phys (2019); <https://doi.org/10.1080/00268976.2019.1662506>

## 17.76 fix langevin/drude command

### 17.76.1 Syntax

```
fix ID group-ID langevin/drude Tcom damp_com seed_com Tdrude damp_drude seed_drude_
↪keyword values ...
```

- ID, group-ID are documented in *fix* command
- langevin/drude = style name of this fix command
- Tcom = desired temperature of the centers of mass (temperature units)
- damp\_com = damping parameter for the thermostat on centers of mass (time units)
- seed\_com = random number seed to use for white noise of the thermostat on centers of mass (positive integer)
- Tdrude = desired temperature of the Drude oscillators (temperature units)
- damp\_drude = damping parameter for the thermostat on Drude oscillators (time units)
- seed\_drude = random number seed to use for white noise of the thermostat on Drude oscillators (positive integer)
- zero or more keyword/value pairs may be appended
- keyword = *zero*

zero value = *no* or *yes*

*no* = do not set total random force on centers of mass to zero

*yes* = set total random force on centers of mass to zero

## 17.76.2 Examples

```
fix 3 all langevin/drude 300.0 100.0 19377 1.0 20.0 83451
fix 1 all langevin/drude 298.15 100.0 19377 5.0 10.0 83451 zero yes
```

Example input scripts available: examples/PACKAGES/drude

## 17.76.3 Description

Apply two Langevin thermostats as described in (*Jiang1*) for thermalizing the reduced degrees of freedom of Drude oscillators. This link describes how to use the *thermalized Drude oscillator model* in LAMMPS and polarizable models in LAMMPS are discussed on the *Howto polarizable* doc page.

Drude oscillators are a way to simulate polarizable atoms, by splitting them into a core and a Drude particle bound by a harmonic bond. The thermalization works by transforming the particles degrees of freedom by these equations. In these equations upper case denotes atomic or center of mass values and lower case denotes Drude particle or dipole values. Primes denote the transformed (reduced) values, while bare letters denote the original values.

Velocities:

$$V' = \frac{M V + m v}{M'}$$

$$v' = v - V$$

Masses:

$$M' = M + m$$

$$m' = \frac{M m}{M'}$$

The Langevin forces are computed as

$$F' = -\frac{M'}{\text{damp}_{\text{com}}} V' + F'_r$$

$$f' = -\frac{m'}{\text{damp}_{\text{drude}}} v' + f'_r$$

$F'_r$  is a random force proportional to  $\sqrt{\frac{2 k_B T_{\text{com}} m'}{dt \text{damp}_{\text{com}}}}$ .  $f'_r$  is a random force proportional to  $\sqrt{\frac{2 k_B T_{\text{drude}} m'}{dt \text{damp}_{\text{drude}}}}$ . Then the real forces acting on the particles are computed from the inverse transform:

$$F = \frac{M}{M'} F' - f'$$

$$f = \frac{m}{M'} F' + f'$$

This fix also thermostats non-polarizable atoms in the group at temperature  $T_{\text{com}}$ , as if they had a massless Drude partner. The Drude particles themselves need not be in the group. The center of mass and the dipole are thermostatted iff the core atom is in the group.

Note that the thermostat effect of this fix is applied to only the translational degrees of freedom of the particles, which is an important consideration if finite-size particles, which have rotational degrees of freedom, are being thermostatted. The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

**Note:** Like the *fix langevin* command, this fix does NOT perform time integration. It only modifies forces to effect thermostating. Thus you must use a separate time integration fix, like *fix nve* or *fix nph* to actually update the velocities

and positions of atoms using the modified forces. Likewise, this fix should not normally be used on atoms that also have their temperature controlled by another fix - e.g. by *fix nvt* or *fix temp/rescale* commands.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostatting.

This fix requires each atom know whether it is a Drude particle or not. You must therefore use the *fix drude* command to specify the Drude status of each atom type.

**Note:** only the Drude core atoms need to be in the group specified for this fix. A Drude electron will be transformed together with its cores even if it is not itself in the group. It is safe to include Drude electrons or non-polarizable atoms in the group. The non-polarizable atoms will simply be thermostatted as if they had a massless Drude partner (electron).

**Note:** Ghost atoms need to know their velocity for this fix to act correctly. You must use the *comm\_modify* command to enable this, e.g.

```
comm_modify vel yes
```

*Tcom* is the target temperature of the centers of mass, which would be used to thermostat the non-polarizable atoms. *Tdrude* is the (normally low) target temperature of the core-Drude particle pairs (dipoles). *Tcom* and *Tdrude* can be specified as an equal-style *variable*. If the value is a variable, it should be specified as *v\_name*, where *name* is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the target temperature.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent temperature.

Like other fixes that perform thermostatting, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostatting is performed on the remaining thermal degrees of freedom, and the bias is added back in.

Note: The temperature thermostatting the core-Drude particle pairs should be chosen low enough, so as to mimic as closely as possible the self-consistent minimization. It must however be high enough, so that the dipoles can follow the local electric field exerted by the neighboring atoms. The optimal value probably depends on the temperature of the centers of mass and on the mass of the Drude particles.

*damp\_com* is the characteristic time for reaching thermal equilibrium of the centers of mass. For example, a value of 100.0 means to relax the temperature of the centers of mass in a timespan of (roughly) 100 time units (tau or fs or ps - see the *units* command). *damp\_drude* is the characteristic time for reaching thermal equilibrium of the dipoles. It is typically a few timesteps.

The number *seed\_com* and *seed\_drude* are positive integers. They set the seeds of the Marsaglia random number generators used for generating the random forces on centers of mass and on the dipoles. Each processor uses the input seed to generate its own unique seed and its own stream of random numbers. Thus the dynamics of the system will not be identical on two runs on different numbers of processors.

The keyword *zero* can be used to eliminate drift due to the thermostat on centers of mass. Because the random forces on different centers of mass are independent, they do not sum exactly to zero. As a result, this fix applies a small random force to the entire system, and the momentum of the total center of mass of the system undergoes a slow random walk. If the keyword *zero* is set to *yes*, the total random force on the centers of mass is set exactly to zero by subtracting off an equal part of it from each center of mass in the group. As a result, the total center of mass of a system with zero initial momentum will not drift over time.

The actual temperatures of cores and Drude particles, in center-of-mass and relative coordinates, respectively, can be calculated using the *compute temp/drude* command.

---

Usage example for rigid bodies in the NPT ensemble:

```
comm_modify vel yes
fix TEMP all langevin/drude 300. 100. 1256 1. 20. 13977 zero yes
fix NPH ATOMS rigid/nph/small molecule iso 1. 1. 500.
fix NVE DRUDES nve
compute TDRUDE all temp/drude
thermo_style custom step cpu etotal ke pe ebond ecoul elong press vol temp c_TDRUDE[1] c_
↪TDRUDE[2]
```

Comments:

- Drude particles should not be in the rigid group, otherwise the Drude oscillators will be frozen and the system will lose its polarizability.
  - *zero yes* avoids a drift of the center of mass of the system, but is a bit slower.
  - Use two different random seeds to avoid unphysical correlations.
  - Temperature is controlled by the fix *langevin/drude*, so the time-integration fixes do not thermostat. Don't forget to time-integrate both cores and Drude particles.
  - Pressure is time-integrated only once by using *nve* for Drude particles and *nph* for atoms/cores (or vice versa). Do not use *nph* for both.
  - The temperatures of cores and Drude particles are calculated by *compute temp/drude*
  - Contrary to the alternative thermostating using Nose-Hoover thermostat fix *npt* and *fix drude/transform*, the *fix\_modify* command is not required here, because the fix *nph* computes the global pressure even if its group is *ATOMS*. This is what we want. If we thermostatted *ATOMS* using *npt*, the pressure should be the global one, but the temperature should be only that of the cores. That's why the command *fix\_modify* should be called in that case.
- 

#### 17.76.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. Because the state of the random number generator is not saved in restart files, this means you cannot do "exact" restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a temperature *compute* you have defined to this fix which will be used in its thermostating procedure, as described above. For consistency, the group used by the compute should include the group of this fix and the Drude particles.

This fix is not invoked during *energy minimization*.



## 17.76.5 Restrictions

none

## 17.76.6 Related commands

*fix langevin, fix drude, fix drude/transform, compute temp/drude, pair\_style thole*

## 17.76.7 Default

The option defaults are zero = no.

---

(Jiang<sup>1</sup>) Jiang, Hardy, Phillips, MacKerell, Schulten, and Roux, J Phys Chem Lett, 2, 87-92 (2011).

## 17.77 fix langevin/eff command

### 17.77.1 Syntax

```
fix ID group-ID langevin/eff Tstart Tstop damp seed keyword values ...
```

- ID, group-ID are documented in *fix* command
- langevin/eff = style name of this fix command
- Tstart,Tstop = desired temperature at start/end of run (temperature units)
- damp = damping parameter (time units)
- seed = random number seed to use for white noise (positive integer)
- zero or more keyword/value pairs may be appended

keyword = *scale* or *tally* or *zero*

*scale* values = type ratio

type = atom type (1-N)

ratio = factor by which to scale the damping coefficient

*tally* values = *no* or *yes*

*no* = do not tally the energy added/subtracted to atoms

*yes* = do tally the energy added/subtracted to atoms

*zero* value = *no* or *yes*

*no* = do not set total random force to zero

*yes* = set total random force to zero

## 17.77.2 Examples

```
fix 3 boundary langevin/eff 1.0 1.0 10.0 699483
fix 1 all langevin/eff 1.0 1.1 10.0 48279 scale 3 1.5
```

## 17.77.3 Description

Apply a Langevin thermostat as described in (*Schneider*) to a group of nuclei and electrons in the *electron force field* model. Used with *fix nve/eff*, this command performs Brownian dynamics (BD), since the total force on each atom will have the form:

$$F = F_c + F_f + F_r$$
$$F_f = - \frac{m}{\text{damp}} v$$
$$F_r \propto \sqrt{\frac{k_B T m}{dt \text{ damp}}}$$

$F_c$  is the conservative force computed via the usual inter-particle interactions (*pair\_style*). The  $F_f$  and  $F_r$  terms are added by this fix on a per-particle basis.

The operation of this fix is exactly like that described by the *fix langevin* command, except that the thermostating is also applied to the radial electron velocity for electron particles.

## 17.77.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. Because the state of the random number generator is not saved in restart files, this means you cannot do “exact” restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a temperature *compute* you have defined to this fix which will be used in its thermostating procedure, as described above. For consistency, the group used by this fix and by the compute should be the same.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*, but only if the *tally* keyword is set to *yes*. See the *thermo\_style* page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”. Note that calculation of this quantity also requires setting the *tally* keyword to *yes*.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

## 17.77.5 Restrictions

none

This fix is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 17.77.6 Related commands

*fix langevin*

## 17.77.7 Default

The option defaults are scale = 1.0 for all types and tally = no.

**(Dunweg)** Dunweg and Paul, Int J of Modern Physics C, 2, 817-27 (1991).

**(Schneider)** Schneider and Stoll, Phys Rev B, 17, 1302 (1978).

## 17.78 fix langevin/spin command

### 17.78.1 Syntax

```
fix ID group-ID langevin/spin T Tdamp seed
```

- ID, group-ID are documented in *fix* command
- langevin/spin = style name of this fix command
- T = desired temperature of the bath (temperature units, K in metal units)
- Tdamp = transverse magnetic damping parameter (adim)
- seed = random number seed to use for white noise (positive integer)

### 17.78.2 Examples

```
fix 2 all langevin/spin 300.0 0.01 21
```

### 17.78.3 Description

Apply a Langevin thermostat as described in ([Mayergoyz](#)) to the magnetic spins associated to the atoms. Used with *fix nve/spin*, this command performs Brownian dynamics (BD). A random torque and a transverse dissipation are applied to each spin  $i$  according to the following stochastic differential equation:

$$\frac{d\vec{s}_i}{dt} = \frac{1}{(1 + \lambda^2)} ((\vec{\omega}_i + \vec{\eta}) \times \vec{s}_i + \lambda \vec{s}_i \times (\vec{\omega}_i \times \vec{s}_i))$$

with  $\lambda$  the transverse damping, and  $\eta$  a random vector. This equation is referred to as the stochastic Landau-Lifshitz-Gilbert (sLLG) equation.

The components of  $\eta$  are drawn from a Gaussian probability law. Their amplitude is defined as a proportion of the temperature of the external thermostat T (in K in metal units).

More details about this implementation are reported in (*Tranchida*).

Note: due to the form of the sLLG equation, this fix has to be defined just before the nve/spin fix (and after all other magnetic fixes). As an example:

```
fix 1 all precession/spin zeeman 0.01 0.0 0.0 1.0
fix 2 all langevin/spin 300.0 0.01 21
fix 3 all nve/spin lattice moving
```

is correct, but defining a force/spin command after the langevin/spin command would give an error message.

Note: The random # *seed* must be a positive integer. A Marsaglia random number generator is used. Each processor uses the input seed to generate its own unique seed and its own stream of random numbers. Thus the dynamics of the system will not be identical on two runs on different numbers of processors.

---

#### 17.78.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. Because the state of the random number generator is not saved in restart files, this means you cannot do “exact” restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior.

This fix is not invoked during *energy minimization*.

#### 17.78.5 Restrictions

The *langevin/spin* fix is part of the SPIN package. This style is only enabled if LAMMPS was built with this package. See the *Build package* page for more info.

The numerical integration has to be performed with *fix nve/spin* when *fix langevin/spin* is enabled.

This fix has to be the last defined magnetic fix before the time integration fix (e.g. *fix nve/spin*).

#### 17.78.6 Related commands

*fix nve/spin*, *fix precession/spin*

#### 17.78.7 Default

none

---

(**Mayergoyz**) I.D. Mayergoyz, G. Bertotti, C. Serpico (2009). Elsevier (2009)

(**Tranchida**) Tranchida, Plimpton, Thibaudau and Thompson, Journal of Computational Physics, 372, 406-425, (2018).

## 17.79 fix latte command

### 17.79.1 Syntax

```
fix ID group-ID latte keyword value ...
```

- ID, group-ID are documented in *fix* command
- latte = style name of this fix command
- zero or more keyword/value pairs may be appended

keyword = *coulomb* or *exclude*

*coulomb* value = peID

peID = ID of compute used to calculate per-atom energy

*exclude* value = fixID

fixID = ID of fix which potentially excludes atoms before calling LATTE

### 17.79.2 Examples

```
fix dftb all latte
fix dftb all exclude GCMC
```

### 17.79.3 Description

This fix style is a wrapper on the self-consistent charge transfer density functional based tight binding (DFTB) code LATTE. If you download and build LATTE, it can be called as a library by LAMMPS via this fix to run dynamics or perform energy minimization using DFTB forces and energies computed by LATTE.

LATTE is principally developed and supported by Marc Cawkwell and co-workers at Los Alamos National Laboratory (LANL). See the full list of contributors in the `src/LATTE/README` file.

To use this fix, the LATTE program needs to be compiled as a library and linked with LAMMPS. LATTE can be downloaded (or cloned) from <https://github.com/lanl/LATTE>. Instructions on how to download and build LATTE on your system can be found in the `lib/latte/README`. Note that you can also use the “make lib-latte” command from the LAMMPS `src` directory to automate this process.

Once LAMMPS is built with the LATTE package, you can run the example input scripts for molecular dynamics or energy minimization that are found in `examples/latte`.

A step-by-step tutorial can be followed at: [LAMMPS-LATTE tutorial](#)

Currently, LAMMPS must be run in serial or as a single MPI task, to use this fix. This is because the version of the LATTE library LAMMPS uses does not support MPI. On the LAMMPS side, this is typically not a bottleneck, since LATTE will be doing 99% or more of the work to compute quantum-accurate forces. On the LATTE side, the LATTE library does support threaded parallelism via OpenMP. You must build the LATTE library with OpenMP support, then set the `OMP_NUM_THREADS` environment variable before performing a LAMMPS + LATTE simulation to tell LATTE how many threads to invoke.

---

**Note:** NEB calculations can be done using this fix using multiple replicas and running LAMMPS in parallel. However, each replica must be run on a single MPI task. For details, see the *neb* command page and the *-partition command-line switch*

---

The *coulomb* argument is not yet supported by fix latte (as of Sept 2022). Eventually it will be used to enable LAMMPS to calculate a Coulomb potential as an alternative to LATTE performing the calculation.

The *exclude* argument allows this fix to work in tandem with another fix which may decide to delete one or more atoms of molecules. The specified fixID is the ID of the other fix.

The one current example of such a fix is the *fix gcmc* command which performs Monte Carlo insertions and deletions. If a trial deletion is performed, then LAMMPS needs to only pass LATTE the atoms which remain. Fix gcmc does not actually remove any atoms until after the new energy is computed (in this case by LATTE), and a Monte Carlo accept/reject decision is made for the trial deletion.

---

LATTE is a code for performing self-consistent charge transfer tight-binding (SC-TB) calculations of total energies and the forces acting on atoms in molecules and solids. This tight-binding method is becoming more and more popular and widely used in chemistry, biochemistry, material science, etc.

The SC-TB formalism is derived from an expansion of the Kohn-Sham density functional to second order in charge fluctuations about a reference charge of overlapping atom-centered densities and bond integrals are parameterized using a Slater-Koster tight-binding approach. This procedure, which usually is referred to as the DFTB method has been described in detail by (*Elstner*) and (*Finnis*) and coworkers.

The work of the LATTE developers follows that of Elstner closely with respect to the physical model. However, the development of LATTE is geared principally toward large-scale, long duration, microcanonical quantum-based Born-Oppenheimer molecular dynamics (QMD) simulations. One of the main bottlenecks of an electronic structure calculation is the solution of the generalized eigenvalue problem which scales with the cube of the system size  $O(N^3)$ .

The Theoretical and Computer sciences divisions at Los Alamos National Laboratory have accumulated large experience addressing this issue by calculating the density matrix directly instead of using diagonalization. We typically use a recursive sparse Fermi-operator expansion using second-order spectral projection functions (SP2-algorithm), which was introduced by Niklasson in 2002 (*Niklasson2002*), (*Rubensson*), (*Mniszewski*). When the matrices involved in the recursive expansion are sufficiently sparse, the calculation of the density matrix scales linearly as a function of the system size  $O(N)$ .

Another important feature is the extended Lagrangian framework for Born-Oppenheimer molecular dynamics (XL-BOMD) (*Niklasson2008*) (*Niklasson2014*), (*Niklasson2017*) that allows for a drastic reduction or even a complete removal of the iterative self-consistent field optimization. Often only a single density matrix calculation per molecular dynamics time step is required, yet total energy stability is well maintained. The SP2 and XL-BOMD techniques enables stable linear scaling MD simulations with a very small computational overhead. This opens a number of opportunities in many different areas of chemistry and materials science, as we now can simulate larger system sizes and longer time scales (*Cawkwell2012*), (*Negre2016*).

---

#### 17.79.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy computed by LATTE to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy yes*.

The *fix\_modify virial* option is supported by this fix to add the contribution computed by LATTE to the global pressure of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify virial yes*.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the potential energy discussed above. The scalar value calculated by this fix is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The DFTB forces computed by LATTE via this fix are used during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the potential energy associated with the DFTB forces to be included in the total potential energy of the system (the quantity being minimized), you **MUST** not disable the *fix\_modify energy* option for this fix.

---

### 17.79.5 Restrictions

This fix is part of the LATTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

You must use metal units, as set by the *units* command to use this fix.

LATTE does not currently compute per-atom energy or per-atom virial contributions. So they will not show up as part of the calculations performed by the *compute pe/atom* or *compute stress/atom* commands.

### 17.79.6 Related commands

none

### 17.79.7 Default

none

---

**(Elstner)** M. Elstner, D. Poresag, G. Jungnickel, J. Elsner, M. Haugk, T. Frauenheim, S. Suhai, and G. Seifert, Phys. Rev. B, 58, 7260 (1998).

**(Elstner)** M. Elstner, D. Poresag, G. Jungnickel, J. Elsner, M. Haugk, T. Frauenheim, S. Suhai, and G. Seifert, Phys. Rev. B, 58, 7260 (1998).

**(Finnis)** M. W. Finnis, A. T. Paxton, M. Methfessel, and M. van Schilfgarde, Phys. Rev. Lett., 81, 5149 (1998).

**(Mniszewski)** S. M. Mniszewski, M. J. Cawkwell, M. E. Wall, J. Mohd-Yusof, N. Bock, T. C. Germann, and A. M. N. Niklasson, J. Chem. Theory Comput., 11, 4644 (2015).

**(Niklasson2002)** A. M. N. Niklasson, Phys. Rev. B, 66, 155115 (2002).

**(Rubensson)** E. H. Rubensson, A. M. N. Niklasson, SIAM J. Sci. Comput. 36 (2), 147-170, (2014).

**(Niklasson2008)** A. M. N. Niklasson, Phys. Rev. Lett., 100, 123004 (2008).

**(Niklasson2014)** A. M. N. Niklasson and M. Cawkwell, J. Chem. Phys., 141, 164123, (2014).

**(Niklasson2017)** A. M. N. Niklasson, J. Chem. Phys., 147, 054103 (2017).

**(Cawkwell2012)** A. M. N. Niklasson, M. J. Cawkwell, Phys. Rev. B, 86 (17), 174308 (2012).

**(Negre2016)** C. F. A. Negre, S. M. Mniszewski, M. J. Cawkwell, N. Bock, M. E. Wall, and A. M. N. Niklasson, J. Chem. Theory Comp., 12, 3063 (2016).

## 17.80 fix lb/fluid command

### 17.80.1 Syntax

```
fix ID group-ID lb/fluid nevery viscosity density keyword values ...
```

- ID, group-ID are documented in *fix* command
- lb/fluid = style name of this fix command
- nevery = update the lattice-Boltzmann fluid every this many timesteps (should normally be 1)
- viscosity = the fluid viscosity (units of mass/(time\*length)).
- density = the fluid density.
- zero or more keyword/value pairs may be appended
- keyword = *dx* or *dm* or *noise* or *stencil* or *read\_restart* or *write\_restart* or *zwall\_velocity* or *pressurebcx* or *bodyforce* or *D3Q19* or *dumpxdf* or *dof* or *scaleGamma* or *a0* or *npits* or *wp* or *sw*

*dx* values = *dx\_LB* = the lattice spacing.

*dm* values = *dm\_LB* = the lattice-Boltzmann mass unit.

*noise* values = Temperature seed

Temperature = fluid temperature.

seed = random number generator seed (positive integer)

*stencil* values = 2 (trilinear stencil, the default), 3 (3-point immersed boundary [↪stencil](#)), or 4 (4-point Keys' interpolation stencil)

*read\_restart* values = restart file = name of the restart file to use to restart a [↪fluid run](#).

*write\_restart* values = N = write a restart file every N MD timesteps.

*zwall\_velocity* values = *velocity\_bottom* *velocity\_top* = velocities along the [↪y-direction](#) of the bottom and top walls (located at *z=zmin* and *z=zmax*).

*pressurebcx* values = *pgradav* = imposes a pressure jump at the (periodic) x-boundary [↪of pgradav\\*Lx\\*1000](#).

*bodyforce* values = *bodyforcex* *bodyforcey* *bodyforcez* = the x,y and z components of a [↪constant body force](#) added to the fluid.

*D3Q19* values = none (used to switch from the default D3Q15, 15 velocity lattice, to [↪the D3Q19, 19 velocity lattice](#)).

*dumpxdf* values = N file timeI

N = output the force and torque every N timesteps

file = output file name

timeI = 1 (use simulation time to index xdf file), 0 (use output frame number [↪to index xdf file](#))

*dof* values = *dof* = specify the number of degrees of freedom for temperature [↪calculation](#)

*scaleGamma* values = type gammaFactor

type = atom type (1-N)

gammaFactor = factor to scale the *setGamma* gamma value by, for the specified [↪atom type](#).

*a0* values = *a0\_real* = the square of the speed of sound in the fluid.

*npits* values = *npits* *h\_p* *l\_p* *l\_pp* *l\_e*

*npits* = number of pit regions

*h\_p* = z-height of pit regions (floor to bottom of slit)

*l\_p* = x-length of pit regions

*l\_pp* = x-length of slit regions between consecutive pits



`l_e` = x-length of slit regions at ends  
`wp` values = `w_p` = y-width of slit regions (defaults to full width if not present or ↪  
↪if `sw` active)  
`sw` values = none (turns on y-sidewalls (in xz plane) if `npits` option active)

## 17.80.2 Examples

```

fix 1 all lb/fluid 1 1.0 0.0009982071 dx 1.2 dm 0.001
fix 1 all lb/fluid 1 1.0 0.0009982071 dx 1.2 dm 0.001 noise 300.0 2761
fix 1 all lb/fluid 1 1.0 1.0 dx 4.0 dm 10.0 dumpxdmf 500 fflow 0 pressurebcx 0.01 npits ↪
↪2 20 40 5 0 wp 30

```

## 17.80.3 Description

Changed in version 24Mar2022.

Implement a lattice-Boltzmann fluid on a uniform mesh covering the LAMMPS simulation domain. Note that this fix was updated in 2022 and is not backward compatible with the previous version. If you need the previous version, please download an older version of LAMMPS. The MD particles described by *group-ID* apply a velocity dependent force to the fluid.

The lattice-Boltzmann algorithm solves for the fluid motion governed by the Navier Stokes equations,

$$\begin{aligned} \partial_t \rho + \partial_\beta (\rho u_\beta) &= 0 \\ \partial_t (\rho u_\alpha) + \partial_\beta (\rho u_\alpha u_\beta) &= \partial_\beta \sigma_{\alpha\beta} + F_\alpha + \partial_\beta (\eta_{\alpha\beta\gamma\nu} \partial_\gamma u_\nu) \end{aligned}$$

with,

$$\eta_{\alpha\beta\gamma\nu} = \eta \left[ \delta_{\alpha\gamma} \delta_{\beta\nu} + \delta_{\alpha\nu} \delta_{\beta\gamma} - \frac{2}{3} \delta_{\alpha\beta} \delta_{\gamma\nu} \right] + \Lambda \delta_{\alpha\beta} \delta_{\gamma\nu}$$

where  $\rho$  is the fluid density,  $u$  is the local fluid velocity,  $\sigma$  is the stress tensor,  $F$  is a local external force, and  $\eta$  and  $\Lambda$  are the shear and bulk viscosities respectively. Here, we have implemented

$$\sigma_{\alpha\beta} = -P_{\alpha\beta} = -\rho a_0 \delta_{\alpha\beta}$$

with  $a_0$  set to  $\frac{1}{3} \frac{dx}{dt}^2$  by default. You should not normally need to change this default.

The algorithm involves tracking the time evolution of a set of partial distribution functions which evolve according to a velocity discretized version of the Boltzmann equation,

$$(\partial_t + e_{i\alpha} \partial_\alpha) f_i = -\frac{1}{\tau} (f_i - f_i^{eq}) + W_i$$

where the first term on the right hand side represents a single time relaxation towards the equilibrium distribution function, and  $\tau$  is a parameter physically related to the viscosity. On a technical note, we have implemented a 15 velocity model (D3Q15) as default; however, the user can switch to a 19 velocity model (D3Q19) through the use of the *D3Q19* keyword. Physical variables are then defined in terms of moments of the distribution functions,

$$\begin{aligned} \rho &= \sum_i f_i \\ \rho u_\alpha &= \sum_i f_i e_{i\alpha} \end{aligned}$$

Full details of the lattice-Boltzmann algorithm used can be found in *Denniston et al.*.

The fluid is coupled to the MD particles described by *group-ID* through a velocity dependent force. The contribution to the fluid force on a given lattice mesh site  $j$  due to MD particle  $\alpha$  is calculated as:

$$\mathbf{F}_{j\alpha} = \gamma (\mathbf{v}_n - \mathbf{u}_f) \zeta_{j\alpha}$$

where  $\mathbf{v}_n$  is the velocity of the MD particle,  $\mathbf{u}_f$  is the fluid velocity interpolated to the particle location, and  $\gamma$  is the force coupling constant. This force, as with most forces in LAMMPS, and hence the velocities, are calculated at the half-time step.  $\zeta$  is a weight assigned to the grid point, obtained by distributing the particle to the nearest lattice sites.

The force coupling constant,  $\gamma$ , is calculated according to

$$\gamma = \frac{2m_u m_v}{m_u + m_v} \left( \frac{1}{\Delta t} \right)$$

Here,  $m_v$  is the mass of the MD particle,  $m_u$  is a representative fluid mass at the particle location, and  $\Delta t$  is the time step. The fluid mass  $m_u$  that the MD particle interacts with is calculated internally. This coupling is chosen to constrain the particle and associated fluid velocity to match at the end of the time step. As with other constraints, such as *shake*, this constraint can remove degrees of freedom from the simulation which are accounted for internally in the algorithm.

---

**Note:** While this fix applies the force of the particles on the fluid, it does not apply the force of the fluid to the particles. There is only one option to include this hydrodynamic force on the particles, and that is through the use of the *lb/viscous* fix. This fix adds the hydrodynamic force to the total force acting on the particles, after which any of the built-in LAMMPS integrators can be used to integrate the particle motion. If the *lb/viscous* fix is NOT used to add the hydrodynamic force to the total force acting on the particles, this physically corresponds to a situation in which an infinitely massive particle is moving through the fluid (since collisions between the particle and the fluid do not act to change the particle's velocity). In this case, setting *scaleGamma* to -1 for the corresponding particle type will explicitly take this limit (of infinite particle mass) in computing the force coupling for the fluid force.

---

Physical parameters describing the fluid are specified through *viscosity* and *density*. These parameters should all be given in terms of the mass, distance, and time units chosen for the main LAMMPS run, as they are scaled by the LB timestep, lattice spacing, and mass unit, inside the fix.

The *dx* keyword allows the user to specify a value for the LB grid spacing and the *dm* keyword allows the user to specify the LB mass unit. Inside the fix, parameters are scaled by the lattice-Boltzmann timestep,  $dt_{LB}$ , grid spacing,  $dx_{LB}$ , and mass unit,  $dm_{LB}$ .  $dt_{LB}$  is set equal to  $nevery \cdot dt_{MD}$ , where  $dt_{MD}$  is the MD timestep. By default,  $dm_{LB}$  is set equal to 1.0, and  $dx_{LB}$  is chosen so that  $\frac{\tau}{dt} = \frac{3\eta dt}{\rho dx^2}$  is approximately equal to 1.

---

**Note:** Care must be taken when choosing both a value for  $dx_{LB}$ , and a simulation domain size. This fix uses the same subdivision of the simulation domain among processors as the main LAMMPS program. In order to uniformly cover the simulation domain with lattice sites, the lengths of the individual LAMMPS sub-domains must all be evenly divisible by  $dx_{LB}$ . If the simulation domain size is cubic, with equal lengths in all dimensions, and the default value for  $dx_{LB}$  is used, this will automatically be satisfied.

---

If the *noise* keyword is used, followed by a positive temperature value, and a positive integer random number seed, the thermal LB algorithm of *Adhikari et al.* is used.

If the keyword *stencil* is used, the value sets the number of interpolation points used in each direction. For this, the user has the choice between a trilinear stencil (*stencil* 2), which provides a support of 8 lattice sites, or the 3-point immersed boundary method stencil (*stencil* 3), which provides a support of 27 lattice sites, or the 4-point Keys' interpolation stencil (*stencil* 4), which provides a support of 64 lattice sites. The trilinear stencil is the default as it is better suited for simulation of objects close to walls or other objects, due to its smaller support. The 3-point stencil provides smoother motion of the lattice and is suitable for particles not likely to be too close to walls or other objects.

If the keyword *write\_restart* is used, followed by a positive integer, N, a binary restart file is printed every N LB timesteps. This restart file only contains information about the fluid. Therefore, a LAMMPS restart file should also be written in order to print out full details of the simulation.

---

**Note:** When a large number of lattice grid points are used, the restart files may become quite large.

---

In order to restart the fluid portion of the simulation, the keyword *read\_restart* is specified, followed by the name of the binary *lb\_fluid* restart file to be used.

If the *zwall\_velocity* keyword is used y-velocities are assigned to the lower and upper walls. This keyword requires the presence of walls in the z-direction. This is set by assigning fixed boundary conditions in the z-direction. If fixed boundary conditions are present in the z-direction, and this keyword is not used, the walls are assumed to be stationary.

If the *pressurebcx* keyword is used, a pressure jump (implemented by a step jump in density) is imposed at the (periodic) x-boundary. The value set specifies what would be the resulting equilibrium average pressure gradient in the x-direction if the system had a constant cross-section (i.e. resistance to flow). It is converted to a pressure jump by multiplication by the system size in the x-direction. As this value should normally be quite small, it is also assumed to be scaled by 1000.

If the *bodyforce* keyword is used, a constant body force is added to the fluid, defined by its x, y and z components.

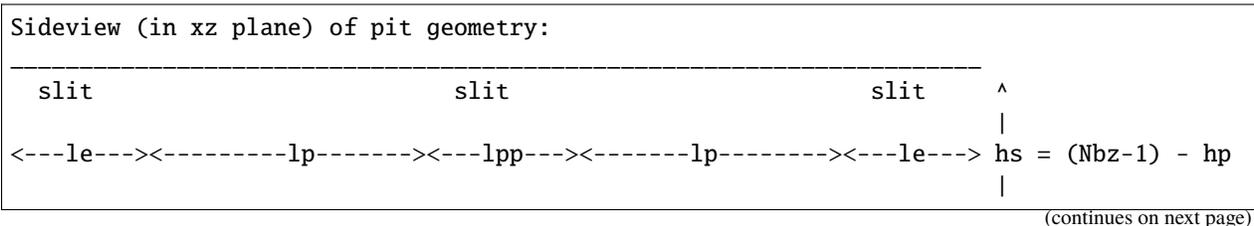
If the keyword *D3Q19* is used, the 19 velocity (D3Q19) lattice is used by the lattice-Boltzmann algorithm. By default, the 15 velocity (D3Q15) lattice is used.

If the *dumpxdmf* keyword is used, followed by a positive integer, N, and a file name, the fluid densities and velocities at each lattice site are output to an xdmf file every N timesteps. This is a binary file format that can be read by visualization packages such as [Paraview](#). The xdmf file format contains a time index for each frame dump and the value *timeI* = 1 uses simulation time while 0 uses the output frame number to index xdmf file. The later can be useful if the *dump vtk* command is used to output the particle positions at the same timesteps and you want to visualize both the fluid and particle data together in [Paraview](#).

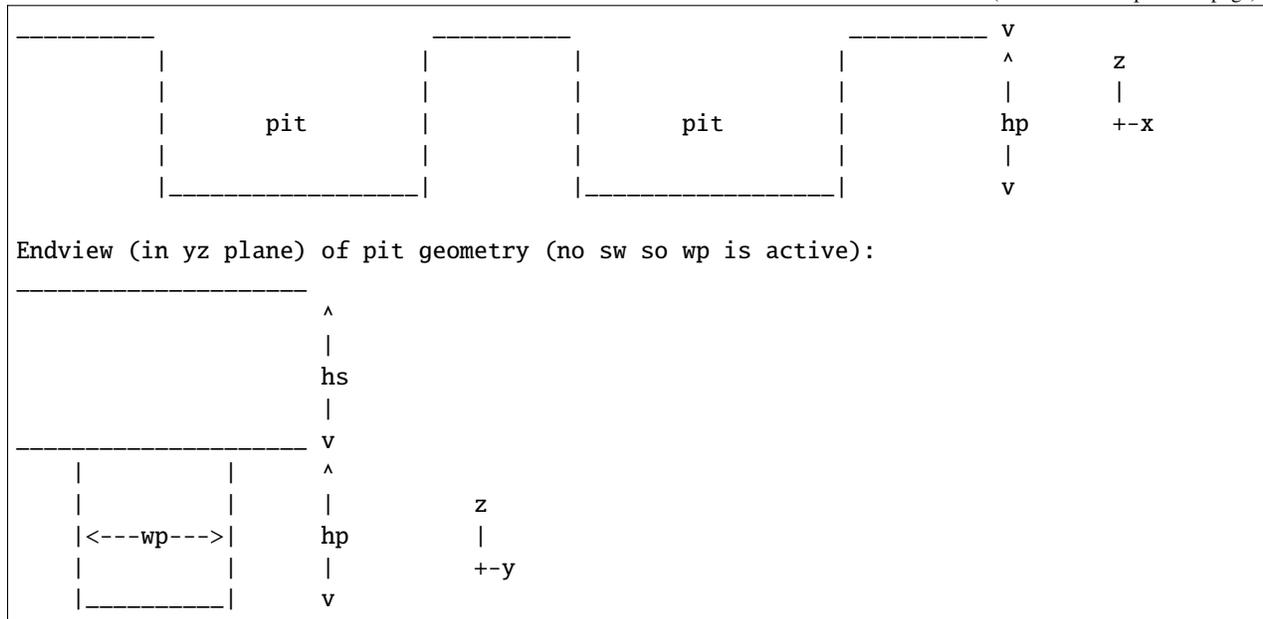
The *scaleGamma* keyword allows the user to scale the  $\gamma$  value by a factor, *gammaFactor*, for a given atom type. Setting *scaleGamma* to -1 for the corresponding particle type will explicitly take the limit of infinite particle mass in computing the force coupling for the fluid force (see note above).

If the *a0* keyword is used, the value specified is used for the square of the speed of sound in the fluid. If this keyword is not present, the speed of sound squared is set equal to  $\frac{1}{3} \left( \frac{dx_{LB}}{dt_{LB}} \right)^2$ . Setting  $a0 > \left( \frac{dx_{LB}}{dt_{LB}} \right)^2$  is not allowed, as this may lead to instabilities. As the speed of sound should usually be much larger than any fluid velocity of interest, its value does not normally have a significant impact on the results. As such, it is usually best to use the default for this option.

The *npits* keyword (followed by integer arguments: *npits*, *h\_p*, *l\_p*, *l\_pp*, *l\_e*) sets the fluid domain to the pits geometry. These arguments should only be used if you actually want something more complex than a rectangular/cubic geometry. The *npits* value sets the number of pits regions (arranged along x). The remaining arguments are sizes measured in multiples of *dx\_lb*: *h\_p* is the z-height of the pit regions, *l\_p* is the x-length of the pit regions, *l\_pp* is the length of the region between consecutive pits (referred to as a “slit” region), and *l\_e* is the x-length of the slit regions at each end of the channel. The pit geometry must fill the system in the x-direction but can be longer, in which case it is truncated (which enables asymmetric entrance/exit end sections). The additional *wp* keyword allows the width (in y-direction) of the pit to be specified (the default is full width) and the *sw* keyword indicates that there should be sidewalls in the y-direction (default is periodic in y-direction). These parameters are illustrated below:



(continued from previous page)



For further details, as well as descriptions and results of several test runs, see *Denniston et al.*. Please include a citation to this paper if the `lb_fluid` fix is used in work contributing to published research.

#### 17.80.4 Restart, fix\_modify, output, run start/stop, minimize info

Due to the large size of the fluid data, this fix writes its own binary restart files, if requested, independent of the main LAMMPS *binary restart files*; no information about `lb_fluid` is written to the main LAMMPS *binary restart files*.

None of the `fix_modify` options are relevant to this fix.

The fix computes a global scalar which can be accessed by various *output commands*. The scalar is the current temperature of the group of particles described by *group-ID* along with the fluid constrained to move with them. The temperature is computed via the kinetic energy of the group and fluid constrained to move with them and the total number of degrees of freedom (calculated internally). If the particles are not integrated independently (such as via *fix NVE*) but have additional constraints imposed on them (such as via integration using *fix rigid*) the degrees of freedom removed from these additional constraints will not be properly accounted for. In this case, the user can specify the total degrees of freedom independently using the *dof* keyword.

The fix also computes a global array of values which can be accessed by various *output commands*. There are 5 entries in the array. The first entry is the temperature of the fluid, the second entry is the total mass of the fluid plus particles, the third through fifth entries give the x, y, and z total momentum of the fluid plus particles.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.80.5 Restrictions

This fix is part of the LATBOLTZ package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix can only be used with an orthogonal simulation domain.

The boundary conditions for the fluid are specified independently to the particles. However, these should normally be specified consistently via the main LAMMPS *boundary* command (p p p, p p f, and p f f are the only consistent possibilities). Shrink-wrapped boundary conditions are not permitted with this fix.

This fix must be used before any of *fix lb/viscous* and *fix lb/momentum* as the fluid needs to be initialized before any of these routines try to access its properties. In addition, in order for the hydrodynamic forces to be added to the particles, this fix must be used in conjunction with the *lb/viscous* fix.

This fix needs to be used in conjunction with a standard LAMMPS integrator such as *fix NVE* or *fix rigid*.

## 17.80.6 Related commands

*fix lb/viscous*, *fix lb/momentum*

## 17.80.7 Default

$dx$  is chosen such that  $\frac{\tau}{dt_{LB}} = \frac{3\eta dt_{LB}}{\rho dx_{LB}^2}$  is approximately equal to 1.  $dm$  is set equal to 1.0.  $a0$  is set equal to  $\frac{1}{3} \left( \frac{dx_{LB}}{dt_{LB}} \right)^2$ . The trilinear stencil is used as the default interpolation method. The D3Q15 lattice is used for the lattice-Boltzmann algorithm.

---

**(Denniston et al.)** Denniston, C., Afrasiabian, N., Cole-Andre, M.G., Mackay, F. E., Ollila, S.T.T., and Whitehead, T., LAMMPS lb/fluid fix version 2: Improved Hydrodynamic Forces Implemented into LAMMPS through a lattice-Boltzmann fluid, Computer Physics Communications 275 (2022) 108318 .

**(Mackay and Denniston)** Mackay, F. E., and Denniston, C., Coupling MD particles to a lattice-Boltzmann fluid through the use of conservative forces, J. Comput. Phys. 237 (2013) 289-298.

**(Adhikari et al.)** Adhikari, R., Stratford, K., Cates, M. E., and Wagner, A. J., Fluctuating lattice Boltzmann, Europhys. Lett. 71 (2005) 473-479.

## 17.81 fix lb/momentum command

### 17.81.1 Syntax

```
fix ID group-ID lb/momentum nevery keyword values ...
```

- ID, group-ID are documented in the *fix* command
- lb/momentum = style name of this fix command
- nevery = adjust the momentum every this many timesteps
- zero or more keyword/value pairs may be appended
- keyword = *linear*

```
linear values = xflag yflag zflag  
xflag,yflag,zflag = 0/1 to exclude/include each dimension.
```

## 17.81.2 Examples

```
fix 1 sphere lb/momentum  
fix 1 all lb/momentum linear 1 1 0
```

## 17.81.3 Description

This fix is based on the *fix momentum* command, and was created to be used in place of that command, when a lattice-Boltzmann fluid is present.

Zero the total linear momentum of the system, including both the atoms specified by group-ID and the lattice-Boltzmann fluid every nevery timesteps. If there are no atoms specified by group-ID only the fluid momentum is affected. This is accomplished by adjusting the particle velocities and the fluid velocities at each lattice site.

---

**Note:** This fix only considers the linear momentum of the system.

---

By default, the subtraction is performed for each dimension. This can be changed by specifying the keyword *linear*, along with a set of three flags set to 0/1 in order to exclude/ include the corresponding dimension.

## 17.81.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.81.5 Restrictions

Can only be used if a lattice-Boltzmann fluid has been created via the *fix lb/fluid* command, and must come after this command.

This fix is part of the LATBOLTZ package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 17.81.6 Related commands

*fix momentum*, *fix lb/fluid*

### 17.81.7 Default

Zeros the total system linear momentum in each dimension.

## 17.82 `fix lb/viscous` command

### 17.82.1 Syntax

```
fix ID group-ID lb/viscous
```

- ID, group-ID are documented in *fix* command
- lb/viscous = style name of this fix command

### 17.82.2 Examples

```
fix 1 flow lb/viscous
```

### 17.82.3 Description

This fix is similar to the *fix viscous* command, and is to be used in place of that command when a lattice-Boltzmann fluid is present using the *fix lb/fluid*. This should be used in conjunction with one of the built-in LAMMPS integrators, such as *fix NVE* or *fix rigid*.

This fix adds a viscous force to each atom to cause it move with the same velocity as the fluid (an equal and opposite force is applied to the fluid via *fix lb/fluid*). When *fix lb/fluid* is called with the noise option, the atoms will also experience random forces which will thermalize them to the same temperature as the fluid. In this way, the combination of this fix with *fix lb/fluid* and a LAMMPS integrator like *fix NVE* is analogous to *fix langevin* except here the fluid is explicit. The temperature of the particles can be monitored via the scalar output of *fix lb/fluid*.

For details of this fix, as well as descriptions and results of several test runs, see *Denniston et al.*. Please include a citation to this paper if this fix is used in work contributing to published research.

### 17.82.4 Restart, `fix_modify`, output, run start/stop, minimize info

As described in the *fix viscous* documentation:

“No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command. This fix should only be used with damped dynamics minimizers that allow for non-conservative forces. See the *min\_style* command for details.”

### 17.82.5 Restrictions

This fix is part of the LATBOLTZ package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Can only be used if a lattice-Boltzmann fluid has been created via the *fix lb/fluid* command, and must come after this command.

### 17.82.6 Related commands

*fix lb/fluid*

### 17.82.7 Default

none

---

**(Denniston et al.)** Denniston, C., Afrasiabian, N., Cole-Andre, M.G., Mackay, F. E., Ollila, S.T.T., and Whitehead, T., LAMMPS lb/fluid fix version 2: Improved Hydrodynamic Forces Implemented into LAMMPS through a lattice-Boltzmann fluid, Computer Physics Communications 275 (2022) 108318 .

## 17.83 fix lineforce command

### 17.83.1 Syntax

```
fix ID group-ID lineforce x y z
```

- ID, group-ID are documented in *fix* command
- lineforce = style name of this fix command
- x y z = direction of line as a 3-vector

### 17.83.2 Examples

```
fix hold boundary lineforce 0.0 1.0 1.0
```

### 17.83.3 Description

Adjust the forces on each atom in the group so that only the component of force along the linear direction specified by the vector (x,y,z) remains. This is done by subtracting out components of force in the plane perpendicular to the line.

If the initial velocity of the atom is 0.0 (or along the line), then it should continue to move along the line thereafter.



### 17.83.4 Restart, `fix_modify`, `output`, `run start/stop`, `minimize` info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

### 17.83.5 Restrictions

none

### 17.83.6 Related commands

*fix plane-force*

### 17.83.7 Default

none

## 17.84 `fix manifoldforce` command

### 17.84.1 Syntax

```
fix ID group-ID manifoldforce manifold manifold-args ...
```

- ID, group-ID are documented in *fix* command
- manifold = name of the manifold
- manifold-args = parameters for the manifold

### 17.84.2 Examples

```
fix constrain all manifoldforce sphere 5.0
```

### 17.84.3 Description

This fix subtracts each time step from the force the component along the normal of the specified *manifold*. This can be used in combination with *minimize* to remove overlap between particles while keeping them (roughly) constrained to the given manifold, e.g. to set up a run with *fix nve/manifold/rattle*. I have found that only *hftn* and *quickmin* with a very small time step perform adequately though.

### 17.84.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is invoked during *energy minimization*.

---

### 17.84.5 Restrictions

This fix is part of the MANIFOLD package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Only use this with *min\_style hfqn* or *min\_style quickmin*. If not, the constraints will not be satisfied very well at all. A warning is generated if the *min\_style* is incompatible but no error.

---

### 17.84.6 Related commands

*fix nve/manifold/rattle*, *fix nvt/manifold/rattle*

## 17.85 fix mdi/qm command

### 17.85.1 Syntax

<code>fix ID group-ID mdi/qm keyword</code>
---

- ID, group-ID are documented in *fix* command
- mdi/qm = style name of this fix command
- zero or more keyword/value pairs may be appended
- keyword = *virial* or *add* or *every* or *connect* or *elements*

*virial* args = *yes* or *no*

*yes* = request virial tensor from server code

*no* = do not request virial tensor from server code

*add* args = *yes* or *no*

*yes* = add returned value from server code to LAMMPS quantities

*no* = do not add returned values to LAMMPS quantities

*every* args = *Nevery*

*Nevery* = request values from server code once every *Nevery* steps

*connect* args = *yes* or *no*

*yes* = perform a one-time connection to the MDI engine code

*no* = do not perform the connection operation

*elements* args = *N\_1 N\_2 ... N\_ntypes*

*N\_1, N\_2, ... N\_ntypes* = atomic number for each of *ntypes* LAMMPS atom types

## 17.85.2 Examples

```
fix 1 all mdi/qm
fix 1 all mdi/qm virial yes
fix 1 all mdi/qm add no every 100 elements 13 29
```

## 17.85.3 Description

This command enables LAMMPS to act as a client with another server code that will compute the total energy, per-atom forces, and total virial for atom conformations and simulation box size/shapes that LAMMPS sends it.

Typically the server code will be a quantum mechanics (QM) code, hence the name of the fix. However this is not required, the server code could be another classical molecular dynamics code or LAMMPS itself. The server code must support use of the [MDI Library](#) as explained below.

These are example use cases for this fix, discussed further below:

- perform an ab initio MD (AIMD) simulation with quantum forces
- perform an energy minimization with quantum forces
- perform a nudged elastic band (NEB) calculation with quantum forces
- perform a QM calculation for a series of independent systems which LAMMPS reads or generates

The code coupling performed by this command is done via the [MDI Library](#). LAMMPS runs as an MDI driver (client), and sends MDI commands to an external MDI engine code (server), e.g. a QM code which has support for MDI. See the [Howto mdi](#) page for more information about how LAMMPS can operate as either an MDI driver or engine.

The `examples/mdi` directory contains input scripts using this fix in the various use cases discussed below. In each case, two instances of LAMMPS are used, once as an MDI driver, once as an MDI engine (surrogate for a QM code). The `examples/mdi/README` file explains how to launch two codes so that they communicate via the MDI library using either MPI or sockets. Any QM code that supports MDI could be used in place of LAMMPS acting as a QM surrogate. See the [Howto mdi](#) page for a current list (March 2022) of such QM codes.

Note that an engine code can support MDI in either or both of two modes. It can be used as a stand-alone code, launched at the same time as LAMMPS. Or it can be used as a plugin library, which LAMMPS loads. See the [mdi plugin](#) command for how to trigger LAMMPS to load a plugin library. The `examples/mdi/README` file explains how to launch the two codes in either mode.

The *virial* keyword setting of *yes* or *no* determines whether LAMMPS will request the QM code to also compute and return a 6-element symmetric virial tensor for the system.

The *add* keyword setting of *yes* or *no* determines whether the energy and forces and virial returned by the QM code will be added to the LAMMPS internal energy and forces and virial or not. If the setting is *no* then the default [fix\\_modify energy](#) and [fix\\_modify virial](#) settings are also set to *no* and your input scripts should not set them to *yes*. See more details on these `fix_modify` settings below.

Whatever the setting for the *add* keyword, the QM energy, forces, and virial will be stored by the fix, so they can be accessed by other commands. See details below.

The *every* keyword determines how often the QM code will be invoked during a dynamics run with the current LAMMPS simulation box and configuration of atoms. The QM code will be called once every *Nevery* timesteps.

The *connect* keyword determines whether this fix performs a one-time connection to the QM code. The default is *yes*. The only time a *no* is needed is if this command is used multiple times in an input script. E.g. if it used inside a loop which also uses the [clear](#) command to destroy the system (including any defined fixes). See the `examples/mdi/in.series.driver` script as an example of this, where LAMMPS is using the QM code to compute energy and

forces for a series of system configurations. In this use case *connect no* is used along with the *mdi connect and exit* command to one-time initiate/terminate the connection outside the loop.

The *elements* keyword allows specification of what element each LAMMPS atom type corresponds to. This is specified by the atomic number of the element, e.g. 13 for Al. An atomic number must be specified for each of the ntypes LAMMPS atom types. Ntypes is typically specified via the *create\_box* command or in the data file read by the *read\_data* command.

If this keyword is specified, then this fix will send the MDI ">ELEMENTS" command to the engine, to insure the two codes are consistent in their definition of atomic species. If this keyword is not specified, then this fix will send the MDI >TYPES command to the engine. This is fine if both the LAMMPS driver and the MDI engine are initialized so that the atom type values are consistent in both codes.

---

The following 3 example use cases are illustrated in the examples/mdi directory. See its README file for more details.

(1) To run an ab initio MD (AIMD) dynamics simulation, or an energy minimization with QM forces, or a multi-replica NEB calculation, use *add yes* and *every 1* (the defaults). This is so that every time LAMMPS needs energy and forces, the QM code will be invoked.

Both LAMMPS and the QM code should define the same system (simulation box, atoms and their types) in their respective input scripts. Note that on this scenario, it may not be necessary for LAMMPS to define a pair style or use a neighbor list.

LAMMPS will then perform the timestepping or minimization iterations for the simulation. At the point in each timestep or iteration when LAMMPS needs the force on each atom, it communicates with the engine code. It sends the current simulation box size and shape (if they change dynamically, e.g. during an NPT simulation), and the current atom coordinates. The engine code computes quantum forces on each atom and the total energy of the system and returns them to LAMMPS.

Note that if the AIMD simulation is an NPT or NPH model, or the energy minimization includes *fix box relax* to equilibrate the box size/shape, then LAMMPS computes a pressure. This means the *virial* keyword should be set to *yes* so that the QM contribution to the pressure can be included.

(2) To run dynamics with a LAMMPS interatomic potential, and evaluate the QM energy and forces once every 1000 steps, use *add no* and *every 1000*. This could be useful for using an MD run to generate randomized configurations which are then passed to the QM code to produce training data for a machine learning potential. A *dump custom* command could be invoked every 1000 steps to dump the atom coordinates and QM forces to a file. Likewise the QM energy and virial could be output with the *thermo\_style custom* command.

(3) To do a QM evaluation of energy and forces for a series of  $N$  independent systems (simulation box and atoms), use *add no* and *every 1*. Write a LAMMPS input script which loops over the  $N$  systems. See the *Howto multiple* doc page for details on looping and removing old systems. The series of systems could be initialized by reading them from data files with *read\_data* commands. Or, for example, by using the *lattice*, *create\_atoms*, *delete\_atoms*, and/or *displace\_atoms random* commands to generate a series of different systems. At the end of the loop perform *run 0* and *write\_dump* commands to invoke the QM code and output the QM energy and forces. As in (2) this be useful to produce QM data for training a machine learning potential.

---

## 17.85.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy computed by the QM code to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy yes*, unless the *add* keyword is set to *no*, in which case the default setting is *no*.

The *fix\_modify virial* option is supported by this fix to add the contribution computed by the QM code to the global pressure of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify virial yes*, unless the *add* keyword is set to *no*, in which case the default setting is *no*.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the energy returned by the QM code. The scalar value calculated by this fix is “extensive”.

This fix also computes a global vector with of length 6 which contains the symmetric virial tensor values returned by the QM code. It can likewise be accessed by various *output commands*.

The ordering of values in the symmetric virial tensor is as follows: *vxx*, *vyy*, *vzz*, *vxy*, *vxz*, *vyz*. The values will be in pressure *units*.

This fix also computes a peratom array with 3 columns which contains the peratom forces returned by the QM code. It can likewise be accessed by various *output commands*.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

Assuming the *add* keyword is set to *yes* (the default), the forces computed by the QM code are used during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the potential energy associated with the QM forces to be included in the total potential energy of the system (the quantity being minimized), you **MUST** not disable the *fix\_modify energy* option for this fix, which means the *add* keyword should also be set to *yes* (the default).

---

## 17.85.5 Restrictions

This command is part of the MDI package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The QM code does not currently compute and return per-atom energy or per-atom virial contributions. So they will not show up as part of the calculations performed by the *compute pe/atom* or *compute stress/atom* commands.

To use LAMMPS as an MDI driver in conjunction with other MDI-enabled codes (MD or QM codes), the *units* command should be used to specify *real* or *metal* units. This will ensure the correct unit conversions between LAMMPS and MDI units. The other code will also perform similar unit conversions into its preferred units.

LAMMPS can also be used as an MDI driver in other unit choices it supports, e.g. *lj*, but then no unit conversion is performed.

## 17.85.6 Related commands

*mdi plugin, mdi engine*

## 17.85.7 Default

The default for the optional keywords are virial = no, add = yes, every = 1, connect = yes.

## 17.86 fix meso/move command

### 17.86.1 Syntax

```
fix ID group-ID meso/move style args keyword values ...
```

- ID, group-ID are documented in *fix* command
- meso/move = style name of this fix command
- style = *linear* or *wiggle* or *rotate* or *variable*

*linear* args = Vx Vy Vz

Vx,Vy,Vz = components of velocity vector (velocity units), any component can be specified as NULL

*wiggle* args = Ax Ay Az period

Ax,Ay,Az = components of amplitude vector (distance units), any component can be specified as NULL

period = period of oscillation (time units)

*rotate* args = Px Py Pz Rx Ry Rz period

Px,Py,Pz = origin point of axis of rotation (distance units)

Rx,Ry,Rz = axis of rotation vector

period = period of rotation (time units)

*variable* args = v\_dx v\_dy v\_dz v\_vx v\_vy v\_vz

v\_dx,v\_dy,v\_dz = 3 variable names that calculate x,y,z displacement as function of time, any component can be specified as NULL

v\_vx,v\_vy,v\_vz = 3 variable names that calculate x,y,z velocity as function of time, any component can be specified as NULL

- zero or more keyword/value pairs may be appended
- keyword = *units*

*units* value = *box* or *lattice*

### 17.86.2 Examples

```
fix 1 boundary meso/move wiggle 3.0 0.0 0.0 1.0 units box
fix 2 boundary meso/move rotate 0.0 0.0 0.0 0.0 0.0 1.0 5.0
fix 2 boundary meso/move variable v_myx v_myv NULL v_VX v_VY NULL
```

### 17.86.3 Description

Perform updates of position, velocity, internal energy and local density for mesoscopic particles in the group each timestep using the specified settings or formulas, without regard to forces on the particles. This can be useful for boundary, solid bodies or other particles, whose movement can influence nearby particles.

The operation of this fix is exactly like that described by the *fix move* command, except that particles' density, internal energy and extrapolated velocity are also updated.

---

**Note:** The particles affected by this fix should not be time integrated by other fixes (e.g. *fix sph*, *fix sph/stationary*), since that will change their positions and velocities twice.

---



---

**Note:** As particles move due to this fix, they will pass through periodic boundaries and be remapped to the other side of the simulation box, just as they would during normal time integration (e.g. via the *fix sph* command). It is up to you to decide whether periodic boundaries are appropriate with the kind of particle motion you are prescribing with this fix.

---



---

**Note:** As discussed below, particles are moved relative to their initial position at the time the fix is specified. These initial coordinates are stored by the fix in “unwrapped” form, by using the image flags associated with each particle. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each particle. You can reset the image flags (e.g. to 0) before invoking this fix by using the *set image* command.

---

The *linear* style moves particles at a constant velocity, so that their position  $X = (x,y,z)$  as a function of time is given in vector notation as

$$X(t) = X0 + V * \text{delta}$$

where  $X0 = (x0,y0,z0)$  is their position at the time the fix is specified,  $V$  is the specified velocity vector with components  $(Vx,Vy,Vz)$ , and *delta* is the time elapsed since the fix was specified. This style also sets the velocity of each particle to  $V = (Vx,Vy,Vz)$ . If any of the velocity components is specified as NULL, then the position and velocity of that component is time integrated the same as the *fix sph* command would perform, using the corresponding force component on the particle.

Note that the *linear* style is identical to using the *variable* style with an *equal-style variable* that uses the *vdisplace()* function. E.g.

```
variable V equal 10.0
variable x equal vdisplace(0.0,$V)
fix 1 boundary move variable v_x NULL NULL v_V NULL NULL
```

The *wiggle* style moves particles in an oscillatory fashion, so that their position  $X = (x,y,z)$  as a function of time is given in vector notation as

$$X(t) = X0 + A \sin(\omega * \text{delta})$$

where  $X0 = (x0,y0,z0)$  is their position at the time the fix is specified,  $A$  is the specified amplitude vector with components  $(Ax,Ay,Az)$ ,  $\omega$  is  $2 \text{ PI} / \text{period}$ , and *delta* is the time elapsed since the fix was specified. This style also sets the velocity of each particle to the time derivative of this expression. If any of the amplitude components is specified as NULL, then the position and velocity of that component is time integrated the same as the *fix sph* command would perform, using the corresponding force component on the particle.

Note that the *wiggle* style is identical to using the *variable* style with *equal-style variables* that use the `swiggle()` and `cwiggle()` functions. E.g.

```
variable A equal 10.0
variable T equal 5.0
variable omega equal 2.0*PI/$T
variable x equal swiggle(0.0,$A,$T)
variable v equal v_omega*($A-cwiggle(0.0,$A,$T))
fix 1 boundary move variable v_x NULL NULL v_v NULL NULL
```

The *rotate* style rotates particles around a rotation axis  $R = (R_x, R_y, R_z)$  that goes through a point  $P = (P_x, P_y, P_z)$ . The *period* of the rotation is also specified. The direction of rotation for the particles around the rotation axis is consistent with the right-hand rule: if your right-hand thumb points along  $R$ , then your fingers wrap around the axis in the direction of rotation.

This style also sets the velocity of each particle to  $(\omega \text{ cross } R_{\text{perp}})$  where  $\omega$  is its angular velocity around the rotation axis and  $R_{\text{perp}}$  is a perpendicular vector from the rotation axis to the particle.

The *variable* style allows the position and velocity components of each particle to be set by formulas specified via the *variable* command. Each of the 6 variables is specified as an argument to the `fix` as `v_name`, where `name` is the variable name that is defined elsewhere in the input script.

Each variable must be of either the *equal* or *atom* style. *Equal*-style variables compute a single numeric quantity, that can be a function of the timestep as well as of other simulation values. *Atom*-style variables compute a numeric quantity for each particle, that can be a function per-atom quantities, such as the particle's position, as well as of the timestep and other simulation values. Note that this `fix` stores the original coordinates of each particle (see note below) so that per-atom quantity can be used in an atom-style variable formula. See the *variable* command for details.

The first 3 variables (`v_dx`, `v_dy`, `v_dz`) specified for the *variable* style are used to calculate a displacement from the particle's original position at the time the `fix` was specified. The second 3 variables (`v_vx`, `v_vy`, `v_vz`) specified are used to compute a velocity for each particle.

Any of the 6 variables can be specified as `NULL`. If both the displacement and velocity variables for a particular  $x, y, z$  component are specified as `NULL`, then the position and velocity of that component is time integrated the same as the *fix sph* command would perform, using the corresponding force component on the particle. If only the velocity variable for a component is specified as `NULL`, then the displacement variable will be used to set the position of the particle, and its velocity component will not be changed. If only the displacement variable for a component is specified as `NULL`, then the velocity variable will be used to set the velocity of the particle, and the position of the particle will be time integrated using that velocity.

The *units* keyword determines the meaning of the distance units used to define the *linear* velocity and *wiggle* amplitude and *rotate* origin. This setting is ignored for the *variable* style. A *box* value selects standard units as defined by the *units* command, e.g. velocity in Angstroms/fs and amplitude and position in Angstroms for `units = real`. A *lattice* value means the velocity units are in lattice spacings per time and the amplitude and position are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing. Each of these 3 quantities may be dependent on the  $x, y, z$  dimension, since the lattice spacings can be different in  $x, y, z$ .



## 17.86.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the original coordinates of moving particles to *binary restart files*, as well as the initial timestep, so that the motion can be continuous in a restarted simulation. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

---

**Note:** Because the move positions are a function of the current timestep and the initial timestep, you cannot reset the timestep to a different value after reading a restart file, if you expect a fix move command to work in an uninterrupted fashion.

---

None of the *fix\_modify* options are relevant to this fix.

This fix produces a per-atom array which can be accessed by various *output commands*. The number of columns for each atom is 3, and the columns store the original unwrapped x,y,z coords of each particle. The per-atom values can be accessed on any timestep.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

This fix is not invoked during *energy minimization*.

## 17.86.5 Restrictions

This fix is part of the DPD-SMOOTH package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store density and internal energy as defined by the *atom\_style sph* command.

All particles in the group must be mesoscopic SPH/SDPD particles.

## 17.86.6 Related commands

*fix move, fix sph, displace\_atoms*

## 17.86.7 Default

The option default is units = lattice.

## 17.87 fix\_modify AtC commands

### 17.87.1 fix\_modify AtC add\_molecule command

#### Syntax

```
fix_modify <AtC fixID> add_molecule <small|large> <tag> <group-ID>
```

- AtC fixID = ID of *fix atc* instance
- add\_molecule = name of the AtC sub-command
- *small* or *large* = can be *small* if molecule size < cutoff radius, must be *large* otherwise
- tag = tag for tracking a molecule

- *group-ID* = LAMMPS defined group-ID

## Examples

```
group WATERGROUP type 1 2
fix_modify AtC add_molecule small water WATERGROUP
```

## Description

Associates a tag with all molecules corresponding to a specified group.

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC add\_species*
- *fix\_modify AtC remove\_species*
- *fix\_modify AtC remove\_molecule*

## Default

None.

## 17.87.2 fix\_modify AtC add\_species command

### Syntax

```
fix_modify <AtC fixID> add_species <tag> <group|type> <ID>
```

- AtC fixID = ID of *fix atc* instance
- add\_species = name of the AtC sub-command
- tag = tag for tracking a species
- *group* or *type* = LAMMPS defined group or type of atoms
- ID = name of group or type number

## Examples

```
fix_modify AtC add_species gold type 1
group GOLDGROUP type 1
fix_modify AtC add_species gold group GOLDGROUP
```

## Description

Associates a tag with all atoms of a specified type or within a specified group.

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC add\_molecule*
- *fix\_modify AtC remove\_species*
- *fix\_modify AtC remove\_molecule*

## Default

None.

### 17.87.3 fix\_modify AtC atom\_element\_map command

#### Syntax

```
fix_modify <AtC fixID> atom_element_map <eulerian|lagrangian> [<frequency>]
```

- AtC fixID = ID of *fix atc* instance
- atom\_element\_map = name of the AtC sub-command
- *eulerian* or *lagrangian* = frame of reference
- frequency = frequency of updating atom-to-continuum maps based on the current configuration - (only for eulerian)

## Examples

```
fix_modify AtC atom_element_map eulerian 100
```

## Description

Changes frame of reference from *eulerian* to *lagrangian* or vice versa and sets the frequency for which the map from atoms to elements is reformed and all the attendant data is recalculated.

## Restrictions

Cannot change map type after initialization.

## Related AtC commands

- [fix\\_modify AtC command overview](#)

## Default

*lagrangian*

## 17.87.4 fix\_modify AtC atom\_weight command

### Syntax

```
fix_modify <AtC fixID> atom_weight <method> <args>
```

- AtC fixID = ID of *fix atc* instance
- atom\_weight = name of the AtC sub-command
- method = *constant* or *lattice* or *element* or *region* or *group* or *read\_in*
  - *constant* <group-ID> <value>: atoms in specified group are assigned the constant value given
  - *lattice*: volume per atom for specified lattice type (e.g. fcc) and parameter
  - *element*: element volume divided among atoms within element
  - *region*: volume per atom determined based on the atom count in the MD regions and their volumes. Note: meaningful only if atoms completely fill all the regions.
  - *group*: volume per atom determined based on the atom count in a group and its volume
  - *node*: (undocumented)
  - *node\_element*: (undocumented)
  - *read\_in*<filename>: list of values for atoms are read-in from specified file

## Examples

```
fix_modify AtC atom_weight constant myatoms 11.8
fix_modify AtC atom_weight lattice
fix_modify AtC atom_weight read-in atm_wt_file.txt
```

## Description

Command for assigning the value of atomic weights used for atomic integration in atom-continuum coupled simulations.

## Restrictions

The use of the lattice option requires a lattice type and parameter is already specified.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

*lattice*

## 17.87.5 fix\_modify AtC atomic\_charge command

### Syntax

```
fix_modify <AtC fixID> <include|omit> atomic_charge
```

- AtC fixID = ID of *fix atc* instance
- *include* or *omit* = switch to activate/deactivate inclusion of intrinsic atomic charge in ATC
- atomic\_charge = name of the AtC sub-command

## Examples

```
fix_modify AtC include atomic_charge
```

## Description

Determines whether AtC tracks the total charge as a finite element field.

## Restrictions

Required for: *electrostatics*

## Related AtC commands

- *fix\_modify* AtC command overview

## Default

If the atom charge is defined, default is on, otherwise default is off.

## 17.87.6 *fix\_modify* AtC *boundary\_dynamics* command

### Syntax

```
fix_modify <AtC fixID> boundary_dynamics <on|damped_harmonic|prescribed|coupled|none>
```

- AtC fixID = ID of *fix atc* instance
- *boundary\_dynamics* = name of the AtC sub-command
- *on* or *damped\_harmonic* *prescribed* *coupled* *none*

## Description

Sets different schemes for controlling boundary atoms. *on* will integrate the boundary atoms using the velocity-verlet algorithm. *damped\_harmonic* uses a mass/spring/dashpot for the boundary atoms with added arguments of the damping and spring constants followed by the ratio of the boundary type mass to the desired mass. *prescribed* forces the boundary atoms to follow the finite element displacement. *coupled* does the same.

## Restrictions

Boundary atoms must be specified. When using swaps between internal and boundary atoms, the initial configuration must have already correctly partitioned the two.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

*prescribed on*

## 17.87.7 fix\_modify AtC boundary\_faceset command

### Syntax

```
fix_modify <AtC fixID> boundary_faceset <is|add> <faceset_name>
```

- AtC fixID = ID of *fix atc* instance
- boundary\_faceset = name of the AtC sub-command
- *is* or *add* = select whether to select or add a faceset
- faceset\_name = name of the faceset

### Examples

```
fix_modify AtC boundary_faceset is obndy
```

### Description

This command species the faceset name when using a faceset to compute the MD/FE boundary fluxes. The faceset must already exist.

### Restrictions

This is only valid when *fe\_md\_boundary* is set to *faceset*.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC fe\_md\_boundary*
- *fix\_modify AtC mesh create\_faceset box*
- *fix\_modify AtC mesh create\_faceset plane*

**Default**

None.

**17.87.8 fix\_modify AtC boundary type command****Syntax**

```
fix_modify <AtC fixID> boundary type <atom-type-id>
```

- AtC fixID = ID of *fix atc* instance
- boundary type = name of the AtC sub-command
- atom-type-id = type id for atoms that represent a fictitious boundary internal to the FE mesh

**Examples**

```
fix_modify AtC boundary type ghost_atoms
```

**Description**

Command to define the atoms that represent the fictitious boundary internal to the FE mesh. For fully overlapped MD/FE domains with periodic boundary conditions no boundary atoms should be defined.

**Restrictions**

None.

**Related AtC commands**

- *fix\_modify AtC command overview*

**Default**

None.

**17.87.9 fix\_modify AtC consistent\_fe\_initialization command****Syntax**

```
fix_modify <AtC fixID> consistent_fe_initialization <on|off>
```

- AtC fixID = ID of *fix atc* instance
- consistent\_fe\_initialization = name of the AtC sub-command
- *on* or *off* = switch to activate/deactivate the initial setting of the FE intrinsic field to match the projected MD field



## Examples

```
fix_modify AtC consistent_fe_initialization on
```

## Description

Determines whether AtC initializes FE intrinsic fields (e.g., temperature) to match the projected MD values. This is particularly useful for fully overlapping simulations.

## Restrictions

Can be used with: *thermal*, *two\_temperature*. Cannot be used with time filtering on. Does not include boundary nodes.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

Default is *off*

## 17.87.10 fix\_modify AtC control localized\_lambda command

### Syntax

```
fix_modify <AtC fixID> control localized_lambda <on|off>
```

- AtC fixID = ID of *fix atc* instance
- control localized\_lambda = name of the AtC sub-command
- *on* or *off* = Toggles state of localization algorithm

## Examples

```
fix_modify AtC control localized_lambda on
```

## Description

Turns the localization algorithms *on* or *off* for control algorithms to restrict the influence of FE coupling or boundary conditions to a region near the boundary of the MD region. Control algorithms will not affect atoms in elements not possessing faces on the boundary of the region. Flux-based control is localized via row-sum lumping while quantity control is done by solving a truncated matrix equation.

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

off

## 17.87.11 fix\_modify AtC control momentum command

### Syntax

```
fix_modify <AtC fixID> control <physics_type> <solution_parameter> <value>
fix_modify AtC control momentum none
fix_modify AtC control momentum rescale <frequency>
fix_modify AtC control momentum glc_displacement
fix_modify AtC control momentum glc_velocity
fix_modify AtC control momentum hoover
fix_modify AtC control momentum flux [faceset face_set_id, interpolate]
```

- AtC fixID = ID of *fix atc* instance
- control = name of the AtC sub-command
- physics\_type = *thermal* or *momentum*
- solution\_parameter = *max\_iterations* or *tolerance*
- value = solution\_parameter value
- momentum option = *none* or *rescale* or *glc\_displacement* or *glc\_velocity* *hoover* or *flux*
- frequency = time step frequency for applying displacement and velocity rescaling
- faceset\_id = id of boundary face set (optional, only for *faceset*)

### Examples

```
fix_modify AtC control momentum none
fix_modify AtC control momentum flux faceset bndy_faces
fix_modify AtC control momentum glc_velocity
```

## Description

The general version of *control* sets the numerical parameters for the matrix solvers used in the specified control algorithm. Many solution approaches require iterative solvers, and these methods enable users to provide the maximum number of iterations and the relative tolerance.

The *control momentum* version sets the momentum exchange mechanism from the finite elements to the atoms, managed through a control algorithm. *rescale* computes a scale factor for each atom to match the finite element temperature. *hoover* is a Gaussian least-constraint isokinetic thermostat enforces that the nodal restricted atomic temperature matches the finite element temperature. *flux* is a similar mode, but rather adds energy to the atoms based on conservation of energy.

*correction\_max\_iterations* sets the maximum number of iterations to compute the second order in time correction term for lambda with the fractional step method. The method uses the same tolerance as the controller's matrix solver.

## Restrictions

Only for be used with the specific controllers *thermal* or *momentum*. They are ignored if a lumped solution is requested.

*control momentum* is only for be used with specific transfers: elastic *rescale* not valid with time filtering activated

## Related AtC commands

- *fix\_modify* AtC command overview
- *fix\_modify* AtC control thermal

## Default

- *max\_iterations* is the number of rows in the matrix.
- *tolerance* is 1.0e-10.

## 17.87.12 fix\_modify AtC control thermal command

### Syntax

```
fix_modify <AtC fixID> control <physics_type> <solution_parameter> <value>
fix_modify <AtC fixID> control thermal <control_type> <optional_args>
fix_modify <AtC fixID> control thermal rescale <frequency>
fix_modify <AtC fixID> control thermal flux <boundary_integration_type> <faceset_id>
fix_modify <AtC fixID> control thermal correction_max_iterations <max_iterations>
```

- AtC fixID = ID of *fix atc* instance
- control = name of the AtC sub-command
- physics\_type = *thermal* or *momentum*
- solution\_parameter = *max\_iterations* or *tolerance*
- value = solution\_parameter value
- thermal control\_type = *none* or *rescale* or *hoover* or *flux*
- frequency = time step frequency for applying velocity rescaling

- `boundary_integration_type` = *faceset* or *interpolate* (optional)
- `faceset_id` = id of boundary face set (optional, only for *faceset*)
- `correction_max_iterations` = maximum number of iterations that will be used by iterative matrix solvers for *thermal* physics type

## Examples

```
fix_modify AtC control thermal none
fix_modify AtC control thermal rescale 10
fix_modify AtC control thermal hoover
fix_modify AtC control thermal flux
fix_modify AtC control thermal flux faceset bndy_faces
fix_modify AtC control thermal correction_max_iterations 10
```

## Description

The general version of *control* sets the numerical parameters for the matrix solvers used in the specified control algorithm. Many solution approaches require iterative solvers, and these methods enable users to provide the maximum number of iterations and the relative tolerance.

The *control thermal* version sets the energy exchange mechanism from the finite elements to the atoms, managed through a control algorithm. *rescale* computes a scale factor for each atom to match the finite element temperature. *hoover* is a Gaussian least-constraint isokinetic thermostat enforces that the nodal restricted atomic temperature matches the finite element temperature. *flux* is a similar mode, but rather adds energy to the atoms based on conservation of energy. *hoover* and *flux* allow the prescription of sources or fixed temperatures on the atoms.

*correction\_max\_iterations* sets the maximum number of iterations to compute the second order in time correction term for lambda with the fractional step method. The method uses the same tolerance as the controller's matrix solver.

## Restrictions

Only for be used with the specific controllers *thermal* or *momentum*. They are ignored if a lumped solution is requested.

*control thermal* is only for be used with specific transfers: *thermal (rescale, hoover, flux)*, *two\_temperature (flux)*. *rescale* not valid with time filtering activated

*correction\_max\_iterations* is only for use with *thermal* physics using the fractional step method.

## Related AtC commands

- [fix\\_modify AtC command overview](#)
- [fix\\_modify AtC control momentum](#)

## Default

- *max\_iterations* is the number of rows in the matrix.
- *tolerance* is 1.0e-10.
- *rescale* frequency is 1
- *flux* boundary\_integration\_type is *interpolate*
- *correction\_max\_iterations* is 20

## 17.87.13 fix\_modify AtC decomposition command

### Syntax

```
fix_modify <AtC fixID> decomposition <type>
```

- AtC fixID = ID of *fix atc* instance
- decomposition = name of the AtC sub-command
- type = *replicated\_memory* or *distributed\_memory*

### Examples

```
fix_modify AtC decomposition distributed_memory
```

### Description

Command for assigning the distribution of work and memory for parallel runs. With *replicated\_memory* the nodal information is replicated on each processor, and with *distributed\_memory* only the owned nodal information kept on each processor. The *replicated\_memory* option is most appropriate for simulations where the number of nodes is much smaller than the number of atoms.

### Restrictions

None.

### Related AtC commands

- *fix\_modify AtC command overview*

## Default

replicated\_memory

## 17.87.14 fix\_modify AtC extrinsic electron\_integration command

### Syntax

```
fix_modify <AtC fixID> extrinsic electron_integration <integration_type> [<num_subcycle_
↪steps>]
```

- AtC fixID = ID of *fix atc* instance
- extrinsic electron\_integration = name of the AtC sub-command
- integration\_type = *explicit* or *implicit* or *steady*
- num\_subcycle\_steps = number of subcycle steps for the electron time integration (optional)

### Examples

```
fix_modify AtC extrinsic electron_integration implicit
fix_modify AtC extrinsic electron_integration explicit 100
```

### Description

Switches between integration schemes for the electron temperature. The number of subcycling steps used to integrate the electron temperature for one LAMMPS timestep can be manually adjusted to capture fast electron dynamics.

### Restrictions

For use only with the two\_temperature type of the AtC fix (see *fix atc* command)

### Related AtC commands

- *fix\_modify AtC command overview*

### Default

*implicit* and *subcycle\_steps* = 1

## 17.87.15 `fix_modify AtC equilibrium_start` command

### Syntax

```
fix_modify <AtC fixID> equilibrium_start <on|off>
```

- AtC fixID = ID of *fix atc* instance
- equilibrium\_start = name of the AtC sub-command
- *exponential* or *step* or *no\_filter* = select type of filter

### Examples

```
fix_modify AtC equilibrium_start on
```

### Description

Starts filtered calculations assuming they start in equilibrium, i.e. perfect finite element force balance.

### Restrictions

Only for use with these specific transfers: thermal, two\_temperature

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC filter*
- *fix\_modify AtC filter scale*

### Default

None.

## 17.87.16 `fix_modify AtC extrinsic exchange` command

### Syntax

```
fix_modify <AtC fixID> extrinsic exchange <on|off>
```

- AtC fixID = ID of *fix atc* instance
- extrinsic exchange = name of the AtC sub-command
- *on* or *off* = set state of energy exchange

## Examples

```
fix_modify AtC extrinsic exchange on
```

## Description

Switches energy exchange between the MD system and the electron system on or off

## Restrictions

For use only with the two\_temperature type of the AtC fix (see *fix atc* command)

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

*on*

## 17.87.17 fix\_modify AtC fe\_md\_boundary command

### Syntax

```
fix_modify <AtC fixID> fe_md_boundary <faceset|interpolate|no_boundary>
```

- AtC fixID = ID of *fix atc* instance
- fe\_md\_boundary = name of the AtC sub-command
- *faceset* or *interpolate* or *no\_boundary*

## Examples

```
fix_modify AtC fe_md_boundary interpolate
```

## Description

Specifies different methods for computing fluxes between between the MD and FE integration regions. Faceset defines a faceset separating the MD and FE regions and uses finite element face quadrature to compute the flux. Interpolate uses a reconstruction scheme to approximate the flux, which is more robust but less accurate if the MD/FE boundary does correspond to a faceset. No boundary results in no fluxes between the systems being computed.



## Restrictions

If *faceset* is used, all the AtC non-boundary atoms must lie within and completely fill the domain enclosed by the faceset.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC mesh create\_faceset box*
- *fix\_modify AtC mesh create\_faceset plane*

## Default

*interpolate*

## 17.87.18 fix\_modify AtC filter scale command

### Syntax

```
fix_modify <AtC fixID> filter scale <scale>
```

- AtC fixID = ID of *fix atc* instance
- filter scale = name of the AtC sub-command
- scale = characteristic times scale of the filter

### Examples

```
fix_modify AtC filter scale 10.0
```

### Description

Sets the time scale for MD dynamics filter to construct a more appropriate continuous field.

### Restrictions

Only for use with these specific transfers: thermal, two\_temperature

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC filter*
- *fix\_modify AtC filter type*

### Default

0.0

## 17.87.19 fix\_modify AtC filter type command

### Syntax

```
fix_modify <AtC fixID> filter type <exponential|step|no_filter>
```

- AtC fixID = ID of *fix atc* instance
- filter type = name of the AtC sub-command
- *exponential* or *step* or *no\_filter* = select type of filter

### Examples

```
fix_modify AtC filter type exponential
```

### Description

Specifies the type of time filter used.

### Restrictions

Only for use with these specific transfers: thermal, two\_temperature

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC filter*
- *fix\_modify AtC filter scale*

**Default**

None.

**17.87.20 fix\_modify AtC fix command****Syntax**

```
fix_modify <AtC fixID> fix <field> <nodeset> <constant|function>
```

- AtC fixID = ID of *fix atc* instance
- fix = name of the AtC sub-command
- field = field kind name valid for type of physics: *temperature* or *electron\_temperature*
- nodeset = name of set of nodes to apply boundary condition
- *constant* or *function* = value or name of function followed by its parameters

**Examples**

```
fix_modify AtC fix temperature groupNAME 10.
fix_modify AtC fix temperature groupNAME 0 0 0 10.0 0 0 1.0
```

**Description**

Creates a constraint on the values of the specified field at specified nodes.

**Restrictions**

The keyword *all* is reserved and thus not available as nodeset name.

**Related AtC commands**

- [fix\\_modify AtC command overview](#)
- [fix\\_modify AtC unfix](#)

**Default**

None.

### 17.87.21 `fix_modify AtC fix_flux` command

#### Syntax

```
fix_modify <AtC fixID> fix_flux <field> <face_set> <value|function>
```

- AtC fixID = ID of *fix atc* instance
- fix\_flux = name of the AtC sub-command
- field = field kind name valid for type of physics: *temperature* or *electron\_temperature*
- face\_set = name of set of element faces
- *value* or *function* = value or name of function followed by its parameters

#### Examples

```
fix_modify AtC fix_flux temperature faceSet 10.0
```

#### Description

Command for fixing normal fluxes e.g. `heat_flux`. This command only prescribes the normal component of the physical flux, e.g. heat (energy) flux. The units are in AtC units, i.e. derived from the LAMMPS length, time, and mass scales.

#### Restrictions

Only normal fluxes (Neumann data) can be prescribed.

#### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC unfix\_flux*

#### Default

None.

### 17.87.22 `fix_modify AtC computes` command

#### Syntax

```
fix_modify <AtC fixID> computes <add|delete> <per-atom compute-ID> <volume|number>
```

- AtC fixID = ID of *fix atc* instance
- computes = name of the AtC sub-command
- *add* or *delete* = add or delete the calculation of an equivalent continuum field for the specified per-atom compute as volume or number density quantity

- per-atom compute-ID = ID of a per-atom compute; fields can be calculated for all per-atom computes available in LAMMPS
- *volume* or *number* = select whether the created field is a per-unit-volume quantity or a per-atom quantity as weighted by kernel functions

## Examples

```
compute virial all stress/atom
fix_modify AtC computes add virial volume
fix_modify AtC computes delete virial

compute centrosymmetry all centro/atom
fix_modify AtC computes add centrosymmetry number
```

## Description

Calculates continuum fields corresponding to specified per-atom *computes* created by LAMMPS.

## Restrictions

Must be used with *fix atc hardy*. The per-atom compute must be specified before the corresponding continuum field can be requested.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC fields*
- *compute*

## Default

None.

## 17.87.23 fix\_modify AtC fields command

### Syntax

```
fix_modify <AtC fixID> fields <all|none>
fix_modify <AtC fixID> fields <add|delete> <list_of_fields>
```

- AtC fixID = ID of *fix atc* instance
- fields = name of the AtC sub-command
- *all* or *none* = output all or no fields
- *add* or *delete* = add or delete the listed output fields
- list\_of\_fields = one or more of the fields listed below:

- density : mass per unit volume
- displacement : displacement vector
- momentum : momentum per unit volume
- velocity : defined by momentum divided by density
- projected\_velocity : simple kernel estimation of atomic velocities
- temperature : temperature derived from the relative atomic kinetic energy
- kinetic\_temperature : temperature derived from the full kinetic energy
- number\_density : simple kernel estimation of number of atoms per unit volume
- stress : Cauchy stress tensor for eulerian analysis (atom\_element\_map), or first Piola-Kirchhoff stress tensor for lagrangian analysis
- transformed\_stress : first Piola-Kirchhoff stress tensor for eulerian analysis (atom\_element\_map), or Cauchy stress tensor for lagrangian analysis
- heat\_flux : spatial heat flux vector for eulerian, or referential heat flux vector for lagrangian
- potential\_energy : potential energy per unit volume
- kinetic\_energy : kinetic energy per unit volume
- thermal\_energy : thermal energy (kinetic energy - continuum kinetic energy) per unit volume
- internal\_energy : total internal energy (potential + thermal) per unit volume
- energy : total energy (potential + kinetic) per unit volume
- number\_density : number of atoms per unit volume
- eshelby\_stress : configurational stress (energy-momentum) tensor defined by [Eshelby]
- vacancy\_concentration : volume fraction of vacancy content
- type\_concentration : volume fraction of a specific atom type

## Examples

```
fix_modify AtC fields add velocity temperature
```

## Description

Allows modification of the fields calculated and output by the AtC transfer class. The commands are cumulative, e.g.:

```
fix_modify AtC fields none  
fix_modify AtC fields add velocity temperature
```

will only output the velocity and temperature fields.

## Restrictions

Must be used with *fix atc hardy*. Currently, the stress and heat flux formulas are only correct for central force potentials, e.g. Lennard-Jones and EAM but not Stillinger-Weber.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC gradients*
- *fix\_modify AtC rates*
- *fix\_modify AtC computes*

## Default

By default, no fields are output.

## References

### 17.87.24 *fix\_modify* AtC gradients command

#### Syntax

```
fix_modify <AtC fixID> gradients <add|delete> <list_of_fields>
```

- AtC fixID = ID of *fix atc* instance
- gradients = name of the AtC sub-command
- *add* or *delete* = select whether to add or delete calculation of gradients for the listed output fields
- list\_of\_fields = one or more of the fields listed below:
  - density : mass per unit volume
  - displacement : displacement vector
  - momentum : momentum per unit volume
  - velocity : defined by momentum divided by density
  - projected\_velocity : simple kernel estimation of atomic velocities
  - temperature : temperature derived from the relative atomic kinetic energy
  - kinetic\_temperature : temperature derived from the full kinetic energy
  - number\_density : simple kernel estimation of number of atoms per unit volume
  - stress : Cauchy stress tensor for eulerian analysis (atom\_element\_map), or first Piola-Kirchhoff stress tensor for lagrangian analysis
  - transformed\_stress : first Piola-Kirchhoff stress tensor for eulerian analysis (atom\_element\_map), or Cauchy stress tensor for lagrangian analysis
  - heat\_flux : spatial heat flux vector for eulerian, or referential heat flux vector for lagrangian
  - potential\_energy : potential energy per unit volume

- `kinetic_energy` : kinetic energy per unit volume
- `thermal_energy` : thermal energy (kinetic energy - continuum kinetic energy) per unit volume
- `internal_energy` : total internal energy (potential + thermal) per unit volume
- `energy` : total energy (potential + kinetic) per unit volume
- `number_density` : number of atoms per unit volume
- `eshelby_stress` : configurational stress (energy-momentum) tensor defined by [Eshelby]
- `vacancy_concentration` : volume fraction of vacancy content
- `type_concentration` : volume fraction of a specific atom type

## Examples

```
fix_modify AtC gradients add temperature velocity stress  
fix_modify AtC gradients delete velocity
```

## Description

Requests calculation and output of gradients of the fields from the AtC transfer class. These gradients will be with regard to spatial or material coordinate for Eulerian or Lagrangian analysis, respectively, as specified by *fix\_modify AtC atom\_element\_map*

## Restrictions

Must be used with *fix atc hardy*.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC atom\_element\_map*
- *fix\_modify AtC fields*
- *fix\_modify AtC rates*

## Default

None.



## 17.87.25 fix\_modify AtC kernel command

### Syntax

```
fix_modify <AtC fixID> kernel <type> <parameters>
```

- AtC fixID = ID of *fix atc* instance
- kernel = name of the AtC sub-command
- type = *step* or *cell* or *cubic\_bar* or *cubic\_cylinder* or *cubic\_sphere* or *quartic\_bar* or *quartic\_cylinder* or *quartic\_sphere*
- the following parameter(s) are required for each kernel:
  - *step* : <radius>
  - *cell* : <hx> <hy> <hz> or <h>
  - *cubic\_bar* : <half\_width>
  - *cubic\_cylinder* : <radius>
  - *cubic\_sphere* : <radius>
  - *quartic\_bar* : <half\_width>
  - *quartic\_cylinder* : <radius>
  - *quartic\_sphere* : <radius>

### Examples

```
fix_modify AtC kernel cell 1.0 1.0 1.0
fix_modify AtC kernel quartic_sphere 10.0
```

### Description

Sets the localization kernel type and parameters for *fix atc hardy*.

### Restrictions

Must be used with *fix atc hardy*. For bar kernel types, half-width oriented along x-direction. For cylinder kernel types, cylindrical axis is assumed to be in z-direction.

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC fields*
- *fix\_modify AtC gradients*
- *fix\_modify AtC rates*
- *fix\_modify AtC computes*

## Default

None.

## 17.87.26 `fix_modify AtC on_the_fly` command

### Syntax

```
fix_modify <AtC fixID> on_the_fly <bond|kernel> <on|off>
```

- AtC fixID = ID of *fix atc* instance
- on\_the\_fly = name of the AtC sub-command
- *bond* or *kernel* = specifies on-the-fly calculation of *bond* or *kernel* matrix elements
- *on* or *off* = activate or discontinue on-the-fly mode

### Examples

```
fix_modify AtC on_the_fly bond on  
fix_modify AtC on_the_fly kernel  
fix_modify AtC on_the_fly kernel off
```

### Description

Overrides normal mode of pre-calculating and storing bond pair-to-node and kernel atom-to-node matrices. If activated, it will calculate elements of these matrices during repeated calls of field computations (i.e. “on-the-fly”) and not store them for future use. The *on* flag is optional - if omitted, `on_the_fly` will be activated for the specified matrix. Can be deactivated using the *off* flag.

### Restrictions

Must be used with *fix atc hardy*.

### Related AtC commands

- [fix\\_modify AtC command overview](#)

### Default

By default, on-the-fly calculation is not active (i.e. *off*). However, THE code does a memory allocation check to determine if it can store all needed bond and kernel matrix elements. If this allocation fails, on-the-fly will be activated.

## 17.87.27 fix\_modify AtC rates command

### Syntax

```
fix_modify <AtC fixID> rates <add|delete> <list_of_fields>
```

- AtC fixID = ID of *fix atc* instance
- rates = name of the AtC sub-command
- *add* or *delete* = select whether to add or delete calculation of rates for the listed output fields
- list\_of\_fields = one or more of the fields listed below:
  - density : mass per unit volume
  - displacement : displacement vector
  - momentum : momentum per unit volume
  - velocity : defined by momentum divided by density
  - projected\_velocity : simple kernel estimation of atomic velocities
  - temperature : temperature derived from the relative atomic kinetic energy
  - kinetic\_temperature : temperature derived from the full kinetic energy
  - number\_density : simple kernel estimation of number of atoms per unit volume
  - stress : Cauchy stress tensor for eulerian analysis (atom\_element\_map), or first Piola-Kirchhoff stress tensor for lagrangian analysis
  - transformed\_stress : first Piola-Kirchhoff stress tensor for eulerian analysis (atom\_element\_map), or Cauchy stress tensor for lagrangian analysis
  - heat\_flux : spatial heat flux vector for eulerian, or referential heat flux vector for lagrangian
  - potential\_energy : potential energy per unit volume
  - kinetic\_energy : kinetic energy per unit volume
  - thermal\_energy : thermal energy (kinetic energy - continuum kinetic energy) per unit volume
  - internal\_energy : total internal energy (potential + thermal) per unit volume
  - energy : total energy (potential + kinetic) per unit volume
  - number\_density : number of atoms per unit volume
  - eshelby\_stress : configurational stress (energy-momentum) tensor defined by [Eshelby]
  - vacancy\_concentration : volume fraction of vacancy content
  - type\_concentration : volume fraction of a specific atom type

## Examples

```
fix_modify AtC rates add temperature velocity stress
fix_modify AtC rates delete stress
```

## Description

Requests calculation and output of rates (time derivatives) of the fields from the AtC transfer class. For Eulerian analysis (see *fix\_modify AtC atom\_element\_map*) these rates are the partial time derivatives of the nodal fields, not the full (material) time derivatives.

## Restrictions

Must be used with *fix atc hardy*.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC atom\_element\_map*
- *fix\_modify AtC fields*
- *fix\_modify AtC fields*

## Default

None.

## 17.87.28 fix\_modify AtC initial command

### Syntax

```
fix_modify <AtC fixID> initial <field> <nodeset> <constant|function>
```

- AtC fixID = ID of *fix atc* instance
- initial = name of the AtC sub-command
- field = field kind name valid for type of physics: *temperature* or *electron\_temperature*
- nodeset = name of set of nodes to apply initial condition
- *constant* or *function* = value or name of function followed by its parameters

## Examples

```
fix_modify AtC initial temperature groupNAME 10.
```

## Description

Sets the initial values for the specified field at the specified nodes.

## Restrictions

The keyword *all* is reserved and thus not available as nodeset name.

## Related AtC commands

- [fix\\_modify AtC command overview](#)

## Default

None.

## 17.87.29 fix\_modify AtC internal\_element\_set command

### Syntax

```
fix_modify <AtC fixID> internal_element_set <element_set_name>
```

- AtC fixID = ID of *fix atc* instance
- internal\_element\_set = name of the AtC sub-command
- element\_set\_name = name of element set defining internal region, or *off*

## Examples

```
fix_modify AtC internal_element_set myElementSet
fix_modify AtC internal_element_set off
```

## Description

Enables AtC to base the region for internal atoms to be an element set. If no ghost atoms are used, all the AtC atoms must be constrained to remain in this element set by the user, e.g., with walls. If boundary atoms are used in conjunction with Eulerian atom maps AtC will partition all atoms of a boundary or internal type to be of type internal if they are in the internal region or to be of type boundary otherwise.

## Restrictions

If boundary atoms are used in conjunction with Eulerian atom maps, the Eulerian reset frequency must be an integer multiple of the Lammmps reneighbor frequency.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC atom\_element\_map*
- *fix\_modify AtC boundary type*

## Default

*off*

## 17.87.30 fix\_modify AtC internal\_quadrature command

### Syntax

```
fix_modify <AtC fixID> internal_quadrature <on|off> [region]
```

- AtC fixID = ID of *fix atc* instance
- internal\_quadrature = name of the AtC sub-command
- on or off = select whether internal quadrature is enabled or not
- region = treat finite elements as within MD region (optional)

### Examples

```
fix_modify AtC internal_quadrature off
```

### Description

Command to use or not use atomic quadrature on internal elements fully filled with atoms. By turning the internal quadrature off these elements do not contribute to the governing PDE and the fields at the internal nodes follow the weighted averages of the atomic data.

Optional region tag specifies which finite element nodes will be treated as being within the MD region. This option is only valid with internal\_quadrature off.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

on.

## 17.87.31 fix\_modify AtC kernel\_bandwidth command

### Syntax

```
fix_modify <AtC fixID> kernel_bandwidth <value>
```

- AtC fixID = ID of *fix atc* instance
- kernel\_bandwidth = name of the AtC sub-command
- value = new bandwidth value

### Examples

```
fix_modify AtC kernel_bandwidth 8
```

### Description

Sets a maximum parallel bandwidth for the kernel functions during parallel communication. If the command is not issued, the default will be to assume the bandwidth of the kernel matrix corresponds to the number of sampling locations.

### Restrictions

Only is used if kernel functions are being used.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

Number of sample locations.

### 17.87.32 `fix_modify AtC control lumped_lambda_solve` command

#### Syntax

```
fix_modify <AtC fixID> control lumped_lambda_solve <on|off>
```

- AtC fixID = ID of *fix atc* instance
- control lumped\_lambda\_solve = name of the AtC sub-command
- *on* or *off* = Toggles state of lumped matrix

#### Examples

```
fix_modify AtC control lumped_lambda_solve on
```

#### Description

Command select whether to use or not use lumped matrix for lambda solve.

#### Restrictions

None.

#### Related AtC commands

- *fix\_modify AtC command overview*

#### Default

off

### 17.87.33 `fix_modify AtC control mask_direction` command

#### Syntax

```
fix_modify <AtC fixID> control mask_direction <direction> <on|off>
```

- AtC fixID = ID of *fix atc* instance
- control mask\_direction = name of the AtC sub-command
- direction = select direction
- *on* or *off* = Toggles state



## Examples

```
fix_modify AtC control mask_direction 0 on
```

## Description

Command to mask out certain dimensions from the atomic regulator

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*

## 17.87.34 fix\_modify AtC mass\_matrix command

### Syntax

```
fix_modify <AtC fixID> mass_matrix <fe|md_fe>
```

- AtC fixID = ID of *fix atc* instance
- mass\_matrix = name of the AtC sub-command
- *fe* or *md\_fe* = activate/deactivate using the FE mass matrix in the MD region

## Examples

```
fix_modify AtC mass_matrix fe
```

## Description

Determines whether AtC uses the FE mass matrix based on Gaussian quadrature or based on atomic quadrature in the MD region. This is useful for fully overlapping simulations to improve efficiency.

## Restrictions

Should not be used unless the FE region is contained within the MD region, otherwise the method will be unstable and inaccurate.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

*md\_fe*

## 17.87.35 fix\_modify AtC material command

### Syntax

```
fix_modify <AtC fixID> material <elementset_name> <material_id>
```

- AtC fixID = ID of *fix atc* instance
- material = name of the AtC sub-command
- elementset\_name = name of the elementset
- material\_id = ID of the material

### Examples

```
fix_modify AtC material gap_region 1
```

### Description

Sets the material model in *elementset\_name* to be of type *material\_id*.

### Restrictions

The element set must already be created and the material must be specified in the material file given the the atc fix on construction

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

All elements default to the first material in the material file.

## 17.87.36 `fix_modify AtC mesh add_to_nodeset` command

### Syntax

```
fix_modify <AtC fixID> mesh add_to_nodeset <id> <xmin> <xmax> <ymin> <ymax> <zmin> <zmax>
```

- AtC fixID = ID of *fix atc* instance
- mesh create\_nodeset = name of the AtC sub-command
- id = id to assign to the collection of FE nodes
- <xmin> <xmax> <ymin> <ymax> <zmin> <zmax> = coordinates of the bounding box that contains the desired nodes to be added

### Examples

```
fix_modify AtC mesh add_to_nodeset lbc -12.1 -11.9 -12 12 -12 12
```

### Description

Command to add nodes to an already existing FE nodeset.

### Restrictions

None

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC mesh create\_nodeset*

### Default

Coordinates are assumed to be in lattice units.

## 17.87.37 `fix_modify AtC mesh create` command

### Syntax

```
fix_modify <AtC fixID> mesh create <nx> <ny> <nz> <region-ID> <f|p> <f|p> <f|p>
```

- AtC fixID = ID of *fix atc* instance
- mesh create = name of the AtC sub-command
- nx ny nz = number of elements in x-, y-, and z-direction
- region-ID = ID of region that is to be meshed
- f or p = periodicity flags for x-, y-, and z-direction

## Examples

```
fix_modify AtC mesh create 10 1 1 feRegion p p p
```

## Description

Creates a uniform mesh in a rectangular region.

## Restrictions

Creates only uniform rectangular grids in a rectangular region

## Related AtC commands

- [fix\\_modify AtC command overview](#)
- [fix\\_modify AtC mesh quadrature](#)

## Default

When created, the mesh defaults to gauss2 (2-point Gaussian) quadrature. Use the [fix\\_modify AtC mesh quadrature](#) command to change the quadrature style.

## 17.87.38 fix\_modify AtC mesh create\_elementset command

### Syntax

```
fix_modify <AtC fixID> mesh create_elementset <id> <xmin> <xmax> <ymin> <ymax> <zmin>  
↔<zmax>
```

- AtC fixID = ID of *fix atc* instance
- mesh create\_elementset = name of the AtC sub-command
- id = id to assign to the collection of FE nodes
- <xmin> <xmax> <ymin> <ymax> <zmin> <zmax> = coordinates of the bounding box that contains only the desired elements

## Examples

```
fix_modify AtC mesh create_elementset middle -4.1 4.1 -100 100 -100 1100
```

## Description

Command to assign an id to a set of FE elements to be used subsequently in defining material and mesh-based operations.

## Restrictions

Only viable for rectangular grids.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC mesh delete\_elements*
- *fix\_modify AtC mesh nodeset\_to\_elementset*

## Default

Coordinates are assumed to be in lattice units.

## 17.87.39 fix\_modify AtC mesh create\_faceset box command

### Syntax

```
fix_modify <AtC fixID> mesh create_faceset <id> box <xmin> <xmax> <ymin> <ymax> <zmin>
↔<zmax> <in|out> [units]
```

- AtC fixID = ID of *fix atc* instance
- mesh create\_faceset = name of the AtC sub-command
- id = id to assign to the collection of FE faces
- box = use bounding box to define FE faces
- <xmin> <xmax> <ymin> <ymax> <zmin> <zmax> = coordinates of the bounding box that is coincident with the desired FE faces
- <in|out> = “in” gives inner faces to the box, “out” gives the outer faces to the box
- units = option to specify real as opposed to lattice units

### Examples

```
fix_modify AtC mesh create_faceset obndy box -4.0 4.0 -12 12 -12 12 out
```

## Description

Command to assign an id to a set of FE faces.

## Restrictions

Only viable for rectangular grids.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC mesh create\_faceset plane*

## Default

The default options are units = lattice and the use of outer faces.

## 17.87.40 fix\_modify AtC mesh create\_faceset plane command

### Syntax

```
fix_modify <AtC fixID> mesh create_faceset <id> plane <x|y|z> <val1> <x|y|z> <lval2>  
↔<uval2> [units]
```

- AtC fixID = ID of *fix atc* instance
- mesh create\_faceset = name of the AtC sub-command
- id = id to assign to the collection of FE faces
- plane = use plane to define faceset
- <val1>,<lval2>,<uval2> = plane is specified as the x|y|z=val1 plane bounded by the segments x|y|z = [lval2,uval2]
- units = option to specify real as opposed to lattice units

## Examples

```
fix_modify AtC mesh create_faceset xyplane plane y 0 x -4 0
```

## Description

Command to assign an id to a set of FE faces.

## Restrictions

Only viable for rectangular grids.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC mesh create\_faceset box*

## Default

The default options are units = lattice.

## 17.87.41 fix\_modify AtC mesh create\_nodeset command

### Syntax

```
fix_modify <AtC fixID> mesh create_nodeset <id> <xmin> <xmax> <ymin> <ymax> <zmin> <zmax>
```

- AtC fixID = ID of *fix atc* instance
- mesh create\_nodeset = name of the AtC sub-command
- id = id to assign to the collection of FE nodes
- <xmin> <xmax> <ymin> <ymax> <zmin> <zmax> = coordinates of the bounding box that contains only the desired nodes

### Examples

```
fix_modify AtC mesh create_nodeset lbc -12.1 -11.9 -12 12 -12 12
```

### Description

Command to assign an id to a set of FE nodes to be used subsequently in defining boundary conditions.

### Restrictions

None

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC mesh add\_to\_nodeset*

### Default

Coordinates are assumed to be in lattice units.

## 17.87.42 `fix_modify AtC mesh delete_elements` command

### Syntax

```
fix_modify <AtC fixID> mesh delete_elements <id>
```

- AtC fixID = ID of *fix atc* instance
- mesh create\_elementset = name of the AtC sub-command
- id = id of the element set

### Examples

```
fix_modify AtC mesh delete_elements gap
```

### Description

Deletes a group of elements from the mesh.

### Restrictions

None.

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC mesh create\_elementset*
- *fix\_modify AtC mesh nodeset\_to\_elementset*



## Default

None.

## 17.87.43 `fix_modify AtC mesh nodeset_to_elementset` command

### Syntax

```
fix_modify <AtC fixID> mesh nodeset_to_elementset <nodeset_id> <elementset_id> <max/min>
```

- AtC fixID = ID of *fix atc* instance
- mesh nodeset\_to\_elementset = name of the AtC sub-command
- nodeset\_id = id of desired nodeset from which to create the elementset
- elementset\_id = id to assign to the collection of FE elements
- <max/min> = flag to choose either the maximal or minimal elementset

### Examples

```
fix_modify AtC mesh nodeset_to_elementset myNodeset myElementset min
```

### Description

Command to create an elementset from an existing nodeset. Either the minimal element set of elements with all nodes in the set, or maximal element set with all elements with at least one node in the set, can be created.

### Restrictions

None.

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC mesh create\_elementset*
- *fix\_modify AtC mesh delete\_elements*

### Default

Unless specified, the maximal element set is created.

## 17.87.44 `fix_modify` AtC mesh output command

### Syntax

```
fix_modify <AtC fixID> mesh output <file_prefix>
```

- AtC fixID = ID of *fix atc* instance
- mesh output = name of the AtC sub-command
- file\_prefix = prefix of various generated output files

### Examples

```
fix_modify AtC mesh output meshData
```

### Description

Command to output mesh and associated data: nodesets, facesets, and elementsets. This data is only output once upon initialization since currently the mesh is static. Creates binary (EnSight, “gold” format) output of mesh data.

### Restrictions

None.

### Related AtC commands

- *fix\_modify AtC command overview*

### Default

None.

## 17.87.45 `fix_modify` AtC mesh quadrature command

### Syntax

```
fix_modify <AtC fixID> mesh quadrature <quad>
```

- AtC fixID = ID of *fix atc* instance
- mesh quadrature = name of the AtC sub-command
- quad = *nodal* or *gauss1* or *gauss2* or *gauss3* or *face*

## Examples

```
fix_modify AtC mesh quadrature face
```

## Description

(Re-)assigns the quadrature style for an existing mesh. When a mesh is created its quadrature method defaults to gauss2. Use this call to change it after the fact.

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC mesh create*

## Default

None.

## 17.87.46 fix\_modify AtC mesh read command

### Syntax

```
fix_modify <AtC fixID> mesh read <f|p> <f|p> <f|p>
```

- AtC fixID = ID of *fix atc* instance
- mesh read = name of the AtC sub-command
- filename = name of the file containing the mesh to be read
- f or p = periodicity flags for x-, y-, and z-direction (optional)

## Examples

```
fix_modify AtC mesh read myComponent.mesh p p p
fix_modify AtC mesh read myOtherComponent.exo
```

## Description

Reads a mesh from a text or exodus file, and assigns periodic boundary conditions if needed.

## Restrictions

None

## Related AtC commands

- *fix\_modify* AtC command overview
- *fix\_modify* AtC mesh create
- *fix\_modify* AtC mesh write

## Default

Periodicity flags are set to false (f) by default.

## 17.87.47 fix\_modify AtC mesh write command

### Syntax

```
fix_modify <AtC fixID> mesh write <f|p> <f|p> <f|p>
```

- AtC fixID = ID of *fix atc* instance
- mesh write = name of the AtC sub-command
- filename = name of the file containing the mesh to be write

### Examples

```
fix_modify AtC mesh write myMesh.mesh
```

## Description

Writes a mesh to a text file.

## Restrictions

None

## Related AtC commands

- [fix\\_modify AtC command overview](#)
- [fix\\_modify AtC mesh create](#)
- [fix\\_modify AtC mesh read](#)

## Default

None.

## 17.87.48 fix\_modify AtC output command

### Syntax

```
fix_modify <AtC fixID> output <filename_prefix> <frequency> [text|full_
↪text|binary|vector_components|tensor_components]
fix_modify <AtC fixID> output index [step|time]
```

- AtC fixID = ID of *fix atc* instance
- *output* or *output index* = name of the AtC sub-command
- filename\_prefix = prefix for data files (for *output*)
- frequency = frequency of output in timesteps (for *output*)
- optional keywords for *output*:
  - text = creates text output of index, step and nodal variable values for unique nodes
  - full\_text = creates text output index, nodal id, step, nodal coordinates and nodal variable values for unique and image nodes
  - binary = creates binary EnSight output
  - vector\_components = outputs vectors as scalar components
  - tensor\_components = outputs tensor as scalar components (for use with ParaView)
- *step* or *time* = index output by step or by time (for *output index*)

### Examples

```
fix_modify AtC output heatFE 100
fix_modify AtC output hardyFE 1 text tensor_components
fix_modify AtC output hardyFE 10 text binary tensor_components
fix_modify AtC output index step
```

## Description

Creates text and/or binary (EnSight, “gold” format) output of nodal/mesh data which is transfer/physics specific. Output indexing by step or time is possible.

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix atc command*

## Default

No default format. Output indexed by time.

## 17.87.49 fix\_modify AtC output boundary\_integral command

### Syntax

```
fix_modify <AtC fixID> output boundary_integral <fieldname> faceset [name]
```

- AtC fixID = ID of *fix atc* instance
- output boundary\_integral = name of the AtC sub-command
- fieldname = name of hardy field
- faceset = required keyword
- name= name of faceset

### Examples

```
fix_modify AtC output boundary_integral stress faceset loop1
```

## Description

Calculates a surface integral of the given field dotted with the outward normal of the faces and puts output in the “GLOBALS” file.

## Restrictions

Must be used with the hardy/field type of *fix atc*

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

None.

## 17.87.50 *fix\_modify* AtC output contour\_integral command

### Syntax

```
fix_modify <AtC fixID> output contour_integral <fieldname> faceset <name> [axis [x|y|z]]
```

- AtC fixID = ID of *fix atc* instance
- output contour\_integral = name of the AtC sub-command
- fieldname = name of hardy field
- faceset = required keyword
- name = name of faceset
- *axis x* or *axis y* or *axis z* = (optional)

### Examples

```
fix_modify AtC output contour_integral stress faceset loop1
```

### Description

Calculates a surface integral of the given field dotted with the outward normal of the faces and puts output in the “GLOBALS” file.

### Restrictions

Must be used with the hardy/field type of *fix atc*

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

None.

## 17.87.51 fix\_modify AtC output nodeset command

### Syntax

```
fix_modify <AtC fixID> output nodeset <nodeset_name> <operation>
```

- AtC fixID = ID of *fix atc* instance
- output nodeset = name of the AtC sub-command
- nodeset\_name= name of nodeset to be operated on
- operation = *sum*
  - *sum* = creates nodal sum over nodes in specified nodeset

### Examples

```
fix_modify AtC output nodeset nset1 sum
```

### Description

Performs operation over the nodes belonging to specified nodeset and outputs resulting variable values to GLOBALS file.

### Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix atc command*



## Default

None.

## 17.87.52 `fix_modify` AtC output volume\_integral command

### Syntax

```
fix_modify <AtC fixID> output volume_integral <elementset_name> <field>
```

- AtC fixID = ID of *fix atc* instance
- output volume\_integral = name of the AtC sub-command
- elementset\_name= name of elementset to be integrated over
- fieldname = name of field to integrate

### Examples

```
fix_modify AtC output volume_integral eset1 mass_density
```

### Description

Performs volume integration of specified field over elementset and outputs resulting variable values to GLOBALS file.

### Restrictions

None.

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix atc command*

## Default

None.

## 17.87.53 `fix_modify` AtC pair\_interactions command

## 17.87.54 `fix_modify` AtC bond\_interactions command

### Syntax

```
fix_modify <AtC fixID> pair_interactions <on|off>  
fix_modify <AtC fixID> bond_interactions <on|off>
```

- AtC fixID = ID of *fix atc* instance
- *pair\_interactions* or *bond\_interactions* = name of the AtC sub-command
- *on* or *off* = activate or deactivate

## Examples

```
fix_modify AtC pair_interactions off
fix_modify AtC bond_interactions on
```

## Description

Include bonds and/or pairs in stress and heat flux computations.

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

*pair\_interactions*: on, *bond\_interactions*: off

## 17.87.55 fix\_modify AtC poisson\_solver command

### Syntax

```
fix_modify <AtC fixID> poisson_solver mesh create <nx> <ny> <nz> <region-ID> <f|p> <f|p>
↔<f|p>
```

- AtC fixID = ID of *fix atc* instance
- *poisson\_solver* = name of the AtC sub-command
- *nx ny nz* = number of elements in x, y, and z
- *region-id* = id of region to be meshed
- *f* or *p* = periodicity flags for x, y, and z

## Examples

```
fix_modify AtC poisson_solver mesh create 10 1 1 feRegion p p p
```

## Description

Creates a uniform mesh in a rectangular region.

## Restrictions

Creates only uniform rectangular grids in rectangular regions.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

None.

## 17.87.56 fix\_modify AtC read\_restart command

### Syntax

```
fix_modify <AtC fixID> read_restart <file_name>
```

- AtC fixID = ID of *fix atc* instance
- read\_restart = name of the AtC sub-command
- file\_name = name of AtC restart file

## Examples

```
fix_modify AtC read_restart restart.mydata.AtC
```

## Description

Reads the current state of the AtC fields from a named text-based restart file.

## Restrictions

The restart file only contains fields and their time derivatives. The reference positions of the atoms and the commands that initialize the fix are not saved e.g. an identical mesh containing the same atoms will have to be recreated.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC write\_restart*

## Default

None.

## 17.87.57 fix\_modify AtC remove\_molecule command

### Syntax

```
fix_modify <AtC fixID> remove_molecule <tag>
```

- AtC fixID = ID of *fix atc* instance
- remove\_molecule = name of the AtC sub-command
- tag = tag for tracking a molecule

### Examples

```
fix_modify AtC remove_molecule water
```

### Description

Removes tag designated for tracking a specified set of molecules.

### Restrictions

None.

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC add\_species*
- *fix\_modify AtC add\_molecule*
- *fix\_modify AtC remove\_species*

**Default**

None.

**17.87.58 fix\_modify AtC remove\_source command****Syntax**

```
fix_modify <AtC fixID> remove_source <field> <element_set>
```

- AtC fixID = ID of *fix atc* instance
- remove\_source = name of the AtC sub-command
- field = field kind name valid for type of physics: *temperature* or *electron\_temperature*
- element\_set = name of set of elements

**Examples**

```
fix_modify AtC remove_source temperature groupNAME
```

**Description**

Remove a domain source.

**Restrictions**

The keyword *all* is reserved and thus not available as element\_set name.

**Related AtC commands**

- *fix\_modify AtC command overview*
- *fix\_modify AtC source*

**Default**

None.

**17.87.59 fix\_modify AtC remove\_species command****Syntax**

```
fix_modify <AtC fixID> remove_species <tag>
```

- AtC fixID = ID of *fix atc* instance
- remove\_species = name of the AtC sub-command

- tag = tag for tracking a species

### Examples

```
fix_modify AtC remove_species gold
```

### Description

Removes tag designated for tracking a specified species.

### Restrictions

None.

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC add\_species*
- *fix\_modify AtC add\_molecule*
- *fix\_modify AtC remove\_molecule*

### Default

None.

## 17.87.60 fix\_modify AtC reset\_atomic\_reference\_positions command

### Syntax

```
fix_modify <AtC fixID> reset_atomic_reference_positions
```

- AtC fixID = ID of *fix atc* instance
- reset\_atomic\_reference\_positions = name of the AtC sub-command

### Examples

```
fix_modify AtC reset_atomic_reference_positions
```

## Description

Resets the atomic positions ATC uses to perform point to field operations. It can be used to use perfect lattice sites in ATC but a thermalized or deformed lattice in LAMMPS.

## Restrictions

None.

## Related AtC commands

- [fix\\_modify AtC command overview](#)

## Default

None

## 17.87.61 fix\_modify AtC reset\_time command

### Syntax

```
fix_modify <AtC fixID> reset_time <value>
```

- AtC fixID = ID of *fix atc* instance
- reset\_time = name of the AtC sub-command
- value = new time value

### Examples

```
fix_modify AtC reset_time 0.0
```

## Description

Resets the simulation time counter.

## Restrictions

None.

## Related AtC commands

- [fix\\_modify AtC command overview](#)

## Default

None

## 17.87.62 fix\_modify AtC sample\_frequency command

### Syntax

```
fix_modify <AtC fixID> sample_frequency <freq>
```

- AtC fixID = ID of *fix atc* instance
- sample\_frequency = name of the AtC sub-command
- freq = frequency to sample fields in number of steps

### Examples

```
fix_modify AtC sample_frequency 10
```

### Description

Specifies a frequency at which fields are computed for the case where time filters are being applied.

### Restrictions

Must be used with *fix atc hardy* and is only relevant when time filters are being used.

## Related AtC commands

- [fix\\_modify AtC command overview](#)

## Default

None.



### 17.87.63 `fix_modify AtC set reference_potential_energy` command

#### Syntax

```
fix_modify <AtC fixID> set reference_potential_energy [<value|filename>]
```

- AtC fixID = ID of *fix atc* instance
- set reference\_potential\_energy = name of the AtC sub-command
- value = optional user specified zero point for PE in native LAMMPS energy units
- filename = optional user specified string for file of nodal PE values to be read-in

#### Examples

```
fix_modify AtC set reference_potential_energy
fix_modify AtC set reference_potential_energy -0.05
fix_modify AtC set reference_potential_energy myPEvalues
```

#### Description

Used to set various quantities for the post-processing algorithms. It sets the zero point for the potential energy density using the value provided for all nodes, or from the current configuration of the lattice if no value is provided, or values provided within the specified filename.

#### Restrictions

Must be used with *fix atc hardy* or *fix atc field*.

#### Related AtC commands

- [fix\\_modify AtC command overview](#)

#### Default

Defaults to the LAMMPS zero point i.e. isolated atoms.

### 17.87.64 `fix_modify AtC source` command

#### Syntax

```
fix_modify <AtC fixID> source <field> <element_set> <value|function>
```

- AtC fixID = ID of *fix atc* instance
- source = name of the AtC sub-command
- field = field kind name valid for type of physics: *temperature* or *electron\_temperature*
- element\_set = name of set of elements

- *value* or *function* = value or name of function followed by its parameters

## Examples

```
fix_modify AtC source temperature middle temporal_ramp 10.0 0.0
```

## Description

Add domain sources to the mesh. The units are consistent with LAMMPS's units for mass, length and time and are defined by the PDE being solved, e.g. for thermal transfer the balance equation is for energy and source is energy per time.

## Restrictions

The keyword *all* is reserved and thus not available as `element_set` name.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC remove\_source*

## Default

None.

## 17.87.65 fix\_modify AtC source\_integration command

### Syntax

```
fix_modify <AtC fixID> source_integration <fe|atom>
```

- AtC `fixID` = ID of *fix atc* instance
- `source_integration` = name of the AtC sub-command
- *fe* or *atom* = (undocumented)

## Examples

```
fix_modify AtC source_integration atom
```

## Description

(undocumented)

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

Default is *fe*

## 17.87.66 fix\_modify AtC temperature\_definition command

### Syntax

```
fix_modify <AtC fixID> temperature_definition <kinetic|total>
```

- AtC fixID = ID of *fix atc* instance
- temperature\_definition = name of the AtC sub-command
- *kinetic* or *total* = (undocumented)

### Examples

```
fix_modify AtC temperature_definition kinetic
```

## Description

Change the definition for the atomic temperature used to create the finite element temperature. The kinetic option is based only on the kinetic energy of the atoms while the total option uses the total energy (kinetic + potential) of an atom.

## Restrictions

This command is only valid when using thermal coupling. Also, while not a formal restriction, the user should ensure that associating a potential energy with each atom makes physical sense for the total option to be meaningful.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

*kinetic*

## 17.87.67 fix\_modify AtC filter command

### Syntax

```
fix_modify <AtC fixID> filter <on|off|equilibrate>
```

- AtC fixID = ID of *fix atc* instance
- filter = name of the AtC sub-command
- *on* or *off* or *equilibrate* = Select state of filter

### Examples

```
fix_modify AtC filter on
```

### Description

Filters the MD dynamics to construct a more appropriate continuous field. Equilibrating first filters the time derivatives without changing the dynamics to provide a better initial condition to the filtered dynamics.

### Restrictions

Only for use with these specific transfers: thermal, two\_temperature

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC filter scale*
- *fix\_modify AtC equilibrium\_start*

## Default

off

## 17.87.68 fix\_modify AtC time\_integration command

### Syntax

```
fix_modify <AtC fixID> time_integration <descriptor>
```

- AtC fixID = ID of *fix atc* instance
- time\_integration = name of the AtC sub-command
- descriptor = *gear* or *fractional\_step* or *verlet*

### Examples

```
fix_modify AtC time_integration fractional_step
```

### Description

Command to select the thermal or momentum time integration.

---

Options for thermal time integration:

#### *gear*

atomic velocity update with second order Verlet, nodal temperature update with third or fourth order Gear, thermostats based on controlling power

#### *fractional\_step*

atomic velocity update with second order Verlet, mixed nodal temperature update, 3/4 Gear for continuum and 2 Verlet for atomic contributions, thermostats based on controlling discrete energy changes

---

Options for momentum time integration:

#### *verlet*

atomic velocity update with second order Verlet, nodal temperature update with second order Verlet, kinetostats based on controlling force

#### *fractional\_step*

atomic velocity update with second order Verlet, mixed nodal momentum update, second order Verlet for continuum and exact second order Verlet for atomic contributions, kinetostats based on controlling discrete momentum changes

#### *gear*

atomic velocity update with second order Verlet, nodal temperature update with third or fourth order Gear, kinetostats based on controlling power.

---

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*

## Default

None.

## 17.87.69 fix\_modify AtC track\_displacement command

### Syntax

```
fix_modify <AtC fixID> track_displacement <on|off>
```

- AtC fixID = ID of *fix atc* instance
- track\_displacement = name of the AtC sub-command
- *on* or *off* = (undocumented)

### Examples

```
fix_modify AtC track_displacement on
```

### Description

Determines whether displacement is tracked or not. For solids problems this is a useful quantity, but for fluids it is not relevant.

### Restrictions

Some constitutive models require the displacement field.

### Related AtC commands

- *fix\_modify AtC command overview*

**Default**

*on*

**17.87.70 fix\_modify AtC unfix command****Syntax**

```
fix_modify <AtC fixID> unfix <field> <nodeset>
```

- AtC fixID = ID of *fix atc* instance
- unfix = name of the AtC sub-command
- field = field kind name valid for type of physics: *temperature* or *electron\_temperature*
- nodeset = name of set of nodes to apply boundary condition

**Examples**

```
fix_modify AtC unfix temperature groupNAME
```

**Description**

Removes constraint on field values for specified nodes.

**Restrictions**

The keyword *all* is reserved and thus not available as nodeset name.

**Related AtC commands**

- [fix\\_modify AtC command overview](#)
- [fix\\_modify AtC fix](#)

**Default**

None.

**17.87.71 fix\_modify AtC unfix\_flux command****Syntax**

```
fix_modify <AtC fixID> unfix_flux <field> <face_set> <value|function>
```

- AtC fixID = ID of *fix atc* instance
- unfix\_flux = name of the AtC sub-command

- field = field kind name valid for type of physics: *temperature* or *electron\_temperature*
- face\_set = name of set of element faces

### Examples

```
fix_modify AtC unfix_flux temperature faceSet
```

### Description

Command for removing prescribed normal fluxes e.g. heat\_flux, stress.

### Restrictions

None.

### Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC fix\_flux*

### Default

None.

## 17.87.72 fix\_modify AtC write\_atom\_weights command

### Syntax

```
fix_modify <AtC fixID> write_atom_weights <filename> <frequency>
```

- AtC fixID = ID of *fix atc* instance
- write\_atom\_weights = name of the AtC sub-command
- filename = name of file that atomic weights are written to
- frequency = how often writes will occur

### Examples

```
fix_modify AtC write_atom_weights atm_wt_file.txt 10
```



## Description

Command for writing the values of atomic weights to a specified file.

## Restrictions

None.

## Related AtC commands

- [fix\\_modify AtC command overview](#)

## Default

None

## 17.87.73 fix\_modify AtC write\_restart command

### Syntax

```
fix_modify <AtC fixID> write_restart <file_name>
```

- AtC fixID = ID of *fix atc* instance
- write\_restart = name of the AtC sub-command
- file\_name = name of AtC restart file

### Examples

```
fix_modify AtC write_restart restart.mydata.AtC
```

## Description

Dumps the current state of the fields to a named text-based restart file. This done when the command is invoked and not repeated, unlike the otherwise similar LAMMPS command.

## Restrictions

None.

## Related AtC commands

- *fix\_modify AtC command overview*
- *fix\_modify AtC read\_restart*

## Default

None.

# 17.88 fix mol/swap command

## 17.88.1 Syntax

```
fix ID group-ID mol/swap N X itype jtype seed T keyword value ...
```

- ID, group-ID are documented in *fix* command
- atom/swap = style name of this fix command
- N = invoke this fix every N steps
- X = number of swaps to attempt every N steps
- itype,jtype = two atom types to swap with each other
- seed = random # seed (positive integer)
- T = scaling temperature of the MC swaps (temperature units)
- zero or more keyword/value pairs may be appended to args
- keyword = *ke*

*ke* value = *no* or *yes*

*no* = no conservation of kinetic energy after atom swaps

*yes* = kinetic energy is conserved after atom swaps

## 17.88.2 Examples

```
fix 2 all mol/swap 100 1 2 3 29494 300.0 ke no
fix mySwap fluid mol/swap 500 10 1 2 482798 1.0
```

## 17.88.3 Description

This fix performs Monte Carlo swaps of two specified atom types within a randomly selected molecule. Two possible use cases are as follows.

First, consider a mixture of some molecules with atoms of itype and other molecules with atoms of jtype. The fix will select a random molecule and attempt to swap all the itype atoms to jtype for the first kind of molecule, or all the jtype atoms to itype for the second kind. Because the swap will only take place if it is energetically favorable, the fix can be used to determine the miscibility of 2 different kinds of molecules much more quickly than just dynamics would do it.

Second, consider diblock co-polymers with two types of monomers *itype* and *jtype*. The fix will select a random molecule and attempt to do a *itype* <-> *jtype* swap of all those monomers within the molecule. Thus the fix can be used to find the energetically favorable fractions of two flavors of diblock co-polymers.

Intra-molecular swaps of atom types are attempted every *N* timesteps. On that timestep, *X* swaps are attempted. For each attempt a single molecule ID is randomly selected. The range of possible molecule IDs from *loID* to *hiID* is pre-computed before each run begins. The *loID*/*hiID* is set for the molecule with the smallest/largest ID which has any *itype* or *jtype* atoms in it. Note that if you define a system with many molecule IDs between *loID* and *hiID* which have no *itype* or *jtype* atoms, then the fix will be inefficient at performing swaps. Also note that if atoms with molecule ID = 0 exist, they are not considered molecules by this fix; they are assumed to be solvent atoms or molecules.

Candidate atoms for swapping must also be in the fix group. Atoms within the selected molecule which are not *itype* or *jtype* are ignored.

When an atom is swapped from *itype* to *jtype* (or vice versa), if charges are defined, the charge values for *itype* versus *jtype* atoms are also swapped. This requires that all *itype* atoms in the system have the same charge value. Likewise all *jtype* atoms in the system must have the same charge value. If this is not the case, LAMMPS issues a warning that it cannot swap charge values.

If the *ke* keyword is set to yes, which is the default, and the masses of *itype* and *jtype* atoms are different, then when a swap occurs, the velocity of the swapped atom is rescaled by the sqrt of the mass ratio, so as to conserve the kinetic energy of the atom.

---

The potential energy of the entire system is computed before and after each swap is performed within a single molecule. The specified temperature *T* is used in the Metropolis criterion to accept or reject the attempted swap. If the swap is rejected all swapped values are reversed.

The potential energy calculations can include systems and models with the following features:

- manybody pair styles, including EAM
- hybrid pair styles
- long-range electrostatics (*k*space)
- triclinic systems
- potential energy contributions from other fixes

For the last bullet point, fixes can have an associated potential energy. Examples of such fixes include: *efield*, *gravity*, *addforce*, *langevin*, *restrain*, *temp/berendsen*, *temp/rescale*, and *wall fixes*. For that energy to be included in the total potential energy of the system (the quantity used for the swap accept/reject decision), you MUST enable the *fix\_modify energy* option for that fix. The doc pages for individual *fix* commands specify if this should be done.

---

**Note:** One comment on computational efficiency. If the cutoff lengths defined for the pair style are different for *itype* versus *jtype* atoms (for any of their interactions with any other atom type), then a new neighbor list needs to be generated for every attempted swap. This is potentially expensive if *N* is small or *X* is large.

---

### 17.88.4 Restart, *fix\_modify*, output, run start/stop, minimize info

This fix writes the state of the fix to *binary restart files*. This includes information about the random number generator seed, the next timestep for MC exchanges, the number of exchange attempts and successes etc. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

---

**Note:** For this to work correctly, the timestep must **not** be changed after reading the restart with *reset\_timestep*. The fix will try to detect it and stop with an error.

---

None of the *fix\_modify* options are relevant to this fix.

This fix computes a global vector of length 2, which can be accessed by various *output commands*. The vector values are the following global cumulative quantities:

- 1 = swap attempts
- 2 = swap accepts

The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.88.5 Restrictions

This fix is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

### 17.88.6 Related commands

*fix atom/swap*, *fix gcmc*

### 17.88.7 Default

The option default is *ke = yes*.

## 17.89 *fix momentum* command

Accelerator Variants: *momentum/kk*

## 17.90 fix momentum/chunk command

### 17.90.1 Syntax

```
fix ID group-ID momentum N keyword values ...
```

- ID, group-ID are documented in *fix* command
- momentum = style name of this fix command
- N = adjust the momentum every this many timesteps one or more keyword/value pairs may be appended

```
fix ID group-ID momentum/chunk N chunkID keyword values ...
```

- ID, group-ID are documented in *fix* command
- momentum/chunk = style name of this fix command
- N = adjust the momentum per chunk every this many timesteps
- chunkID = ID of *compute chunk/atom* command
  - one or more keyword/value settings may be appended to each of the fix commands:
- keyword = *linear* or *angular* or *rescale*
  - linear* values = xflag yflag zflag  
xflag,yflag,zflag = 0/1 to exclude/include each dimension
  - angular* values = none
  - rescale* values = none

### 17.90.2 Examples

```
fix 1 all momentum 1 linear 1 1 0
fix 1 all momentum 1 linear 1 1 1 rescale
fix 1 all momentum 100 linear 1 1 1 angular
fix 1 all momentum/chunk 100 molchunk linear 1 1 1 angular
```

### 17.90.3 Description

Fix momentum zeroes the linear and/or angular momentum of the group of atoms every N timesteps by adjusting the velocities of the atoms. Fix momentum/chunk works equivalently, but operates on a per-chunk basis.

One (or both) of the *linear* or *angular* keywords **must** be specified.

If the *linear* keyword is used, the linear momentum is zeroed by subtracting the center-of-mass velocity of the group or chunk from each atom. This does not change the relative velocity of any pair of atoms. One or more dimensions can be excluded from this operation by setting the corresponding flag to 0.

If the *angular* keyword is used, the angular momentum is zeroed by subtracting a rotational component from each atom.

This command can be used to insure the entire collection of atoms (or a subset of them) does not drift or rotate during the simulation due to random perturbations (e.g. *fix langevin* thermostating).

The *rescale* keyword enables conserving the kinetic energy of the group or chunk of atoms by rescaling the velocities after the momentum was removed.

Note that the *velocity* command can be used to create initial velocities with zero aggregate linear and/or angular momentum.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 17.90.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.90.5 Restrictions

Fix momentum/chunk is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 17.90.6 Related commands

*fix recenter*, *velocity*

#### 17.90.7 Default

none

### 17.91 fix move command

#### 17.91.1 Syntax

```
fix ID group-ID move style args keyword values ...
```

- ID, group-ID are documented in *fix* command
- move = style name of this fix command
- style = *linear* or *wiggle* or *rotate* or *transrot* or *variable*

```

linear args = Vx Vy Vz
  Vx,Vy,Vz = components of velocity vector (velocity units), any component can be
  ↪specified as NULL
wiggle args = Ax Ay Az period
  Ax,Ay,Az = components of amplitude vector (distance units), any component can be
  ↪specified as NULL
  period = period of oscillation (time units)
rotate args = Px Py Pz Rx Ry Rz period
  Px,Py,Pz = origin point of axis of rotation (distance units)
  Rx,Ry,Rz = axis of rotation vector
  period = period of rotation (time units)
transrot args = Vx Vy Vz Px Py Pz Rx Ry Rz period
  Vx,Vy,Vz = components of velocity vector (velocity units)
  Px,Py,Pz = origin point of axis of rotation (distance units)
  Rx,Ry,Rz = axis of rotation vector
  period = period of rotation (time units)
variable args = v_dx v_dy v_dz v_vx v_vy v_vz
  v_dx,v_dy,v_dz = 3 variable names that calculate x,y,z displacement as function
  ↪of time, any component can be specified as NULL
  v_vx,v_vy,v_vz = 3 variable names that calculate x,y,z velocity as function of
  ↪time, any component can be specified as NULL

```

- zero or more keyword/value pairs may be appended
- keyword = *units*

*units* value = *box* or *lattice*

## 17.91.2 Examples

```

fix 1 boundary move wiggle 3.0 0.0 0.0 1.0 units box
fix 2 boundary move rotate 0.0 0.0 0.0 0.0 0.0 1.0 5.0
fix 2 boundary move variable v_myx v_myx NULL v_VX v_VY NULL
fix 3 boundary move transrot 0.1 0.1 0.0 0.0 0.0 0.0 0.0 0.0 1.0 5.0 units box

```

## 17.91.3 Description

Perform updates of position and velocity for atoms in the group each timestep using the specified settings or formulas, without regard to forces on the atoms. This can be useful for boundary or other atoms, whose movement can influence nearby atoms.

---

**Note:** The atoms affected by this fix should not normally be time integrated by other fixes (e.g. *fix nve*, *fix nvt*), since that will change their positions and velocities twice.

---



---

**Note:** As atoms move due to this fix, they will pass through periodic boundaries and be remapped to the other side of the simulation box, just as they would during normal time integration (e.g. via the *fix nve* command). It is up to you to decide whether periodic boundaries are appropriate with the kind of atom motion you are prescribing with this fix.

---



---

**Note:** As discussed below, atoms are moved relative to their initial position at the time the fix is specified. These initial coordinates are stored by the fix in “unwrapped” form, by using the image flags associated with each atom. See

---

the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g. to 0) before invoking this fix by using the *set image* command.

---

The *linear* style moves atoms at a constant velocity, so that their position  $X = (x,y,z)$  as a function of time is given in vector notation as

$$X(t) = X0 + V * \text{delta}$$

where  $X0 = (x0,y0,z0)$  is their position at the time the fix is specified,  $V$  is the specified velocity vector with components  $(Vx,Vy,Vz)$ , and *delta* is the time elapsed since the fix was specified. This style also sets the velocity of each atom to  $V = (Vx,Vy,Vz)$ . If any of the velocity components is specified as NULL, then the position and velocity of that component is time integrated the same as the *fix nve* command would perform, using the corresponding force component on the atom.

Note that the *linear* style is identical to using the *variable* style with an *equal-style variable* that uses the *vdisplace()* function. E.g.

```
variable V equal 10.0
variable x equal vdisplace(0.0,$V)
fix 1 boundary move variable v_x NULL NULL v_V NULL NULL
```

The *wiggle* style moves atoms in an oscillatory fashion, so that their position  $X = (x,y,z)$  as a function of time is given in vector notation as

$$X(t) = X0 + A \sin(\omega * \text{delta})$$

where  $X0 = (x0,y0,z0)$  is their position at the time the fix is specified,  $A$  is the specified amplitude vector with components  $(Ax,Ay,Az)$ ,  $\omega$  is  $2 \text{ PI} / \text{period}$ , and *delta* is the time elapsed since the fix was specified. This style also sets the velocity of each atom to the time derivative of this expression. If any of the amplitude components is specified as NULL, then the position and velocity of that component is time integrated the same as the *fix nve* command would perform, using the corresponding force component on the atom.

Note that the *wiggle* style is identical to using the *variable* style with *equal-style variables* that use the *swiggle()* and *cwiggle()* functions. E.g.

```
variable A equal 10.0
variable T equal 5.0
variable omega equal 2.0*PI/$T
variable x equal swiggle(0.0,$A,$T)
variable v equal v_omega*($A-cwiggle(0.0,$A,$T))
fix 1 boundary move variable v_x NULL NULL v_v NULL NULL
```

The *rotate* style rotates atoms around a rotation axis  $R = (Rx,Ry,Rz)$  that goes through a point  $P = (Px,Py,Pz)$ . The *period* of the rotation is also specified. The direction of rotation for the atoms around the rotation axis is consistent with the right-hand rule: if your right-hand thumb points along  $R$ , then your fingers wrap around the axis in the direction of rotation.

This style also sets the velocity of each atom to  $(\omega \text{ cross } R_{\text{perp}})$  where  $\omega$  is its angular velocity around the rotation axis and  $R_{\text{perp}}$  is a perpendicular vector from the rotation axis to the atom. If the defined *atom\_style* assigns an angular velocity or angular momentum or orientation to each atom (*atom styles* sphere, ellipsoid, line, tri, body), then those properties are also updated appropriately to correspond to the atom’s motion and rotation over time.

The *transrot* style combines the effects of *rotate* and *linear* so that it is possible to prescribe a rotating group of atoms that also moves at a constant velocity. The arguments are for the translation first and then for the rotation. Since the rotation affects all coordinate components, it is not possible to set any of the translation vector components to NULL.



The *variable* style allows the position and velocity components of each atom to be set by formulas specified via the *variable* command. Each of the 6 variables is specified as an argument to the fix as *v\_name*, where *name* is the variable name that is defined elsewhere in the input script.

Each variable must be of either the *equal* or *atom* style. *Equal*-style variables compute a single numeric quantity, that can be a function of the timestep as well as of other simulation values. *Atom*-style variables compute a numeric quantity for each atom, that can be a function per-atom quantities, such as the atom's position, as well as of the timestep and other simulation values. Note that this fix stores the original coordinates of each atom (see note below) so that per-atom quantity can be used in an atom-style variable formula. See the *variable* command for details.

The first 3 variables (*v\_dx*,*v\_dy*,*v\_dz*) specified for the *variable* style are used to calculate a displacement from the atom's original position at the time the fix was specified. The second 3 variables (*v\_vx*,*v\_vy*,*v\_vz*) specified are used to compute a velocity for each atom.

Any of the 6 variables can be specified as NULL. If both the displacement and velocity variables for a particular *x,y,z* component are specified as NULL, then the position and velocity of that component is time integrated the same as the *fix nve* command would perform, using the corresponding force component on the atom. If only the velocity variable for a component is specified as NULL, then the displacement variable will be used to set the position of the atom, and its velocity component will not be changed. If only the displacement variable for a component is specified as NULL, then the velocity variable will be used to set the velocity of the atom, and the position of the atom will be time integrated using that velocity.

The *units* keyword determines the meaning of the distance units used to define the *linear* velocity and *wiggle* amplitude and *rotate* origin. This setting is ignored for the *variable* style. A *box* value selects standard units as defined by the *units* command, e.g. velocity in Angstroms/fs and amplitude and position in Angstroms for *units = real*. A *lattice* value means the velocity units are in lattice spacings per time and the amplitude and position are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing. Each of these 3 quantities may be dependent on the *x,y,z* dimension, since the lattice spacings can be different in *x,y,z*.

---

#### 17.91.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the original coordinates of moving atoms to *binary restart files*, as well as the initial timestep, so that the motion can be continuous in a restarted simulation. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

---

**Note:** Because the move positions are a function of the current timestep and the initial timestep, you cannot reset the timestep to a different value after reading a restart file, if you expect a fix move command to work in an uninterrupted fashion.

---

None of the *fix\_modify* options are relevant to this fix.

This fix produces a per-atom array which can be accessed by various *output commands*. The number of columns for each atom is 3, and the columns store the original unwrapped *x,y,z* coords of each atom. The per-atom values can be accessed on any timestep.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

For *rRESPA time integration*, this fix adjusts the position and velocity of atoms on the outermost rRESPA level.

### 17.91.5 Restrictions

none

### 17.91.6 Related commands

*fix nve, displace\_atoms*

### 17.91.7 Default

none

The option default is units = lattice.

## 17.92 fix mscg command

### 17.92.1 Syntax

```
fix ID group-ID mscg N keyword args ...
```

- ID, group-ID are documented in *fix* command
- mscg = style name of this fix command
- N = invoke this fix every this many timesteps
- zero or more keyword/value pairs may be appended
- keyword = *range* or *name* or *max*
  - range* arg = *on* or *off*
    - on* = range finding functionality is performed
    - off* = force matching functionality is performed
  - name* args = name1 ... nameN
    - name1,...,nameN = string names for each atom type (1-Ntype)
  - max* args = maxb maxa maxd
    - maxb,maxa,maxd = maximum bonds/angles/dihedrals per atom

### 17.92.2 Examples

```
fix 1 all mscg 1
fix 1 all mscg 1 range name A B
fix 1 all mscg 1 max 4 8 20
```

### 17.92.3 Description

This fix applies the Multi-Scale Coarse-Graining (MSCG) method to snapshots from a dump file to generate potentials for coarse-grained simulations from all-atom simulations, using a force-matching technique (*Izvekov, Noid*).

It makes use of the MS-CG library, written and maintained by Greg Voth's group at the University of Chicago, which is freely available on their [MS-CG GitHub site](#). See instructions on obtaining and installing the MS-CG library in the `src/MSCG/README` file, which must be done before you build LAMMPS with this fix command and use the command in a LAMMPS input script.

An example script using this fix is provided the `examples/mscg` directory.

The general workflow for using LAMMPS in conjunction with the MS-CG library to create a coarse-grained model and run coarse-grained simulations is as follows:

1. Perform all-atom simulations on the system to be coarse grained.
2. Generate a trajectory mapped to the coarse-grained model.
3. Create input files for the MS-CG library.
4. Run the range finder functionality of the MS-CG library.
5. Run the force matching functionality of the MS-CG library.
6. Check the results of the force matching.
7. Run coarse-grained simulations using the new coarse-grained potentials.

This fix can perform the range finding and force matching steps 4 and 5 of the above workflow when used in conjunction with the *rerun* command. It does not perform steps 1-3 and 6-7.

Step 2 can be performed using a Python script (what is the name?) provided with the MS-CG library which defines the coarse-grained model and converts a standard LAMMPS dump file for an all-atom simulation (step 1) into a LAMMPS dump file which has the positions of and forces on the coarse-grained beads.

In step 3, an input file named "control.in" is needed by the MS-CG library which sets parameters for the range finding and force matching functionalities. See the `examples/mscg/control.in` file as an example. And see the documentation provided with the MS-CG library for more info on this file.

When this fix is used to perform steps 4 and 5, the MS-CG library also produces additional output files. The range finder functionality (step 4) outputs files defining pair and bonded interaction ranges. The force matching functionality (step 5) outputs tabulated force files for every interaction in the system. Other diagnostic files can also be output depending on the parameters in the MS-CG library input script. Again, see the documentation provided with the MS-CG library for more info.

---

The *range* keyword specifies which MS-CG library functionality should be invoked. If *on*, the step 4 range finder functionality is invoked. *off*, the step 5 force matching functionality is invoked.

If the *name* keyword is used, string names are defined to associate with the integer atom types in LAMMPS. *Ntype* names must be provided, one for each atom type (1-Ntype).

The *max* keyword specifies the maximum number of bonds, angles, and dihedrals a bead can have in the coarse-grained model.

## 17.92.4 Restrictions

This fix is part of the MSCG package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

The MS-CG library uses C++11, which may not be supported by older compilers. The MS-CG library also has some additional numeric library dependencies, which are described in its documentation.

Currently, the MS-CG library is not setup to run in parallel with MPI, so this fix can only be used in a serial LAMMPS build and run on a single processor.

## 17.92.5 Related commands

none

## 17.92.6 Default

The default keyword settings are range off, max 4 12 36.

---

**(Izvekov)** Izvekov, Voth, J Chem Phys 123, 134105 (2005).

**(Noid)** Noid, Chu, Ayton, Krishna, Izvekov, Voth, Das, Andersen, J Chem Phys 128, 134105 (2008).

## 17.93 fix msst command

### 17.93.1 Syntax

```
fix ID group-ID msst dir shockvel keyword value ...
```

- ID, group-ID are documented in *fix* command
- msst = style name of this fix
- dir = *x* or *y* or *z*
- shockvel = shock velocity (strictly positive, distance/time units)
- zero or more keyword value pairs may be appended
- keyword = *q* or *mu* or *p0* or *v0* or *e0* or *tscale* or *beta* or *dftb*
  - q* value = cell mass-like parameter (mass<sup>2</sup>/distance<sup>4</sup> units)
  - mu* value = artificial viscosity (mass/length/time units)
  - p0* value = initial pressure in the shock equations (pressure units)
  - v0* value = initial simulation cell volume in the shock equations (distance<sup>3</sup> units)
  - e0* value = initial total energy (energy units)
  - tscale* value = reduction in initial temperature (unitless fraction between 0.0 and 1.0)
  - dftb* value = *yes* or *no* for whether using MSST in conjunction with DFTB+
  - beta* value = scale factor for improved energy conservation

## 17.93.2 Examples

```
fix 1 all msst y 100.0 q 1.0e5 mu 1.0e5
fix 2 all msst z 50.0 q 1.0e4 mu 1.0e4 v0 4.3419e+03 p0 3.7797e+03 e0 -9.72360e+02
↳tscale 0.01
fix 1 all msst y 100.0 q 1.0e5 mu 1.0e5 dftb yes beta 0.5
```

## 17.93.3 Description

This command performs the Multi-Scale Shock Technique (MSST) integration to update positions and velocities each timestep to mimic a compressive shock wave passing over the system. See [\(Reed\)](#) for a detailed description of this method. The MSST varies the cell volume and temperature in such a way as to restrain the system to the shock Hugoniot and the Rayleigh line. These restraints correspond to the macroscopic conservation laws dictated by a shock front. *shockvel* determines the steady shock velocity that will be simulated.

To perform a simulation, choose a value of *q* that provides volume compression on the timescale of 100 fs to 1 ps. If the volume is not compressing, either the shock speed is chosen to be below the material sound speed or *p0* has been chosen inaccurately. Volume compression at the start can be sped up by using a non-zero value of *tscale*. Use the smallest value of *tscale* that results in compression.

Under some special high-symmetry conditions, the pressure (volume) and/or temperature of the system may oscillate for many cycles even with an appropriate choice of mass-like parameter *q*. Such oscillations have physical significance in some cases. The optional *mu* keyword adds an artificial viscosity that helps break the system symmetry to equilibrate to the shock Hugoniot and Rayleigh line more rapidly in such cases.

The keyword *tscale* is a factor between 0 and 1 that determines what fraction of thermal kinetic energy is converted to compressive strain kinetic energy at the start of the simulation. Setting this parameter to a non-zero value may assist in compression at the start of simulations where it is slow to occur.

If keywords *e0*, *p0*, or *v0* are not supplied, these quantities will be calculated on the first step, after the energy specified by *tscale* is removed. The value of *e0* is not used in the dynamical equations, but is used in calculating the deviation from the Hugoniot.

The keyword *beta* is a scaling term that can be added to the MSST ionic equations of motion to account for drift in the conserved quantity during long timescale simulations, similar to a Berendsen thermostat. See [\(Reed\)](#) and [\(Goldman\)](#) for more details. The value of *beta* must be between 0.0 and 1.0 inclusive. A value of 0.0 means no contribution, a value of 1.0 means a full contribution.

Values of *shockvel* less than a critical value determined by the material response will not have compressive solutions. This will be reflected in lack of significant change of the volume in the MSST.

For all pressure styles, the simulation box stays orthogonal in shape. Parrinello-Rahman boundary conditions (tilted box) are supported by LAMMPS, but are not implemented for MSST.

This fix computes a temperature and pressure and potential energy each timestep. To do this, the fix creates its own computes of style “temp” “pressure”, and “pe”, as if these commands had been issued:

```
compute fix-ID_MSST_temp all temp
compute fix-ID_MSST_press all pressure fix-ID_MSST_temp

compute fix-ID_MSST_pe all pe
```

See the [compute temp](#) and [compute pressure](#) commands for details. Note that the IDs of the new computes are the fix-ID + “\_MSST\_temp” or “MSST\_press” or “\_MSST\_pe”. The group for the new computes is “all”.

The *dftb* keyword is to allow this fix to be used when LAMMPS is being driven by DFTB+, a density-functional tight-binding code. If the keyword *dftb* is used with a value of *yes*, then the MSST equations are altered to account for the electron entropy contribution to the Hugoniot relations and total energy. See (*Reed2*) and (*Goldman*) for details on this contribution. In this case, you must define a *fix external* command in your input script, which is used to callback to DFTB+ during the LAMMPS timestepping. DFTB+ will communicate its info to LAMMPS via that fix.

---

### 17.93.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of all internal variables to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

The progress of the MSST can be monitored by printing the global scalar and global vector quantities computed by the fix.

As mentioned above, the scalar is the cumulative energy change due to the fix. By monitoring the *thermodynamic econserve* output, this can be used to test if the MD timestep is sufficiently small for accurate integration of the dynamic equations.

The global vector contains four values in the following order. The vector values output by this fix are “intensive”.

[*dhugoniot*, *drayleigh*, *lagrangian\_speed*, *lagrangian\_position*]

1. *dhugoniot* is the departure from the Hugoniot (temperature units).
2. *drayleigh* is the departure from the Rayleigh line (pressure units).
3. *lagrangian\_speed* is the laboratory-frame Lagrangian speed (particle velocity) of the computational cell (velocity units).
4. *lagrangian\_position* is the computational cell position in the reference frame moving at the shock speed. This is usually a good estimate of distance of the computational cell behind the shock front.

To print these quantities to the log file with descriptive column headers, the following LAMMPS commands are suggested:

```
fix                msst all msst z
variable dhug      equal f_msst[1]
variable dray      equal f_msst[2]
variable lgr_vel   equal f_msst[3]
variable lgr_pos   equal f_msst[4]
thermo_style       custom step temp ke pe lz pzz econserve v_dhug v_dray v_lgr_vel v_lgr_
↪pos f_msst
```

### 17.93.5 Restrictions

This fix style is part of the SHOCK package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

All cell dimensions must be periodic. This fix can not be used with a triclinic cell. The MSST fix has been tested only for the group-ID all.

### 17.93.6 Related commands

*fix nphug*, *fix deform*

### 17.93.7 Default

The keyword defaults are  $q = 10$ ,  $\mu = 0$ ,  $t_{scale} = 0.01$ ,  $dftb = no$ ,  $\beta = 0.0$ . Note that  $p_0$ ,  $v_0$ , and  $\epsilon_0$  are calculated on the first timestep.

---

**(Reed)** Reed, Fried, and Joannopoulos, Phys. Rev. Lett., 90, 235503 (2003).

**(Reed2)** Reed, J. Phys. Chem. C, 116, 2205 (2012).

**(Goldman)** Goldman, Srinivasan, Hamel, Fried, Gaus, and Elstner, J. Phys. Chem. C, 117, 7885 (2013).

## 17.94 fix mvv/dpd command

## 17.95 fix mvv/edpd command

## 17.96 fix mvv/tdpd command

### 17.96.1 Syntax

```
fix ID group-ID mvv/dpd lambda
```

```
fix ID group-ID mvv/edpd lambda
```

```
fix ID group-ID mvv/tdpd lambda
```

- ID, group-ID are documented in *fix* command
- mvv/dpd, mvv/edpd, mvv/tdpd = style name of this fix command
- lambda = (optional) relaxation parameter (unitless)

## 17.96.2 Examples

```
fix 1 all mvv/dpd
fix 1 all mvv/dpd 0.5
fix 1 all mvv/edpd
fix 1 all mvv/edpd 0.5
fix 1 all mvv/tdpd
fix 1 all mvv/tdpd 0.5
```

## 17.96.3 Description

Perform time integration using the modified velocity-Verlet (MVV) algorithm to update position and velocity (fix mvv/dpd), or position, velocity and temperature (fix mvv/edpd), or position, velocity and concentration (fix mvv/tdpd) for particles in the group each timestep.

The modified velocity-Verlet (MVV) algorithm aims to improve the stability of the time integrator by using an extrapolated version of the velocity for the force evaluation:

$$\begin{aligned}v\left(t + \frac{\Delta t}{2}\right) &= v(t) + \frac{\Delta t}{2} \cdot a(t) \\r(t + \Delta t) &= r(t) + \Delta t \cdot v\left(t + \frac{\Delta t}{2}\right) \\a(t + \Delta t) &= \frac{1}{m} \cdot F\left[r(t + \Delta t), v(t) + \lambda \cdot \Delta t \cdot a(t)\right] \\v(t + \Delta t) &= v\left(t + \frac{\Delta t}{2}\right) + \frac{\Delta t}{2} \cdot a(t + \Delta t)\end{aligned}$$

where the parameter  $\lambda$  depends on the specific choice of DPD parameters, and needs to be tuned on a case-by-case basis. Specification of a *lambda* value is optional. If specified, the setting must be from 0.0 to 1.0. If not specified, a default value of 0.5 is used, which effectively reproduces the standard velocity-Verlet (VV) scheme. For more details, see *Groot*.

Fix *mvv/dpd* updates the position and velocity of each atom. It can be used with the *pair\_style mdpd* command or other pair styles such as *pair dpd*.

Fix *mvv/edpd* updates the per-atom temperature, in addition to position and velocity, and must be used with the *pair\_style edpd* command.

Fix *mvv/tdpd* updates the per-atom chemical concentration, in addition to position and velocity, and must be used with the *pair\_style tdpd* command.

---

## 17.96.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.



## 17.96.5 Restrictions

This fix is part of the DPD-MESO package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 17.96.6 Related commands

*pair\_style mdpd*, *pair\_style edpd*, *pair\_style tdpd*

## 17.96.7 Default

The default value for the optional *lambda* parameter is 0.5.

---

(Groot) Groot and Warren, J Chem Phys, 107: 4423-4435 (1997). DOI: 10.1063/1.474784

## 17.97 fix neb command

### 17.97.1 Syntax

```
fix ID group-ID neb Kspring keyword value
```

- ID, group-ID are documented in *fix* command
- neb = style name of this fix command
- Kspring = spring constant for parallel nudging force (force/distance units or force units, see parallel keyword)
- zero or more keyword/value pairs may be appended
- keyword = *parallel* or *perp* or *end*

*parallel* value = *neigh* or *ideal*

*neigh* = parallel nudging force based on distance to neighbor replicas (Kspring = force/↵distance units)

*ideal* = parallel nudging force based on interpolated ideal position (Kspring = force↵↵units)

*perp* value = *Kspring2*

*Kspring2* = spring constant for perpendicular nudging force (force/distance units)

*end* values = *estyle* *Kspring3*

*estyle* = *first* or *last* or *last/efirst* or *last/efirst/middle*

*first* = apply force to first replica

*last* = apply force to last replica

*last/efirst* = apply force to last replica and set its target energy to that of first↵↵replica

*last/efirst/middle* = same as *last/efirst* plus prevent middle replicas having lower↵↵energy than first replica

*Kspring3* = spring constant for target energy term (1/distance units)

## 17.97.2 Examples

```
fix 1 active neb 10.0
fix 2 all neb 1.0 perp 1.0 end last
fix 2 all neb 1.0 perp 1.0 end first 1.0 end last 1.0
fix 1 all neb 1.0 parallel ideal end last/efirst 1
```

## 17.97.3 Description

Add nudging forces to atoms in the group for a multi-replica simulation run via the *neb* command to perform a nudged elastic band (NEB) calculation for finding the transition state. Hi-level explanations of NEB are given with the *neb* command and on the *Howto replica* doc page. The *fix neb* command must be used with the “neb” command and defines how inter-replica nudging forces are computed. A NEB calculation is divided in two stages. In the first stage *n* replicas are relaxed toward a MEP until convergence. In the second stage, the climbing image scheme (see (*Henkelman2*)) is enabled, so that the replica having the highest energy relaxes toward the saddle point (i.e. the point of highest energy along the MEP), and a second relaxation is performed.

A key purpose of the nudging forces is to keep the replicas equally spaced. During the NEB calculation, the 3N-length vector of interatomic force  $F_i = -\text{Grad}(V)$  for each replica *I* is altered. For all intermediate replicas (i.e. for  $1 < I < N$ , except the climbing replica) the force vector becomes:

$$F_i = -\text{Grad}(V) + (\text{Grad}(V) \text{ dot } T') T' + F_{\text{nudge\_parallel}} + F_{\text{nudge\_perp}}$$

$T'$  is the unit “tangent” vector for replica *I* and is a function of  $R_i$ ,  $R_{i-1}$ ,  $R_{i+1}$ , and the potential energy of the 3 replicas; it points roughly in the direction of  $(R_{i+1} - R_{i-1})$ ; see the (*Henkelman1*) paper for details.  $R_i$  are the atomic coordinates of replica *I*;  $R_{i-1}$  and  $R_{i+1}$  are the coordinates of its neighbor replicas. The term  $(\text{Grad}(V) \text{ dot } T')$  is used to remove the component of the gradient parallel to the path which would tend to distribute the replica unevenly along the path.  $F_{\text{nudge\_parallel}}$  is an artificial nudging force which is applied only in the tangent direction and which maintains the equal spacing between replicas (see below for more information).  $F_{\text{nudge\_perp}}$  is an optional artificial spring which is applied in a direction perpendicular to the tangent direction and which prevent the paths from forming acute kinks (see below for more information).

In the second stage of the NEB calculation, the interatomic force  $F_i$  for the climbing replica (the replica of highest energy after the first stage) is changed to:

$$F_i = -\text{Grad}(V) + 2 (\text{Grad}(V) \text{ dot } T') T' + F_{\text{nudge\_perp}}$$

and the relaxation procedure is continued to a new converged MEP.

The keyword *parallel* specifies how the parallel nudging force is computed. With a value of *neigh*, the parallel nudging force is computed as in (*Henkelman1*) by connecting each intermediate replica with the previous and the next image:

$$F_{\text{nudge\_parallel}} = K_{\text{spring}} * (|R_{i+1} - R_i| - |R_i - R_{i-1}|)$$

Note that in this case the specified  $K_{\text{spring}}$  is in force/distance units.

With a value of *ideal*, the spring force is computed as suggested in ref (WeinanE) <WeinanE>

$$F_{\text{nudge\_parallel}} = -K_{\text{spring}} * (RD - RD_{\text{ideal}}) / (2 * \text{meanDist})$$

where  $RD$  is the “reaction coordinate” see *neb* section, and  $RD_{\text{ideal}}$  is the ideal  $RD$  for which all the images are equally spaced. I.e.  $RD_{\text{ideal}} = (I-1)*\text{meanDist}$  when the climbing replica is off, where *I* is the replica number). The  $\text{meanDist}$  is the average distance between replicas. Note that in this case the specified  $K_{\text{spring}}$  is in force units.

Note that the *ideal* form of nudging can often be more effective at keeping the replicas equally spaced.

The keyword *perp* specifies if and how a perpendicular nudging force is computed. It adds a spring force perpendicular to the path in order to prevent the path from becoming too strongly kinked. It can significantly improve the convergence of the NEB calculation when the resolution is poor. I.e. when few replicas are used; see (*Maras*) for details.

The perpendicular spring force is given by

$$F_{\text{nudge\_perp}} = K_{\text{spring2}} * F(R_{i-1}, R_i, R_{i+1}) (R_{i+1} + R_{i-1} - 2 R_i)$$

where  $K_{\text{spring2}}$  is the specified value.  $F(R_{i-1}, R_i, R_{i+1})$  is a smooth scalar function of the angle  $R_{i-1}, R_i, R_{i+1}$ . It is equal to 0.0 when the path is straight and is equal to 1 when the angle  $R_{i-1}, R_i, R_{i+1}$  is acute.  $F(R_{i-1}, R_i, R_{i+1})$  is defined in (*Jonsson*).

If  $K_{\text{spring2}}$  is set to 0.0 (the default) then no perpendicular spring force is added.

By default, no additional forces act on the first and last replicas during the NEB relaxation, so these replicas simply relax toward their respective local minima. By using the key word *end*, additional forces can be applied to the first and/or last replicas, to enable them to relax toward a MEP while constraining their energy  $E$  to the target energy  $E_{\text{Target}}$ .

If  $E_{\text{Target}} > E$ , the interatomic force  $F_i$  for the specified replica becomes:

$$F_i = -\text{Grad}(V) + (\text{Grad}(V) \cdot T' + (E - E_{\text{Target}}) * K_{\text{spring3}}) T', \quad \text{when } \text{Grad}(V) \cdot T' < 0$$

$$F_i = -\text{Grad}(V) + (\text{Grad}(V) \cdot T' + (E_{\text{Target}} - E) * K_{\text{spring3}}) T', \quad \text{when } \text{Grad}(V) \cdot T' > 0$$

The “spring” constant on the difference in energies is the specified  $K_{\text{spring3}}$  value.

When *estyle* is specified as *first*, the force is applied to the first replica. When *estyle* is specified as *last*, the force is applied to the last replica. Note that the *end* keyword can be used twice to add forces to both the first and last replicas.

For both these *estyle* settings, the target energy  $E_{\text{Target}}$  is set to the initial energy of the replica (at the start of the NEB calculation).

If the *estyle* is specified as *last/first* or *last/first/middle*, force is applied to the last replica, but the target energy  $E_{\text{Target}}$  is continuously set to the energy of the first replica, as it evolves during the NEB relaxation.

The difference between these two *estyle* options is as follows. When *estyle* is specified as *last/first*, no change is made to the inter-replica force applied to the intermediate replicas (neither first or last). If the initial path is too far from the MEP, an intermediate replica may relax “faster” and reach a lower energy than the last replica. In this case the intermediate replica will be relaxing toward its own local minima. This behavior can be prevented by specifying *estyle* as *last/first/middle* which will alter the inter-replica force applied to intermediate replicas by removing the contribution of the gradient to the inter-replica force. This will only be done if a particular intermediate replica has a lower energy than the first replica. This should effectively prevent the intermediate replicas from over-relaxing.

After converging a NEB calculation using an *estyle* of *last/first/middle*, you should check that all intermediate replicas have a larger energy than the first replica. If this is not the case, the path is probably not a MEP.

Finally, note that the last replica may never reach the target energy if it is stuck in a local minima which has a larger energy than the target energy.

### 17.97.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, as invoked by the *minimize* command via the *neb* command.

### 17.97.5 Restrictions

This command can only be used if LAMMPS was built with the REPLICAS package. See the *Build package* doc page for more info.

### 17.97.6 Related commands

*neb*

### 17.97.7 Default

The option defaults are parallel = neigh, perp = 0.0, ends is not specified (no inter-replica force on the end replicas).

---

**(Henkelman1)** Henkelman and Jonsson, J Chem Phys, 113, 9978-9985 (2000).

**(Henkelman2)** Henkelman, Uberuaga, Jonsson, J Chem Phys, 113, 9901-9904 (2000).

**(WeinanE)** E, Ren, Vanden-Eijnden, Phys Rev B, 66, 052301 (2002).

**(Jonsson)** Jonsson, Mills and Jacobsen, in Classical and Quantum Dynamics in Condensed Phase Simulations, edited by Berne, Ciccotti, and Coker World Scientific, Singapore, 1998, p 385.

**(Maras)** Maras, Trushin, Stukowski, Ala-Nissila, Jonsson, Comp Phys Comm, 205, 13-21 (2016).

## 17.98 fix neb/spin command

### 17.98.1 Syntax

```
fix ID group-ID neb/spin Kspring
```

- ID, group-ID are documented in *fix* command
- neb/spin = style name of this fix command

```
Kspring = spring constant for parallel nudging force  
(force/distance units or force units, see parallel keyword)
```

## 17.98.2 Examples

```
fix 1 active neb/spin 1.0
```

## 17.98.3 Description

Add nudging forces to spins in the group for a multi-replica simulation run via the *neb/spin* command to perform a geodesic nudged elastic band (GNEB) calculation for finding the transition state. Hi-level explanations of GNEB are given with the *neb/spin* command and on the *Howto replica* doc page. The *fix neb/spin* command must be used with the “*neb/spin*” command and defines how inter-replica nudging forces are computed. A GNEB calculation is divided in two stages. In the first stage *n* replicas are relaxed toward a MEP until convergence. In the second stage, the climbing image scheme is enabled, so that the replica having the highest energy relaxes toward the saddle point (i.e. the point of highest energy along the MEP), and a second relaxation is performed.

The nudging forces are calculated as explained in (*BessarabB*). See this reference for more explanation about their expression.

## 17.98.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, as invoked by the *minimize* command via the *neb/spin* command.

## 17.98.5 Restrictions

This command can only be used if LAMMPS was built with the SPIN package. See the *Build package* doc page for more info.

## 17.98.6 Related commands

*neb\_spin*

## 17.98.7 Default

none

---

(*BessarabB*) Bessarab, Uzdin, Jonsson, Comp Phys Comm, 196, 335-347 (2015).

## 17.99 fix nvt command

Accelerator Variants: *nvt/gpu*, *nvt/intel*, *nvt/kk*, *nvt/omp*

## 17.100 fix npt command

Accelerator Variants: *npt/gpu*, *npt/intel*, *npt/kk*, *npt/omp*

## 17.101 fix nph command

Accelerator Variants: *nph/kk*, *nph/omp*

### 17.101.1 Syntax

```
fix ID group-ID style_name keyword value ...
```

- ID, group-ID are documented in *fix* command
- style\_name = *nvt* or *npt* or *nph*
- one or more keyword/value pairs may be appended

keyword = *temp* or *iso* or *aniso* or *tri* or *x* or *y* or *z* or *xy* or *yz* or *xz* or *couple* or ↵  
↵ *tchain* or *pchain* or *mtk* or *tloop* or *ploop* or *nreset* or *drag* or *ptemp* or *dilate* ↵  
↵ or *scalexy* or *scaleyz* or *scalexz* or *flip* or *fixedpoint* or *update*

*temp* values = Tstart Tstop Tdamp

Tstart,Tstop = external temperature at start/end of run

Tdamp = temperature damping parameter (time units)

*iso* or *aniso* or *tri* values = Pstart Pstop Pdamp

Pstart,Pstop = scalar external pressure at start/end of run (pressure units)

Pdamp = pressure damping parameter (time units)

*x* or *y* or *z* or *xy* or *yz* or *xz* values = Pstart Pstop Pdamp

Pstart,Pstop = external stress tensor component at start/end of run (pressure ↵  
↵ units)

Pdamp = stress damping parameter (time units)

*couple* = *none* or *xyz* or *xy* or *yz* or *xz*

*tchain* value = N

N = length of thermostat chain (1 = single thermostat)

*pchain* value = N

N length of thermostat chain on barostat (0 = no thermostat)

*mtk* value = *yes* or *no* = add in MTK adjustment term or not

*tloop* value = M

M = number of sub-cycles to perform on thermostat

*ploop* value = M

M = number of sub-cycles to perform on barostat thermostat

*nreset* value = reset reference cell every this many timesteps

*drag* value = Df

Df = drag factor added to barostat/thermostat (0.0 = no drag)

*ptemp* value = Ttarget

Ttarget = target temperature for barostat

*dilate* value = dilate-group-ID  
 dilate-group-ID = only dilate atoms in this group due to barostat volume changes  
*scalexy* value = *yes* or *no* = scale xy with ly  
*scaleyz* value = *yes* or *no* = scale yz with lz  
*scalexz* value = *yes* or *no* = scale xz with lz  
*flip* value = *yes* or *no* = allow or disallow box flips when it becomes highly skewed  
*fixedpoint* values = x y z  
 x,y,z = perform barostat dilation/contraction around this point (distance units)  
*update* value = *dipole* or *dipole/dlm*  
 dipole = update dipole orientation (only for sphere variants)  
 dipole/dlm = use DLM integrator to update dipole orientation (only for sphere\_↵  
 ↵variants)

### 17.101.2 Examples

```

fix 1 all nvt temp 300.0 300.0 100.0
fix 1 water npt temp 300.0 300.0 100.0 iso 0.0 0.0 1000.0
fix 2 jello npt temp 300.0 300.0 100.0 tri 5.0 5.0 1000.0
fix 2 ice nph x 1.0 1.0 0.5 y 2.0 2.0 0.5 z 3.0 3.0 0.5 yz 0.1 0.1 0.5 xz 0.2 0.2 0.5 xy_↵
↵0.3 0.3 0.5 nreset 1000
  
```

### 17.101.3 Description

These commands perform time integration on Nose-Hoover style non-Hamiltonian equations of motion which are designed to generate positions and velocities sampled from the canonical (nvt), isothermal-isobaric (npt), and isenthalpic (nph) ensembles. This updates the position and velocity for atoms in the group each timestep.

The thermostating and barostating is achieved by adding some dynamic variables which are coupled to the particle velocities (thermostating) and simulation domain dimensions (barostating). In addition to basic thermostating and barostating, these fixes can also create a chain of thermostats coupled to the particle thermostat, and another chain of thermostats coupled to the barostat variables. The barostat can be coupled to the overall box volume, or to individual dimensions, including the *xy*, *xz* and *yz* tilt dimensions. The external pressure of the barostat can be specified as either a scalar pressure (isobaric ensemble) or as components of a symmetric stress tensor (constant stress ensemble). When used correctly, the time-averaged temperature and stress tensor of the particles will match the target values specified by *Tstart/Tstop* and *Pstart/Pstop*.

The equations of motion used are those of Shinoda et al in (*Shinoda*), which combine the hydrostatic equations of Martyna, Tobias and Klein in (*Martyna*) with the strain energy proposed by Parrinello and Rahman in (*Parrinello*). The time integration schemes closely follow the time-reversible measure-preserving Verlet and rRESPA integrators derived by Tuckerman et al in (*Tuckerman*).

The thermostat parameters for fix styles *nvt* and *npt* are specified using the *temp* keyword. Other thermostat-related keywords are *tchain*, *tloop* and *drag*, which are discussed below.

The thermostat is applied to only the translational degrees of freedom for the particles. The translational degrees of freedom can also have a bias velocity removed before thermostating takes place; see the description below. The desired temperature at each timestep is a ramped value during the run from *Tstart* to *Tstop*. The *Tdamp* parameter is specified in time units and determines how rapidly the temperature is relaxed. For example, a value of 10.0 means to relax the temperature in a timespan of (roughly) 10 time units (e.g.  $\tau$  or fs or ps - see the *units* command). The atoms in the fix group are the only ones whose velocities and positions are updated by the velocity/position update portion of the integration.

**Note:** A Nose-Hoover thermostat will not work well for arbitrary values of  $Tdamp$ . If  $Tdamp$  is too small, the temperature can fluctuate wildly; if it is too large, the temperature will take a very long time to equilibrate. A good choice for many models is a  $Tdamp$  of around 100 timesteps. Note that this is NOT the same as 100 time units for most *units* settings. A simple way to ensure this, is via using an *immediate variable* expression accessing the thermo property 'dt', which is the length of the time step. Example:

---

```
fix 1 all nvt temp 300.0 300.0 $(100.0*dt)
```

---

The barostat parameters for fix styles *npt* and *nph* is specified using one or more of the *iso*, *aniso*, *tri*, *x*, *y*, *z*, *xy*, *xz*, *yz*, and *couple* keywords. These keywords give you the ability to specify all 6 components of an external stress tensor, and to couple various of these components together so that the dimensions they represent are varied together during a constant-pressure simulation.

Other barostat-related keywords are *pchain*, *mtk*, *ploop*, *nreset*, *drag*, and *dilate*, which are discussed below.

Orthogonal simulation boxes have 3 adjustable dimensions (*x,y,z*). Triclinic (non-orthogonal) simulation boxes have 6 adjustable dimensions (*x,y,z,xy,xz,yz*). The *create\_box*, *read\_data*, and *read\_restart* commands specify whether the simulation box is orthogonal or non-orthogonal (triclinic) and explain the meaning of the *xy,xz,yz* tilt factors.

The target pressures for each of the 6 components of the stress tensor can be specified independently via the *x*, *y*, *z*, *xy*, *xz*, *yz* keywords, which correspond to the 6 simulation box dimensions. For each component, the external pressure or tensor component at each timestep is a ramped value during the run from *Pstart* to *Pstop*. If a target pressure is specified for a component, then the corresponding box dimension will change during a simulation. For example, if the *y* keyword is used, the *y*-box length will change. If the *xy* keyword is used, the *xy* tilt factor will change. A box dimension will not change if that component is not specified, although you have the option to change that dimension via the *fix deform* command.

Note that in order to use the *xy*, *xz*, or *yz* keywords, the simulation box must be triclinic, even if its initial tilt factors are 0.0.

For all barostat keywords, the *Pdamp* parameter operates like the *Tdamp* parameter, determining the time scale on which pressure is relaxed. For example, a value of 10.0 means to relax the pressure in a timespan of (roughly) 10 time units (e.g.  $\tau$  or fs or ps - see the *units* command).

---

**Note:** A Nose-Hoover barostat will not work well for arbitrary values of *Pdamp*. If *Pdamp* is too small, the pressure and volume can fluctuate wildly; if it is too large, the pressure will take a very long time to equilibrate. A good choice for many models is a *Pdamp* of around 1000 timesteps. However, note that *Pdamp* is specified in time units, and that timesteps are NOT the same as time units for most *units* settings.

---

The relaxation rate of the barostat is set by its inertia  $W$ :

$$W = (N + 1)k_B T_{\text{target}} P_{\text{damp}}^2$$

where  $N$  is the number of atoms,  $k_B$  is the Boltzmann constant, and  $T_{\text{target}}$  is the target temperature of the barostat (*Martyna*). If a thermostat is defined,  $T_{\text{target}}$  is the target temperature of the thermostat. If a thermostat is not defined,  $T_{\text{target}}$  is set to the current temperature of the system when the barostat is initialized. If this temperature is too low the simulation will quit with an error. Note: in previous versions of LAMMPS,  $T_{\text{target}}$  would default to a value of 1.0 for *lj* units and 300.0 otherwise if the system had a temperature of exactly zero.

If a thermostat is not specified by this fix,  $T_{\text{target}}$  can be manually specified using the *Ptemp* parameter. This may be useful if the barostat is initialized when the current temperature does not reflect the steady state temperature of the system. This keyword may also be useful in athermal simulations where the temperature is not well defined.



Regardless of what atoms are in the `fix` group (the only atoms which are time integrated), a global pressure or stress tensor is computed for all atoms. Similarly, when the size of the simulation box is changed, all atoms are re-scaled to new positions, unless the keyword `dilate` is specified with a `dilate-group-ID` for a group that represents a subset of the atoms. This can be useful, for example, to leave the coordinates of atoms in a solid substrate unchanged and controlling the pressure of a surrounding fluid. This option should be used with care, since it can be unphysical to dilate some atoms and not others, because it can introduce large, instantaneous displacements between a pair of atoms (one dilated, one not) that are far from the dilation origin. Also note that for atoms not in the `fix` group, a separate time integration `fix` like `fix nve` or `fix nvt` can be used on them, independent of whether they are dilated or not.

The `couple` keyword allows two or three of the diagonal components of the pressure tensor to be “coupled” together. The value specified with the keyword determines which are coupled. For example, `xz` means the  $P_{xx}$  and  $P_{zz}$  components of the stress tensor are coupled. `xyz` means all 3 diagonal components are coupled. Coupling means two things: the instantaneous stress will be computed as an average of the corresponding diagonal components, and the coupled box dimensions will be changed together in lockstep, meaning coupled dimensions will be dilated or contracted by the same percentage every timestep. The `Pstart`, `Pstop`, `Pdamp` parameters for any coupled dimensions must be identical. `Couple xyz` can be used for a 2d simulation; the `z` dimension is simply ignored.

The `iso`, `aniso`, and `tri` keywords are simply shortcuts that are equivalent to specifying several other keywords together.

The keyword `iso` means couple all 3 diagonal components together when pressure is computed (hydrostatic pressure), and dilate/contract the dimensions together. Using “`iso Pstart Pstop Pdamp`” is the same as specifying these 4 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
couple xyz
```

The keyword `aniso` means `x`, `y`, and `z` dimensions are controlled independently using the  $P_{xx}$ ,  $P_{yy}$ , and  $P_{zz}$  components of the stress tensor as the driving forces, and the specified scalar external pressure. Using “`aniso Pstart Pstop Pdamp`” is the same as specifying these 4 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
couple none
```

The keyword `tri` means `x`, `y`, `z`, `xy`, `xz`, and `yz` dimensions are controlled independently using their individual stress components as the driving forces, and the specified scalar pressure as the external normal stress. Using “`tri Pstart Pstop Pdamp`” is the same as specifying these 7 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
xy 0.0 0.0 Pdamp
yz 0.0 0.0 Pdamp
xz 0.0 0.0 Pdamp
couple none
```

In some cases (e.g. for solids) the pressure (volume) and/or temperature of the system can oscillate undesirably when a Nose/Hoover barostat and thermostat is applied. The optional `drag` keyword will damp these oscillations, although it alters the Nose/Hoover equations. A value of 0.0 (no drag) leaves the Nose/Hoover formalism unchanged. A non-zero value adds a drag term; the larger the value specified, the greater the damping effect. Performing a short run and

monitoring the pressure and temperature is the best way to determine if the drag term is working. Typically a value between 0.2 to 2.0 is sufficient to damp oscillations after a few periods. Note that use of the drag keyword will interfere with energy conservation and will also change the distribution of positions and velocities so that they do not correspond to the nominal NVT, NPT, or NPH ensembles.

An alternative way to control initial oscillations is to use chain thermostats. The keyword *tchain* determines the number of thermostats in the particle thermostat. A value of 1 corresponds to the original Nose-Hoover thermostat. The keyword *pchain* specifies the number of thermostats in the chain thermostating the barostat degrees of freedom. A value of 0 corresponds to no thermostating of the barostat variables.

The *mtk* keyword controls whether or not the correction terms due to Martyna, Tuckerman, and Klein are included in the equations of motion (*Martyna*). Specifying *no* reproduces the original Hoover barostat, whose volume probability distribution function differs from the true NPT and NPH ensembles by a factor of  $1/V$ . Hence using *yes* is more correct, but in many cases the difference is negligible.

The keyword *tloop* can be used to improve the accuracy of integration scheme at little extra cost. The initial and final updates of the thermostat variables are broken up into *tloop* sub-steps, each of length  $dt/tloop$ . This corresponds to using a first-order Suzuki-Yoshida scheme (*Tuckerman*). The keyword *ploop* does the same thing for the barostat thermostat.

The keyword *nreset* controls how often the reference dimensions used to define the strain energy are reset. If this keyword is not used, or is given a value of zero, then the reference dimensions are set to those of the initial simulation domain and are never changed. If the simulation domain changes significantly during the simulation, then the final average pressure tensor will differ significantly from the specified values of the external stress tensor. A value of *nstep* means that every *nstep* timesteps, the reference dimensions are set to those of the current simulation domain.

The *scaleyz*, *scalexz*, and *scalexy* keywords control whether or not the corresponding tilt factors are scaled with the associated box dimensions when barostatting triclinic periodic cells. The default values *yes* will turn on scaling, which corresponds to adjusting the linear dimensions of the cell while preserving its shape. Choosing *no* ensures that the tilt factors are not scaled with the box dimensions. See below for restrictions and default values in different situations. In older versions of LAMMPS, scaling of tilt factors was not performed. The old behavior can be recovered by setting all three scale keywords to *no*.

The *flip* keyword allows the tilt factors for a triclinic box to exceed half the distance of the parallel box length, as discussed below. If the *flip* value is set to *yes*, the bound is enforced by flipping the box when it is exceeded. If the *flip* value is set to *no*, the tilt will continue to change without flipping. Note that if applied stress induces large deformations (e.g. in a liquid), this means the box shape can tilt dramatically and LAMMPS will run less efficiently, due to the large volume of communication needed to acquire ghost atoms around a processor's irregular-shaped sub-domain. For extreme values of tilt, LAMMPS may also lose atoms and generate an error.

The *fixedpoint* keyword specifies the fixed point for barostat volume changes. By default, it is the center of the box. Whatever point is chosen will not move during the simulation. For example, if the lower periodic boundaries pass through (0,0,0), and this point is provided to *fixedpoint*, then the lower periodic boundaries will remain at (0,0,0), while the upper periodic boundaries will move twice as far. In all cases, the particle trajectories are unaffected by the chosen value, except for a time-dependent constant translation of positions.

If the *update* keyword is used with the *dipole* value, then the orientation of the dipole moment of each particle is also updated during the time integration. This option should be used for models where a dipole moment is assigned to finite-size particles, e.g. spheroids via use of the *atom\_style hybrid sphere dipole* command.

The default dipole orientation integrator can be changed to the Dullweber-Leimkuhler-McLachlan integration scheme (*Dullweber*) when using *update* with the value *dipole/dlm*. This integrator is symplectic and time-reversible, giving better energy conservation and allows slightly longer timesteps at only a small additional computational cost.

---

**Note:** Using a barostat coupled to tilt dimensions *xy*, *xz*, *yz* can sometimes result in arbitrarily large values of the tilt dimensions, i.e. a dramatically deformed simulation box. LAMMPS allows the tilt factors to grow a small amount

beyond the normal limit of half the box length (0.6 times the box length), and then performs a box “flip” to an equivalent periodic cell. See the discussion of the *flip* keyword above, to allow this bound to be exceeded, if desired.

The flip operation is described in more detail in the page for *fix deform*. Both the barostat dynamics and the atom trajectories are unaffected by this operation. However, if a tilt factor is incremented by a large amount (1.5 times the box length) on a single timestep, LAMMPS can not accommodate this event and will terminate the simulation with an error. This error typically indicates that there is something badly wrong with how the simulation was constructed, such as specifying values of *Pstart* that are too far from the current stress value, or specifying a timestep that is too large. Triclinic barostatting should be used with care. This also is true for other barostat styles, although they tend to be more forgiving of insults. In particular, it is important to recognize that equilibrium liquids can not support a shear stress and that equilibrium solids can not support shear stresses that exceed the yield stress.

One exception to this rule is if the first dimension in the tilt factor (x for xy) is non-periodic. In that case, the limits on the tilt factor are not enforced, since flipping the box in that dimension does not change the atom positions due to non-periodicity. In this mode, if you tilt the system to extreme angles, the simulation will simply become inefficient due to the highly skewed simulation box.

**Note:** Unlike the *fix temp/berendsen* command which performs thermostating but NO time integration, these fixes perform thermostating/barostatting AND time integration. Thus you should not use any other time integration fix, such as *fix nve* on atoms to which this fix is applied. Likewise, *fix nvt* and *fix npt* should not normally be used on atoms that also have their temperature controlled by another fix - e.g. by *fix langevin* or *fix temp/rescale* commands.

See the *Howto thermostat* and *Howto barostat* doc pages for a discussion of different ways to compute temperature and perform thermostating and barostatting.

These fixes compute a temperature and pressure each timestep. To do this, the thermostat and barostat fixes create their own computes of style “temp” and “pressure”, as if one of these sets of commands had been issued:

For *fix nvt*:

```
compute fix-ID_temp group-ID temp
```

For *fix npt* and *fix nph*:

```
compute fix-ID_temp all temp
compute fix-ID_press all pressure fix-ID_temp
```

For *fix nvt*, the group for the new temperature compute is the same as the fix group. For *fix npt* and *fix nph*, the group for both the new temperature and pressure compute is “all” since pressure is computed for the entire system. In the case of *fix nph*, the temperature compute is not used for thermostating, but just for a kinetic-energy contribution to the pressure. See the *compute temp* and *compute pressure* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix\_ID + underscore + “press”.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of these fix’s temperature or pressure via the *compute\_modify* command. Or you can print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include

a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

These fixes can be used with either the *verlet* or *respa integrators*. When using one of the barostat fixes with *respa*, LAMMPS uses an integrator constructed according to the following factorization of the Liouville propagator (for two rRESPA levels):

$$\begin{aligned} \exp(iL\Delta t) = & \hat{E} \exp\left(iL_{T\text{-baro}}\frac{\Delta t}{2}\right) \exp\left(iL_{T\text{-part}}\frac{\Delta t}{2}\right) \exp\left(iL_{\epsilon,2}\frac{\Delta t}{2}\right) \exp\left(iL_2^{(2)}\frac{\Delta t}{2}\right) \\ & \times \left[ \exp\left(iL_2^{(1)}\frac{\Delta t}{2n}\right) \exp\left(iL_{\epsilon,1}\frac{\Delta t}{2n}\right) \exp\left(iL_1\frac{\Delta t}{n}\right) \exp\left(iL_{\epsilon,1}\frac{\Delta t}{2n}\right) \exp\left(iL_2^{(1)}\frac{\Delta t}{2n}\right) \right]^n \\ & \times \exp\left(iL_2^{(2)}\frac{\Delta t}{2}\right) \exp\left(iL_{\epsilon,2}\frac{\Delta t}{2}\right) \exp\left(iL_{T\text{-part}}\frac{\Delta t}{2}\right) \exp\left(iL_{T\text{-baro}}\frac{\Delta t}{2}\right) \\ & + \mathcal{O}(\Delta t^3) \end{aligned}$$

This factorization differs somewhat from that of Tuckerman et al, in that the barostat is only updated at the outermost rRESPA level, whereas Tuckerman's factorization requires splitting the pressure into pieces corresponding to the forces computed at each rRESPA level. In theory, the latter method will exhibit better numerical stability. In practice, because Pdamp is normally chosen to be a large multiple of the outermost rRESPA timestep, the barostat dynamics are not the limiting factor for numerical stability. Both factorizations are time-reversible and can be shown to preserve the phase space measure of the underlying non-Hamiltonian equations of motion.

---

**Note:** This implementation has been shown to conserve linear momentum up to machine precision under NVT dynamics. Under NPT dynamics, for a system with zero initial total linear momentum, the total momentum fluctuates close to zero. It may occasionally undergo brief excursions to non-negligible values, before returning close to zero. Over long simulations, this has the effect of causing the center-of-mass to undergo a slow random walk. This can be mitigated by resetting the momentum at infrequent intervals using the *fix momentum* command.

---

The *fix npt* and *fix nph* commands can be used with rigid bodies or mixtures of rigid bodies and non-rigid particles (e.g. solvent). But there are also *fix rigid/npt* and *fix rigid/nph* commands, which are typically a more natural choice. See the page for those commands for more discussion of the various ways to do this.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 17.101.4 Restart, fix\_modify, output, run start/stop, minimize info

These fixes writes the state of all the thermostat and barostat variables to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* and *press* options are supported by these fixes. You can use them to assign a *compute* you have defined to this fix which will be used in its thermostating or barostating procedure, as described above. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

---

**Note:** If both the *temp* and *press* keywords are used in a single *thermo\_modify* command (or in two separate commands), then the order in which the keywords are specified is important. Note that a *pressure compute* defines its own temperature compute as an argument when it is specified. The *temp* keyword will override this (for the pressure compute being used by fix npt), but only if the *temp* keyword comes after the *press* keyword. If the *temp* keyword comes before the *press* keyword, then the new pressure compute specified by the *press* keyword will be unaffected by the *temp* setting.

---

The cumulative energy change in the system imposed by these fixes, via either thermostating and/or barostating, is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* page for details.

These fixes compute a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

These fixes compute also compute a global vector of quantities, which can be accessed by various *output commands*. The vector values are “intensive”.

The vector stores internal Nose/Hoover thermostat and barostat variables. The number and meaning of the vector values depends on which fix is used and the settings for keywords *tchain* and *pchain*, which specify the number of Nose/Hoover chains for the thermostat and barostat. If no thermostating is done, then *tchain* is 0. If no barostating is done, then *pchain* is 0. In the following list, “ndof” is 0, 1, 3, or 6, and is the number of degrees of freedom in the barostat. Its value is 0 if no barostat is used, else its value is 6 if any off-diagonal stress tensor component is barostated, else its value is 1 if *couple xyz* is used or *couple xy* for a 2d simulation, otherwise its value is 3.

The order of values in the global vector and their meaning is as follows. The notation means there are *tchain* values for *eta*, followed by *tchain* for *eta\_dot*, followed by *ndof* for *omega*, etc:

- $\text{eta}[\text{tchain}]$  = particle thermostat displacements (unitless)
- $\text{eta\_dot}[\text{tchain}]$  = particle thermostat velocities (1/time units)
- $\text{omega}[\text{ndof}]$  = barostat displacements (unitless)
- $\text{omega\_dot}[\text{ndof}]$  = barostat velocities (1/time units)
- $\text{etap}[\text{pchain}]$  = barostat thermostat displacements (unitless)
- $\text{etap\_dot}[\text{pchain}]$  = barostat thermostat velocities (1/time units)
- $\text{PE\_eta}[\text{tchain}]$  = potential energy of each particle thermostat displacement (energy units)
- $\text{KE\_eta\_dot}[\text{tchain}]$  = kinetic energy of each particle thermostat velocity (energy units)
- $\text{PE\_omega}[\text{ndof}]$  = potential energy of each barostat displacement (energy units)
- $\text{KE\_omega\_dot}[\text{ndof}]$  = kinetic energy of each barostat velocity (energy units)
- $\text{PE\_etap}[\text{pchain}]$  = potential energy of each barostat thermostat displacement (energy units)
- $\text{KE\_etap\_dot}[\text{pchain}]$  = kinetic energy of each barostat thermostat velocity (energy units)

- PE\_strain[1] = scalar strain energy (energy units)

These fixes can ramp their external temperature and pressure over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

These fixes are not invoked during *energy minimization*.

---

### 17.101.5 Restrictions

*X*, *y*, *z* cannot be barostatted if the associated dimension is not periodic. *Xy*, *xz*, and *yz* can only be barostatted if the simulation domain is triclinic and the second dimension in the keyword (*y* dimension in *xy*) is periodic. *Z*, *xz*, and *yz*, cannot be barostatted for 2D simulations. The *create\_box*, *read\_data*, and *read\_restart* commands specify whether the simulation box is orthogonal or non-orthogonal (triclinic) and explain the meaning of the *xy,xz,yz* tilt factors.

For the *temp* keyword, the final *Tstop* cannot be 0.0 since it would make the external *T* = 0.0 at some timestep during the simulation which is not allowed in the Nose/Hoover formulation.

The *scaleyz yes* and *scalexz yes* keyword/value pairs can not be used for 2D simulations. *scaleyz yes*, *scalexz yes*, and *scalexy yes* options can only be used if the second dimension in the keyword is periodic, and if the tilt factor is not coupled to the barostat via keywords *tri*, *yz*, *xz*, and *xy*.

These fixes can be used with dynamic groups as defined by the *group* command. Likewise they can be used with groups to which atoms are added or deleted over time, e.g. a deposition simulation. However, the conservation properties of the thermostat and barostat are defined for systems with a static set of atoms. You may observe odd behavior if the atoms in a group vary dramatically over time or the atom count becomes very small.

### 17.101.6 Related commands

*fix nve*, *fix\_modify*, *run\_style*

### 17.101.7 Default

The keyword defaults are *tchain* = 3, *pchain* = 3, *mtk* = yes, *tloop* = 1, *ploop* = 1, *nreset* = 0, *drag* = 0.0, *dilate* = all, *couple* = none, *flip* = yes, *scaleyz* = *scalexz* = *scalexy* = yes if periodic in second dimension and not coupled to barostat, otherwise no.

---

**(Martyna)** Martyna, Tobias and Klein, J Chem Phys, 101, 4177 (1994).

**(Parrinello)** Parrinello and Rahman, J Appl Phys, 52, 7182 (1981).

**(Tuckerman)** Tuckerman, Alejandre, Lopez-Rendon, Jochim, and Martyna, J Phys A: Math Gen, 39, 5629 (2006).

**(Shinoda)** Shinoda, Shiga, and Mikami, Phys Rev B, 69, 134103 (2004).

**(Dullweber)** Dullweber, Leimkuhler and McLachlan, J Chem Phys, 107, 5840 (1997).

## 17.102 fix nvt/eff command

## 17.103 fix npt/eff command

## 17.104 fix nph/eff command

### 17.104.1 Syntax

```
fix ID group-ID style_name keyword value ...
```

- ID, group-ID are documented in *fix* command
- style\_name = *nvt/eff* or *npt/eff* or *nph/eff*

one or more keyword value pairs may be appended

keyword = *temp* or *iso* or *aniso* or *tri* or *x* or *y* or *z* or *xy* or *yz* or *xz* or *couple* or *tchain* or *pchain* or *mtk* or *tloop* or *ploop* or *nreset* or *drag* or *dilate*

*temp* values = Tstart Tstop Tdamp

Tstart,Tstop = external temperature at start/end of run

Tdamp = temperature damping parameter (time units)

*iso* or *aniso* or *tri* values = Pstart Pstop Pdamp

Pstart,Pstop = scalar external pressure at start/end of run (pressure units)

Pdamp = pressure damping parameter (time units)

*x* or *y* or *z* or *xy* or *yz* or *xz* values = Pstart Pstop Pdamp

Pstart,Pstop = external stress tensor component at start/end of run (pressure units)

Pdamp = stress damping parameter (time units)

*couple* = *none* or *xyz* or *xy* or *yz* or *xz*

*tchain* value = length of thermostat chain (1 = single thermostat)

*pchain* values = length of thermostat chain on barostat (0 = no thermostat)

*mtk* value = *yes* or *no* = add in MTK adjustment term or not

*tloop* value = number of sub-cycles to perform on thermostat

*ploop* value = number of sub-cycles to perform on barostat thermostat

*nreset* value = reset reference cell every this many timesteps

*drag* value = drag factor added to barostat/thermostat (0.0 = no drag)

*dilate* value = *all* or *partial*

### 17.104.2 Examples

```
fix 1 all nvt/eff temp 300.0 300.0 0.1
fix 1 part npt/eff temp 300.0 300.0 0.1 iso 0.0 0.0 1.0
fix 2 part npt/eff temp 300.0 300.0 0.1 tri 5.0 5.0 1.0
fix 2 ice nph/eff x 1.0 1.0 0.5 y 2.0 2.0 0.5 z 3.0 3.0 0.5 yz 0.1 0.1 0.5 xz 0.2 0.2 0.
↪5 xy 0.3 0.3 0.5 nreset 1000
```

### 17.104.3 Description

These commands perform time integration on Nose-Hoover style non-Hamiltonian equations of motion for nuclei and electrons in the group for the *electron force field* model. The fixes are designed to generate positions and velocities sampled from the canonical (nvt), isothermal-isobaric (npt), and isenthalpic (nph) ensembles. This is achieved by adding some dynamic variables which are coupled to the particle velocities (thermostatting) and simulation domain dimensions (barostatting). In addition to basic thermostatting and barostatting, these fixes can also create a chain of thermostats coupled to the particle thermostat, and another chain of thermostats coupled to the barostat variables. The barostat can be coupled to the overall box volume, or to individual dimensions, including the *xy*, *xz* and *yz* tilt dimensions. The external pressure of the barostat can be specified as either a scalar pressure (isobaric ensemble) or as components of a symmetric stress tensor (constant stress ensemble). When used correctly, the time-averaged temperature and stress tensor of the particles will match the target values specified by Tstart/Tstop and Pstart/Pstop.

The operation of these fixes is exactly like that described by the *fix nvt*, *npt*, and *nph* commands, except that the radius and radial velocity of electrons are also updated. Likewise the temperature and pressure calculated by the fix, using the computes it creates (as discussed in the *fix nvt*, *npt*, and *nph* doc page), are performed with computes that include the eFF contribution to the temperature or kinetic energy from the electron radial velocity.

---

**Note:** there are two different pressures that can be reported for eFF when defining the *pair\_style* (see *pair eff/cut* to understand these settings), one (default) that considers electrons do not contribute radial virial components (i.e. electrons treated as incompressible ‘rigid’ spheres) and one that does. The radial electronic contributions to the virials are only tallied if the flexible pressure option is set, and this will affect both global and per-atom quantities. In principle, the true pressure of a system is somewhere in between the rigid and the flexible eFF pressures, but, for most cases, the difference between these two pressures will not be significant over long-term averaged runs (i.e. even though the energy partitioning changes, the total energy remains similar).

---

---

**Note:** currently, there is no available option for the user to set or create temperature distributions that include the radial electronic degrees of freedom with the *velocity* command, so the user must allow for these degrees of freedom to equilibrate (i.e. equi-partitioning of energy) through time integration.

---

### 17.104.4 Restart, fix\_modify, output, run start/stop, minimize info

See the page for the *fix nvt*, *npt*, and *nph* commands for details.

### 17.104.5 Restrictions

This fix is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Other restriction discussed on the page for the *fix nvt*, *npt*, and *nph* commands also apply.

---

**Note:** The temperature for systems (regions or groups) with only electrons and no nuclei is 0.0 (i.e. not defined) in the current temperature calculations, a practical example would be a uniform electron gas or a very hot plasma, where electrons remain delocalized from the nuclei. This is because, even though electron virials are included in the temperature calculation, these are averaged over the nuclear degrees of freedom only. In such cases a corrective term must be added to the pressure to get the correct kinetic contribution.

---



## 17.104.6 Related commands

*fix nvt*, *fix nph*, *fix npt*, *fix\_modify*, *run\_style*

## 17.104.7 Default

The keyword defaults are *tchain* = 3, *pchain* = 3, *mtk* = yes, *tloop* = *ploop* = 1, *nreset* = 0, *drag* = 0.0, *dilate* = all, and *couple* = none.

(**Martyna**) Martyna, Tobias and Klein, J Chem Phys, 101, 4177 (1994).

(**Parrinello**) Parrinello and Rahman, J Appl Phys, 52, 7182 (1981).

(**Tuckerman**) Tuckerman, Alejandre, Lopez-Rendon, Jochim, and Martyna, J Phys A: Math Gen, 39, 5629 (2006).

(**Shinoda**) Shinoda, Shiga, and Mikami, Phys Rev B, 69, 134103 (2004).

## 17.105 fix nvt/uef command

## 17.106 fix npt/uef command

### 17.106.1 Syntax

```
fix ID group-ID style_name erate edot_x edot_y temp Tstart Tstop Tdamp keyword value ...
```

- ID, group-ID are documented in *fix* command
- style\_name = *nvt/uef* or *npt/uef*
- *Tstart*, *Tstop*, and *Tdamp* are documented in the *fix npt* command
- *edot\_x* and *edot\_y* are the strain rates in the x and y directions (1/(time units))
- one or more keyword/value pairs may be appended

keyword = *erate* or *ext* or *strain* or *temp* or *iso* or *x* or *y* or *z* or *tchain* or *pchain*,  
 ↪ or *tloop* or *ploop* or *mtk*

*erate* values = *e\_x* *e\_y* = engineering strain rates (required)

*ext* value = *x* or *y* or *z* or *xy* or *yz* or *xz* = external dimensions

sets the external dimensions used to calculate the scalar pressure

*strain* values = *e\_x* *e\_y* = initial strain

usually not needed, but may be needed to resume a run with a data file.

*temp*, *iso*, *x*, *y*, *z*, *tchain*, *pchain*, *tloop*, *ploop*, *mtk*

keywords documented by the *fix npt* command

## 17.106.2 Examples

```

fix uniax_nvt all nvt/uef temp 400 400 100 erate 0.00001 -0.000005
fix biax_nvt all nvt/uef temp 400 400 100 erate 0.000005 0.000005
fix uniax_npt all npt/uef temp 400 400 300 iso 1 1 3000 erate 0.00001 -0.000005 ext yz
fix biax_npt all npt/uef temp 400 400 100 erate -0.00001 0.000005 x 1 1 3000

```

## 17.106.3 Description

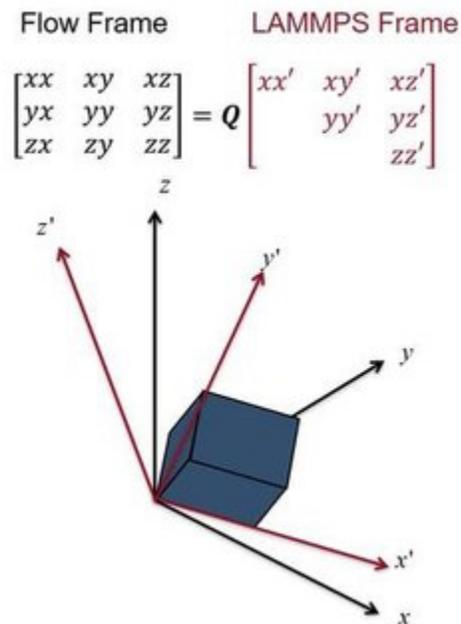
This fix can be used to simulate non-equilibrium molecular dynamics (NEMD) under diagonal flow fields, including uniaxial and bi-axial flow. Simulations under continuous extensional flow may be carried out for an indefinite amount of time. It is an implementation of the boundary conditions from (*Dobson*), and also uses numerical lattice reduction as was proposed by (*Hunt*). The lattice reduction algorithm is from (*Semaev*). The fix is intended for simulations of homogeneous flows, and integrates the SLLOD equations of motion, originally proposed by Hoover and Ladd (see (*Evans and Morris*)). Additional detail about this implementation can be found in (*Nicholson and Rutledge*).

Note that NEMD simulations of a continuously strained system can be performed using the *fix deform*, *fix nvt/sllod*, and *compute temp/deform* commands.

The applied flow field is set by the *eps* keyword. The values *edot\_x* and *edot\_y* correspond to the strain rates in the *xx* and *yy* directions. It is implicitly assumed that the flow field is traceless, and therefore the strain rate in the *zz* direction is equal to  $-(edot_x + edot_y)$ .

**Note:** Due to an instability in the SLLOD equations under extension, *fix momentum* should be used to regularly reset the linear momentum.

The boundary conditions require a simulation box that does not have a consistent alignment relative to the applied flow field. Since LAMMPS utilizes an upper-triangular simulation box, it is not possible to express the evolving simulation box in the same coordinate system as the flow field. This fix keeps track of two coordinate systems: the flow frame, and the upper triangular LAMMPS frame. The coordinate systems are related to each other through the QR decomposition, as is illustrated in the image below.



During most molecular dynamics operations, the system is represented in the LAMMPS frame. Only when the positions and velocities are updated is the system rotated to the flow frame, and it is rotated back to the LAMMPS frame immediately afterwards. For this reason, all vector-valued quantities (except for the tensors from *compute pressure/uef* and *compute temp/uef*) will be computed in the LAMMPS frame. Rotationally invariant scalar quantities like the temperature and hydrostatic pressure are frame-invariant and will be computed correctly. Additionally, the system is in the LAMMPS frame during all of the output steps, and therefore trajectory files made using the dump command will be in the LAMMPS frame unless the *dump cfg/uef* command is used.

Temperature control is achieved with the default Nose-Hoover style thermostat documented in *fix npt*. When this fix is active, only the peculiar velocity of each atom is stored, defined as the velocity relative to the streaming velocity. This is in contrast to *fix nvt/sllod*, which uses a lab-frame velocity, and removes the contribution from the streaming velocity in order to compute the temperature.

Pressure control is achieved using the default Nose-Hoover barostat documented in *fix npt*. There are two ways to control the pressure using this fix. The first method involves using the *ext* keyword along with the *iso* pressure style. With this method, the pressure is controlled by scaling the simulation box isotropically to achieve the average pressure only in the directions specified by *ext*. For example, if the *ext* value is set to *xy*, the average pressure  $(P_{xx}+P_{yy})/2$  will be controlled.

This example command will control the total hydrostatic pressure under uniaxial tension:

```
fix f1 all npt/uef temp 0.7 0.7 0.5 iso 1 1 5 erate -0.5 -0.5 ext xyz
```

This example command will control the average stress in compression directions, which would typically correspond to free surfaces under drawing with uniaxial tension:

```
fix f2 all npt/uef temp 0.7 0.7 0.5 iso 1 1 5 erate -0.5 -0.5 ext xy
```

The second method for pressure control involves setting the normal stresses using the *x*, *y*, and/or *z* keywords. When using this method, the same pressure must be specified via *Pstart* and *Pstop* for all dimensions controlled. Any choice of pressure conditions that would cause LAMMPS to compute a deviatoric stress are not permissible and will result in an error. Additionally, all dimensions with controlled stress must have the same applied strain rate. The *ext* keyword must be set to the default value (*xyz*) when using this method.

For example, the following commands will work:

```
fix f3 all npt/uef temp 0.7 0.7 0.5 x 1 1 5 y 1 1 5 erate -0.5 -0.5
fix f4 all npt/uef temp 0.7 0.7 0.5 z 1 1 5 erate 0.5 0.5
```

The following commands will not work:

```
fix f5 all npt/uef temp 0.7 0.7 0.5 x 1 1 5 z 1 1 5 erate -0.5 -0.5
fix f6 all npt/uef temp 0.7 0.7 0.5 x 1 1 5 z 2 2 5 erate 0.5 0.5
```

These fix computes a temperature and pressure each timestep. To do this, it creates its own computes of style “temp/uef” and “pressure/uef”, as if one of these two sets of commands had been issued:

```
compute fix-ID_temp group-ID temp/uef
compute fix-ID_press group-ID pressure/uef fix-ID_temp

compute fix-ID_temp all temp/uef
compute fix-ID_press all pressure/uef fix-ID_temp
```

See the *compute temp/uef* and *compute pressure/uef* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix\_ID + underscore + “press”.

### 17.106.4 Restart, fix\_modify, output, run start/stop, minimize info

The fix writes the state of all the thermostat and barostat variables, as well as the cumulative strain applied, to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

---

**Note:** It is not necessary to set the *strain* keyword when resuming a run from a restart file. Only for resuming from data files, which do not contain the cumulative applied strain, will this keyword be necessary.

---

This fix can be used with the *fix\_modify temp* and *press* options. The temperature and pressure computes used must be of type *temp/uef* and *pressure/uef*.

This fix computes the same global scalar and vector quantities as *fix npt*.

The fix is not invoked during *energy minimization*.

### 17.106.5 Restrictions

This fix is part of the UEF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Due to requirements of the boundary conditions, when the *strain* keyword is set to zero (or unset), the initial simulation box must be cubic and have style triclinic. If the box is initially of type ortho, use *change\_box* before invoking the fix.

---

**Note:** When resuming from restart files, you may need to use *box tilt large* since lammps has internal criteria from lattice reduction that are not the same as the criteria in the numerical lattice reduction algorithm.

---

### 17.106.6 Related commands

*fix nvt*, *fix nvt/sllod*, *compute temp/uef*, *compute pressure/uef*, *dump cfg/uef*

### 17.106.7 Default

The default keyword values specific to this fix are *exy = xyz*, *strain = 0*. The remaining defaults are the same as for *fix npt* except *tchain = 1*. The reason for this change is given in *fix nvt/sllod*.

---

**(Dobson)** Dobson, J Chem Phys, 141, 184103 (2014).

**(Hunt)** Hunt, Mol Simul, 42, 347 (2016).

**(Semaev)** Semaev, Cryptography and Lattices, 181 (2001).

**(Evans and Morriss)** Evans and Morriss, Phys Rev A, 30, 1528 (1984).

**(Nicholson and Rutledge)** Nicholson and Rutledge, J Chem Phys, 145, 244903 (2016).

## 17.107 fix nph/asphere command

Accelerator Variants: *nph/asphere/omp*

### 17.107.1 Syntax

```
fix ID group-ID nph/asphere args keyword value ...
```

- ID, group-ID are documented in *fix* command
- nph/asphere = style name of this fix command
- additional barostat related keyword/value pairs from the *fix nph* command can be appended

### 17.107.2 Examples

```
fix 1 all nph/asphere iso 0.0 0.0 1000.0
fix 2 all nph/asphere x 5.0 5.0 1000.0
fix 2 all nph/asphere x 5.0 5.0 1000.0 drag 0.2
fix 2 water nph/asphere aniso 0.0 0.0 1000.0 dilate partial
```

### 17.107.3 Description

Perform constant NPH integration to update position, velocity, orientation, and angular velocity each timestep for aspherical or ellipsoidal particles in the group using a Nose/Hoover pressure barostat. P is pressure; H is enthalpy. This creates a system trajectory consistent with the isenthalpic ensemble.

This fix differs from the *fix nph* command, which assumes point particles and only updates their position and velocity.

Additional parameters affecting the barostat are specified by keywords and values documented with the *fix nph* command. See, for example, discussion of the *aniso*, and *dilate* keywords.

The particles in the fix group are the only ones whose velocities and positions are updated by the velocity/position update portion of the NPH integration.

Regardless of what particles are in the fix group, a global pressure is computed for all particles. Similarly, when the size of the simulation box is changed, all particles are re-scaled to new positions, unless the keyword *dilate* is specified with a value of *partial*, in which case only the particles in the fix group are re-scaled. The latter can be useful for leaving the coordinates of particles in a solid substrate unchanged and controlling the pressure of a surrounding fluid.

This fix computes a temperature and pressure each timestep. To do this, the fix creates its own computes of style “temp/asphere” and “pressure”, as if these commands had been issued:

```
compute fix-ID_temp all temp/asphere
compute fix-ID_press all pressure fix-ID_temp
```

See the *compute temp/asphere* and *compute pressure* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix\_ID + underscore + “press”, and the group for the new computes is “all” since pressure is computed for the entire system.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of this fix’s temperature or pressure via the

*compute\_modify* command or print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this fix.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 17.107.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover barostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* and *press* options are supported by this fix. You can use them to assign a *compute* you have defined to this fix which will be used in its thermostating or barostating procedure. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix nph* command.

This fix can ramp its target pressure over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

#### 17.107.5 Restrictions

This fix is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular momentum and a quaternion as defined by the *atom\_style ellipsoid* command.

All particles in the group must be finite-size. They cannot be point particles, but they can be aspherical or spherical as defined by their shape attribute.

## 17.107.6 Related commands

*fix nph*, *fix nve\_asphere*, *fix nvt\_asphere*, *fix npt\_asphere*, *fix\_modify*

## 17.107.7 Default

none

## 17.108 fix nph/body command

### 17.108.1 Syntax

```
fix ID group-ID nph/body args keyword value ...
```

- ID, group-ID are documented in *fix* command
- nph/body = style name of this fix command
- additional barostat related keyword/value pairs from the *fix nph* command can be appended

### 17.108.2 Examples

```
fix 1 all nph/body iso 0.0 0.0 1000.0
fix 2 all nph/body x 5.0 5.0 1000.0
fix 2 all nph/body x 5.0 5.0 1000.0 drag 0.2
fix 2 water nph/body aniso 0.0 0.0 1000.0 dilate partial
```

### 17.108.3 Description

Perform constant NPH integration to update position, velocity, orientation, and angular velocity each timestep for body particles in the group using a Nose/Hoover pressure barostat. P is pressure; H is enthalpy. This creates a system trajectory consistent with the isenthalpic ensemble.

This fix differs from the *fix nph* command, which assumes point particles and only updates their position and velocity.

Additional parameters affecting the barostat are specified by keywords and values documented with the *fix nph* command. See, for example, discussion of the *aniso*, and *dilate* keywords.

The particles in the fix group are the only ones whose velocities and positions are updated by the velocity/position update portion of the NPH integration.

Regardless of what particles are in the fix group, a global pressure is computed for all particles. Similarly, when the size of the simulation box is changed, all particles are re-scaled to new positions, unless the keyword *dilate* is specified with a value of *partial*, in which case only the particles in the fix group are re-scaled. The latter can be useful for leaving the coordinates of particles in a solid substrate unchanged and controlling the pressure of a surrounding fluid.

This fix computes a temperature and pressure each timestep. To do this, the fix creates its own computes of style “temp/body” and “pressure”, as if these commands had been issued:

```
compute fix-ID_temp all temp/body
compute fix-ID_press all pressure fix-ID_temp
```

See the *compute temp/body* and *compute pressure* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix\_ID + underscore + “press”, and the group for the new computes is “all” since pressure is computed for the entire system.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of this fix’s temperature or pressure via the *compute\_modify* command or print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this fix.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 17.108.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover barostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* and *press* options are supported by this fix. You can use them to assign a *compute* you have defined to this fix which will be used in its thermostating or barostating procedure. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix nph* command.

This fix can ramp its target pressure over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.



### 17.108.5 Restrictions

This fix is part of the BODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular momentum and a quaternion as defined by the *atom\_style body* command.

### 17.108.6 Related commands

*fix nph*, *fix nve\_body*, *fix nvt\_body*, *fix npt\_body*, *fix\_modify*

### 17.108.7 Default

none

## 17.109 fix nph/sphere command

Accelerator Variants: *nph/sphere/omp*

### 17.109.1 Syntax

```
fix ID group-ID nph/sphere args keyword value ...
```

- ID, group-ID are documented in *fix* command
- nph/sphere = style name of this fix command
- keyword = *disc*
  - disc* value = none = treat particles as 2d discs, not spheres
- additional barostat related keyword/value pairs from the *fix nph* command can be appended

### 17.109.2 Examples

```
fix 1 all nph/sphere iso 0.0 0.0 1000.0
fix 2 all nph/sphere x 5.0 5.0 1000.0
fix 2 all nph/sphere x 5.0 5.0 1000.0 disc
fix 2 all nph/sphere x 5.0 5.0 1000.0 drag 0.2
fix 2 water nph/sphere aniso 0.0 0.0 1000.0 dilate partial
```

### 17.109.3 Description

Perform constant NPH integration to update position, velocity, and angular velocity each timestep for finite-size spherical particles in the group using a Nose/Hoover pressure barostat. P is pressure; H is enthalpy. This creates a system trajectory consistent with the isenthalpic ensemble.

This fix differs from the *fix nph* command, which assumes point particles and only updates their position and velocity.

If the *disc* keyword is used, then each particle is treated as a 2d disc (circle) instead of as a sphere. This is only possible for 2d simulations, as defined by the *dimension* keyword. The only difference between discs and spheres in this context is their moment of inertia, as used in the time integration.

Additional parameters affecting the barostat are specified by keywords and values documented with the *fix nph* command. See, for example, discussion of the *aniso*, and *dilate* keywords.

The particles in the fix group are the only ones whose velocities and positions are updated by the velocity/position update portion of the NPH integration.

Regardless of what particles are in the fix group, a global pressure is computed for all particles. Similarly, when the size of the simulation box is changed, all particles are re-scaled to new positions, unless the keyword *dilate* is specified with a value of *partial*, in which case only the particles in the fix group are re-scaled. The latter can be useful for leaving the coordinates of particles in a solid substrate unchanged and controlling the pressure of a surrounding fluid.

---

This fix computes a temperature and pressure each timestep. To do this, the fix creates its own computes of style “temp/sphere” and “pressure”, as if these commands had been issued:

```
compute fix-ID_temp all temp/sphere
compute fix-ID_press all pressure fix-ID_temp
```

See the *compute temp/sphere* and *compute pressure* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix\_ID + underscore + “press”, and the group for the new computes is “all” since pressure is computed for the entire system.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of this fix’s temperature or pressure via the *compute\_modify* command or print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this fix.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

### 17.109.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover barostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* and *press* options are supported by this fix. You can use them to assign a *compute* you have defined to this fix which will be used in its thermostating or barostating procedure. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix nph* command.

This fix can ramp its target pressure over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

### 17.109.5 Restrictions

This fix requires that atoms store torque and angular velocity (*omega*) and a radius as defined by the *atom\_style sphere* command.

All particles in the group must be finite-size spheres. They cannot be point particles.

Use of the *disc* keyword is only allowed for 2d simulations, as defined by the *dimension* keyword.

### 17.109.6 Related commands

*fix nph*, *fix nve\_sphere*, *fix nvt\_sphere*, *fix npt\_sphere*, *fix\_modify*

### 17.109.7 Default

none

## 17.110 fix nphug command

Accelerator Variants: *nphug/omp*

### 17.110.1 Syntax

```
fix ID group-ID nphug keyword value ...
```

- ID, group-ID are documented in *fix* command

one or more keyword value pairs may be appended

keyword = *temp* or *iso* or *aniso* or *tri* or *x* or *y* or *z* or *couple* or *tchain* or *pchain*.

↪ or *mtk* or *tloop* or *ploop* or *nreset* or *drag* or *dilate* or *scaleyz* or *scalexz* or.

↪ *scalexy*

```

temp values = Value1 Value2 Tdamp
  Value1, Value2 = Nose-Hoover target temperatures, ignored by Hugoniostat
  Tdamp = temperature damping parameter (time units)
iso or aniso or tri values = Pstart Pstop Pdamp
  Pstart,Pstop = scalar external pressures, must be equal (pressure units)
  Pdamp = pressure damping parameter (time units)
x or y or z or xy or yz or xz values = Pstart Pstop Pdamp
  Pstart,Pstop = external stress tensor components, must be equal (pressure units)
  Pdamp = stress damping parameter (time units)
couple = none or xyz or xy or yz or xz
tchain value = length of thermostat chain (1 = single thermostat)
pchain values = length of thermostat chain on barostat (0 = no thermostat)
mtk value = yes or no = add in MTK adjustment term or not
tloop value = number of sub-cycles to perform on thermostat
ploop value = number of sub-cycles to perform on barostat thermostat
nreset value = reset reference cell every this many timesteps
drag value = drag factor added to barostat/thermostat (0.0 = no drag)
dilate value = all or partial
scaleyz value = yes or no = scale yz with lz
scalexz value = yes or no = scale xz with lz
scalexy value = yes or no = scale xy with ly

```

## 17.110.2 Examples

```

fix myhug all nphug temp 1.0 1.0 10.0 z 40.0 40.0 70.0
fix myhug all nphug temp 1.0 1.0 10.0 iso 40.0 40.0 70.0 drag 200.0 tchain 1 pchain 0

```

## 17.110.3 Description

This command is a variant of the Nose-Hoover *fix npt* fix style. It performs time integration of the Hugoniostat equations of motion developed by Ravelo et al. (*Ravelo*). These equations compress the system to a state with average axial stress or pressure equal to the specified target value and that satisfies the Rankine-Hugoniot (RH) jump conditions for steady shocks.

The compression can be performed either hydrostatically (using keyword *iso*, *aniso*, or *tri*) or uniaxially (using keywords *x*, *y*, or *z*). In the hydrostatic case, the cell dimensions change dynamically so that the average axial stress in all three directions converges towards the specified target value. In the uniaxial case, the chosen cell dimension changes dynamically so that the average axial stress in that direction converges towards the target value. The other two cell dimensions are kept fixed (zero lateral strain).

This leads to the following additional restrictions on the keywords:

- One and only one of the following keywords should be used: *iso*, *aniso*, *tri*, *x*, *y*, *z*
- The specified initial and final target pressures must be the same.
- The keywords *xy*, *xz*, *yz* may not be used.
- The only admissible value for the couple keyword is *xyz*, which has the same effect as keyword *iso*
- The *temp* keyword must be used to specify the time constant for kinetic energy relaxation, but initial and final target temperature values are ignored.

Essentially, a Hugoniosat simulation is an NPT simulation in which the user-specified target temperature is replaced with a time-dependent target temperature  $T_t$  obtained from the following equation:

$$T_t - T = \frac{\left(\frac{1}{2}(P + P_0)(V_0 - V) + E_0 - E\right)}{N_{dof}k_B} = \Delta$$

where  $T$  and  $T_t$  are the instantaneous and target temperatures,  $P$  and  $P_0$  are the instantaneous and reference pressures or axial stresses, depending on whether hydrostatic or uniaxial compression is being performed,  $V$  and  $V_0$  are the instantaneous and reference volumes,  $E$  and  $E_0$  are the instantaneous and reference internal energy (potential plus kinetic),  $N_{dof}$  is the number of degrees of freedom used in the definition of temperature, and  $k_B$  is the Boltzmann constant.  $\Delta$  is the negative deviation of the instantaneous temperature from the target temperature. When the system reaches a stable equilibrium, the value of  $\Delta$  should fluctuate about zero.

The values of  $E_0$ ,  $V_0$ , and  $P_0$  are the instantaneous values at the start of the simulation. These can be overridden using the `fix_modify` keywords `e0`, `v0`, and `p0` described below.

---

**Note:** Unlike the `fix temp/berendsen` command which performs thermostating but NO time integration, this fix performs thermostating/barostatting AND time integration. Thus you should not use any other time integration fix, such as `fix nve` on atoms to which this fix is applied. Likewise, this fix should not be used on atoms that have their temperature controlled by another fix - e.g. by `fix langevin` or `fix temp/rescale` commands.

---

This fix computes a temperature and pressure at each timestep. To do this, the fix creates its own computes of style “temp” and “pressure”, as if one of these two sets of commands had been issued:

```
compute fix-ID_temp group-ID temp
compute fix-ID_press group-ID pressure fix-ID_temp

compute fix-ID_temp all temp
compute fix-ID_press all pressure fix-ID_temp
```

See the `compute temp` and `compute pressure` commands for details. Note that the IDs of the new computes are the `fix-ID + underscore + “temp”` or `fix-ID + underscore + “press”`. The group for the new computes is “all” since pressure is computed for the entire system.

Note that these are NOT the computes used by thermodynamic output (see the `thermo_style` command) with ID = `thermo_temp` and `thermo_press`. This means you can change the attributes of this fix’s temperature or pressure via the `compute_modify` command or print this temperature or pressure during thermodynamic output via the `thermo_style custom` command using the appropriate compute-ID. It also means that changing attributes of `thermo_temp` or `thermo_press` will have no effect on this fix.

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix command-line switch` when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 17.110.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the values of  $E_0$ ,  $V_0$ , and  $P_0$ , as well as the state of all the thermostat and barostat variables to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify* *e0*, *v0* and *p0* keywords can be used to define the values of  $E_0$ ,  $V_0$ , and  $P_0$ . Note the the values for *e0* and *v0* are extensive, and so must correspond to the total energy and volume of the entire system, not energy and volume per atom. If any of these quantities are not specified, then the instantaneous value in the system at the start of the simulation is used.

The *fix\_modify* *temp* and *press* options are supported by this fix. You can use them to assign a *compute* you have defined to this fix which will be used in its thermostating or barostating procedure, as described above. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details. Note that this energy is *\*not\** included in the definition of internal energy  $E$  when calculating the value of  $\Delta$  in the above equation.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

This fix also computes a global vector of quantities, which can be accessed by various *output commands*. The scalar The vector values are “intensive”.

The vector stores three quantities unique to this fix ( $\Delta$ ,  $U_s$ , and  $u_p$ ), followed by all the internal Nose/Hoover thermostat and barostat variables defined for *fix npt*.  $\Delta$  is the deviation of the temperature from the target temperature, given by the above equation.  $U_s$  and  $u_p$  are the shock and particle velocity corresponding to a steady shock calculated from the RH conditions. They have units of distance/time.

#### 17.110.5 Restrictions

This fix style is part of the SHOCK package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

All the usual restrictions for *fix npt* apply, plus the additional ones mentioned above.

#### 17.110.6 Related commands

*fix msst*, *fix npt*, *fix\_modify*

#### 17.110.7 Default

The keyword defaults are the same as those for *fix npt*

---

(Ravelo) Ravelo, Holian, Germann and Lomdahl, Phys Rev B, 70, 014103 (2004).

## 17.111 fix npt/asphere command

Accelerator Variants: *npt/asphere/omp*

### 17.111.1 Syntax

```
fix ID group-ID npt/asphere keyword value ...
```

- ID, group-ID are documented in *fix* command
- npt/asphere = style name of this fix command
- additional thermostat and barostat related keyword/value pairs from the *fix npt* command can be appended

### 17.111.2 Examples

```
fix 1 all npt/asphere temp 300.0 300.0 100.0 iso 0.0 0.0 1000.0
fix 2 all npt/asphere temp 300.0 300.0 100.0 x 5.0 5.0 1000.0
fix 2 all npt/asphere temp 300.0 300.0 100.0 x 5.0 5.0 1000.0 drag 0.2
fix 2 water npt/asphere temp 300.0 300.0 100.0 aniso 0.0 0.0 1000.0 dilate partial
```

### 17.111.3 Description

Perform constant NPT integration to update position, velocity, orientation, and angular velocity each timestep for aspherical or ellipsoidal particles in the group using a Nose/Hoover temperature thermostat and Nose/Hoover pressure barostat. P is pressure; T is temperature. This creates a system trajectory consistent with the isothermal-isobaric ensemble.

This fix differs from the *fix npt* command, which assumes point particles and only updates their position and velocity.

The thermostat is applied to both the translational and rotational degrees of freedom for the aspherical particles, assuming a compute is used which calculates a temperature that includes the rotational degrees of freedom (see below). The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

Additional parameters affecting the thermostat and barostat are specified by keywords and values documented with the *fix npt* command. See, for example, discussion of the *temp*, *iso*, *aniso*, and *dilate* keywords.

The particles in the fix group are the only ones whose velocities and positions are updated by the velocity/position update portion of the NPT integration.

Regardless of what particles are in the fix group, a global pressure is computed for all particles. Similarly, when the size of the simulation box is changed, all particles are re-scaled to new positions, unless the keyword *dilate* is specified with a value of *partial*, in which case only the particles in the fix group are re-scaled. The latter can be useful for leaving the coordinates of particles in a solid substrate unchanged and controlling the pressure of a surrounding fluid.

This fix computes a temperature and pressure each timestep. To do this, the fix creates its own computes of style “temp/asphere” and “pressure”, as if these commands had been issued:

```
compute fix-ID_temp all temp/asphere
compute fix-ID_press all pressure fix-ID_temp
```

See the *compute temp/asphere* and *compute pressure* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix\_ID + underscore + “press”, and the group for the new computes is “all” since pressure is computed for the entire system.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of this fix’s temperature or pressure via the *compute\_modify* command or print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 17.111.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover thermostat and barostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* and *press* options are supported by this fix. You can use them to assign a *compute* you have defined to this fix which will be used in its thermostating or barostating procedure. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix npt* command.

This fix can ramp its target temperature and pressure over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.



### 17.111.5 Restrictions

This fix is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular momentum and a quaternion as defined by the *atom\_style ellipsoid* command.

All particles in the group must be finite-size. They cannot be point particles, but they can be aspherical or spherical as defined by their shape attribute.

### 17.111.6 Related commands

*fix npt*, *fix nve\_asphere*, *fix nvt\_asphere*, *fix\_modify*

### 17.111.7 Default

none

## 17.112 fix npt/body command

### 17.112.1 Syntax

```
fix ID group-ID npt/body keyword value ...
```

- ID, group-ID are documented in *fix* command
- npt/body = style name of this fix command
- additional thermostat and barostat related keyword/value pairs from the *fix npt* command can be appended

### 17.112.2 Examples

```
fix 1 all npt/body temp 300.0 300.0 100.0 iso 0.0 0.0 1000.0
fix 2 all npt/body temp 300.0 300.0 100.0 x 5.0 5.0 1000.0
fix 2 all npt/body temp 300.0 300.0 100.0 x 5.0 5.0 1000.0 drag 0.2
fix 2 water npt/body temp 300.0 300.0 100.0 aniso 0.0 0.0 1000.0 dilate partial
```

### 17.112.3 Description

Perform constant NPT integration to update position, velocity, orientation, and angular velocity each timestep for body particles in the group using a Nose/Hoover temperature thermostat and Nose/Hoover pressure barostat. P is pressure; T is temperature. This creates a system trajectory consistent with the isothermal-isobaric ensemble.

This fix differs from the *fix npt* command, which assumes point particles and only updates their position and velocity.

The thermostat is applied to both the translational and rotational degrees of freedom for the body particles, assuming a compute is used which calculates a temperature that includes the rotational degrees of freedom (see below). The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

Additional parameters affecting the thermostat and barostat are specified by keywords and values documented with the *fix npt* command. See, for example, discussion of the *temp*, *iso*, *aniso*, and *dilate* keywords.

The particles in the fix group are the only ones whose velocities and positions are updated by the velocity/position update portion of the NPT integration.

Regardless of what particles are in the fix group, a global pressure is computed for all particles. Similarly, when the size of the simulation box is changed, all particles are re-scaled to new positions, unless the keyword *dilate* is specified with a value of *partial*, in which case only the particles in the fix group are re-scaled. The latter can be useful for leaving the coordinates of particles in a solid substrate unchanged and controlling the pressure of a surrounding fluid.

---

This fix computes a temperature and pressure each timestep. To do this, the fix creates its own computes of style “temp/body” and “pressure”, as if these commands had been issued:

```
compute fix-ID_temp all temp/body
compute fix-ID_press all pressure fix-ID_temp
```

See the *compute temp/body* and *compute pressure* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix\_ID + underscore + “press”, and the group for the new computes is “all” since pressure is computed for the entire system.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of this fix’s temperature or pressure via the *compute\_modify* command or print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this fix.

Like other fixes that perform thermostatting, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostatting is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

### 17.112.4 Restart, `fix_modify`, `output`, `run start/stop`, `minimize` info

This fix writes the state of the Nose/Hoover thermostat and barostat to *binary restart files*. See the `read_restart` command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The `fix_modify temp` and `press` options are supported by this fix. You can use them to assign a `compute` you have defined to this fix which will be used in its thermostating or barostating procedure. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords `ecouple` and `econserve`. See the `thermo_style` doc page for details.

This fix computes the same global scalar and global vector of quantities as does the `fix npt` command.

This fix can ramp its target temperature and pressure over multiple runs, using the `start` and `stop` keywords of the `run` command. See the `run` command for details of how to do this.

This fix is not invoked during *energy minimization*.

### 17.112.5 Restrictions

This fix is part of the BODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular momentum and a quaternion as defined by the `atom_style body` command.

### 17.112.6 Related commands

`fix npt`, `fix nve_body`, `fix nvt_body`, `fix_modify`

### 17.112.7 Default

none

## 17.113 `fix npt/cauchy` command

### 17.113.1 Syntax

```
fix ID group-ID style_name keyword value ...
```

- ID, group-ID are documented in `fix` command
- style\_name = `npt/cauchy`
- one or more keyword/value pairs may be appended
- keyword = `temp` or `iso` or `aniso` or `tri` or `x` or `y` or `z` or `xy` or `yz` or `xz` or `couple` or `tchain` or `pchain` or `mtk` or `tloop` or `ploop` or `nreset` or `drag` or `dilate` or `scalexy` or `scaleyz` or `scalexz` or `flip` or `fixedpoint`

```

temp values = Tstart Tstop Tdamp
  Tstart,Tstop = external temperature at start/end of run
  Tdamp = temperature damping parameter (time units)
iso or aniso or tri values = Pstart Pstop Pdamp
  Pstart,Pstop = scalar external pressure at start/end of run (pressure units)
  Pdamp = pressure damping parameter (time units)
x or y or z or xy or yz or xz values = Pstart Pstop Pdamp
  Pstart,Pstop = external stress tensor component at start/end of run (pressure
↪units)
  Pdamp = stress damping parameter (time units)
couple = none or xyz or xy or yz or xz
tchain value = N
  N = length of thermostat chain (1 = single thermostat)
pchain values = N
  N length of thermostat chain on barostat (0 = no thermostat)
mtk value = yes or no = add in MTK adjustment term or not
tloop value = M
  M = number of sub-cycles to perform on thermostat
ploop value = M
  M = number of sub-cycles to perform on barostat thermostat
nreset value = reset reference cell every this many timesteps
drag value = Df
  Df = drag factor added to barostat/thermostat (0.0 = no drag)
dilate value = dilate-group-ID
  dilate-group-ID = only dilate atoms in this group due to barostat volume changes
scalexy value = yes or no = scale xy with ly
scaleyz value = yes or no = scale yz with lz
scalexz value = yes or no = scale xz with lz
flip value = yes or no = allow or disallow box flips when it becomes highly skewed
alpha value = strength of Cauchy stress control parameter
continue value = yes or no = whether of not to continue from a previous run
fixedpoint values = x y z
  x,y,z = perform barostat dilation/contraction around this point (distance units)

```

### 17.113.2 Examples

```
fix 1 water npt/cauchy temp 300.0 300.0 100.0 iso 0.0 0.0 1000.0 alpha 0.001
```

### 17.113.3 Description

This command performs time integration on Nose-Hoover style non-Hamiltonian equations of motion which are designed to generate positions and velocities sampled from the isothermal-isobaric (npt) ensembles. This updates the position and velocity for atoms in the group each timestep and the box dimensions.

The thermostatting and barostatting is achieved by adding some dynamic variables which are coupled to the particle velocities (thermostatting) and simulation domain dimensions (barostatting). In addition to basic thermostatting and barostatting, this fix can also create a chain of thermostats coupled to the particle thermostat, and another chain of thermostats coupled to the barostat variables. The barostat can be coupled to the overall box volume, or to individual dimensions, including the *xy*, *xz* and *yz* tilt dimensions. The external pressure of the barostat can be specified as either a scalar pressure (isobaric ensemble) or as components of a symmetric stress tensor (constant stress ensemble). When used correctly, the time-averaged temperature and stress tensor of the particles will match the target values specified by *Tstart/Tstop* and *Pstart/Pstop*.

The equations of motion used are those of Shinoda et al in (*Shinoda*), which combine the hydrostatic equations of Martyna, Tobias and Klein in (*Martyna*) with the strain energy proposed by Parrinello and Rahman in (*Parrinello*). The time integration schemes closely follow the time-reversible measure-preserving Verlet and rRESPA integrators derived by Tuckerman et al in (*Tuckerman*).

---

The thermostat parameters are specified using the *temp* keyword. Other thermostat-related keywords are *tchain*, *tloop* and *drag*, which are discussed below.

The thermostat is applied to only the translational degrees of freedom for the particles. The translational degrees of freedom can also have a bias velocity removed before thermostating takes place; see the description below. The desired temperature at each timestep is a ramped value during the run from *Tstart* to *Tstop*. The *Tdamp* parameter is specified in time units and determines how rapidly the temperature is relaxed. For example, a value of 10.0 means to relax the temperature in a timespan of (roughly) 10 time units (e.g.  $\tau$  or fs or ps - see the *units* command). The atoms in the *fix* group are the only ones whose velocities and positions are updated by the velocity/position update portion of the integration.

---

**Note:** A Nose-Hoover thermostat will not work well for arbitrary values of *Tdamp*. If *Tdamp* is too small, the temperature can fluctuate wildly; if it is too large, the temperature will take a very long time to equilibrate. A good choice for many models is a *Tdamp* of around 100 timesteps. Note that this is NOT the same as 100 time units for most *units* settings.

---

The barostat parameters are specified using one or more of the *iso*, *aniso*, *tri*, *x*, *y*, *z*, *xy*, *xz*, *yz*, and *couple* keywords. These keywords give you the ability to specify all 6 components of an external stress tensor, and to couple various of these components together so that the dimensions they represent are varied together during a constant-pressure simulation.

Other barostat-related keywords are *pchain*, *mtk*, *ploop*, *nreset*, *drag*, and *dilate*, which are discussed below.

Orthogonal simulation boxes have 3 adjustable dimensions (x,y,z). Triclinic (non-orthogonal) simulation boxes have 6 adjustable dimensions (x,y,z,xy,xz,yz). The *create\_box*, *read\_data*, and *read\_restart* commands specify whether the simulation box is orthogonal or non-orthogonal (triclinic) and explain the meaning of the xy,xz,yz tilt factors.

The target pressures for each of the 6 components of the stress tensor can be specified independently via the *x*, *y*, *z*, *xy*, *xz*, *yz* keywords, which correspond to the 6 simulation box dimensions. For each component, the external pressure or tensor component at each timestep is a ramped value during the run from *Pstart* to *Pstop*. If a target pressure is specified for a component, then the corresponding box dimension will change during a simulation. For example, if the *y* keyword is used, the y-box length will change. If the *xy* keyword is used, the xy tilt factor will change. A box dimension will not change if that component is not specified, although you have the option to change that dimension via the *fix deform* command.

Note that in order to use the *xy*, *xz*, or *yz* keywords, the simulation box must be triclinic, even if its initial tilt factors are 0.0.

For all barostat keywords, the *Pdamp* parameter operates like the *Tdamp* parameter, determining the time scale on which pressure is relaxed. For example, a value of 10.0 means to relax the pressure in a timespan of (roughly) 10 time units (e.g.  $\tau$  or fs or ps - see the *units* command).

---

**Note:** A Nose-Hoover barostat will not work well for arbitrary values of *Pdamp*. If *Pdamp* is too small, the pressure and volume can fluctuate wildly; if it is too large, the pressure will take a very long time to equilibrate. A good choice for many models is a *Pdamp* of around 1000 timesteps. However, note that *Pdamp* is specified in time units, and that timesteps are NOT the same as time units for most *units* settings.

---

Regardless of what atoms are in the `fix` group (the only atoms which are time integrated), a global pressure or stress tensor is computed for all atoms. Similarly, when the size of the simulation box is changed, all atoms are re-scaled to new positions, unless the keyword `dilate` is specified with a `dilate-group-ID` for a group that represents a subset of the atoms. This can be useful, for example, to leave the coordinates of atoms in a solid substrate unchanged and controlling the pressure of a surrounding fluid. This option should be used with care, since it can be unphysical to dilate some atoms and not others, because it can introduce large, instantaneous displacements between a pair of atoms (one dilated, one not) that are far from the dilation origin. Also note that for atoms not in the `fix` group, a separate time integration `fix` like `fix nve` or `fix nvt` can be used on them, independent of whether they are dilated or not.

---

The `couple` keyword allows two or three of the diagonal components of the pressure tensor to be “coupled” together. The value specified with the keyword determines which are coupled. For example, `xz` means the  $P_{xx}$  and  $P_{zz}$  components of the stress tensor are coupled. `xyz` means all 3 diagonal components are coupled. Coupling means two things: the instantaneous stress will be computed as an average of the corresponding diagonal components, and the coupled box dimensions will be changed together in lockstep, meaning coupled dimensions will be dilated or contracted by the same percentage every timestep. The `Pstart`, `Pstop`, `Pdamp` parameters for any coupled dimensions must be identical. `Couple xyz` can be used for a 2d simulation; the  $z$  dimension is simply ignored.

---

The `iso`, `aniso`, and `tri` keywords are simply shortcuts that are equivalent to specifying several other keywords together.

The keyword `iso` means couple all 3 diagonal components together when pressure is computed (hydrostatic pressure), and dilate/contract the dimensions together. Using “`iso Pstart Pstop Pdamp`” is the same as specifying these 4 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
couple xyz
```

The keyword `aniso` means  $x$ ,  $y$ , and  $z$  dimensions are controlled independently using the  $P_{xx}$ ,  $P_{yy}$ , and  $P_{zz}$  components of the stress tensor as the driving forces, and the specified scalar external pressure. Using “`aniso Pstart Pstop Pdamp`” is the same as specifying these 4 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
couple none
```

The keyword `tri` means  $x$ ,  $y$ ,  $z$ ,  $xy$ ,  $xz$ , and  $yz$  dimensions are controlled independently using their individual stress components as the driving forces, and the specified scalar pressure as the external normal stress. Using “`tri Pstart Pstop Pdamp`” is the same as specifying these 7 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
xy 0.0 0.0 Pdamp
yz 0.0 0.0 Pdamp
xz 0.0 0.0 Pdamp
couple none
```

---

In some cases (e.g. for solids) the pressure (volume) and/or temperature of the system can oscillate undesirably when a Nose/Hoover barostat and thermostat is applied. The optional `drag` keyword will damp these oscillations, although it alters the Nose/Hoover equations. A value of 0.0 (no drag) leaves the Nose/Hoover formalism unchanged. A non-zero value adds a drag term; the larger the value specified, the greater the damping effect. Performing a short run and

monitoring the pressure and temperature is the best way to determine if the drag term is working. Typically a value between 0.2 to 2.0 is sufficient to damp oscillations after a few periods. Note that use of the drag keyword will interfere with energy conservation and will also change the distribution of positions and velocities so that they do not correspond to the nominal NVT, NPT, or NPH ensembles.

An alternative way to control initial oscillations is to use chain thermostats. The keyword *tchain* determines the number of thermostats in the particle thermostat. A value of 1 corresponds to the original Nose-Hoover thermostat. The keyword *pchain* specifies the number of thermostats in the chain thermostating the barostat degrees of freedom. A value of 0 corresponds to no thermostating of the barostat variables.

The *mtk* keyword controls whether or not the correction terms due to Martyna, Tuckerman, and Klein are included in the equations of motion (*Martyna*). Specifying *no* reproduces the original Hoover barostat, whose volume probability distribution function differs from the true NPT and NPH ensembles by a factor of  $1/V$ . Hence using *yes* is more correct, but in many cases the difference is negligible.

The keyword *tloop* can be used to improve the accuracy of integration scheme at little extra cost. The initial and final updates of the thermostat variables are broken up into *tloop* sub-steps, each of length  $dt/tloop$ . This corresponds to using a first-order Suzuki-Yoshida scheme (*Tuckerman*). The keyword *ploop* does the same thing for the barostat thermostat.

The keyword *nreset* controls how often the reference dimensions used to define the strain energy are reset. If this keyword is not used, or is given a value of zero, then the reference dimensions are set to those of the initial simulation domain and are never changed. If the simulation domain changes significantly during the simulation, then the final average pressure tensor will differ significantly from the specified values of the external stress tensor. A value of *nstep* means that every *nstep* timesteps, the reference dimensions are set to those of the current simulation domain.

The *scaleyz*, *scalexz*, and *scalexy* keywords control whether or not the corresponding tilt factors are scaled with the associated box dimensions when barostatting triclinic periodic cells. The default values *yes* will turn on scaling, which corresponds to adjusting the linear dimensions of the cell while preserving its shape. Choosing *no* ensures that the tilt factors are not scaled with the box dimensions. See below for restrictions and default values in different situations. In older versions of LAMMPS, scaling of tilt factors was not performed. The old behavior can be recovered by setting all three scale keywords to *no*.

The *flip* keyword allows the tilt factors for a triclinic box to exceed half the distance of the parallel box length, as discussed below. If the *flip* value is set to *yes*, the bound is enforced by flipping the box when it is exceeded. If the *flip* value is set to *no*, the tilt will continue to change without flipping. Note that if applied stress induces large deformations (e.g. in a liquid), this means the box shape can tilt dramatically and LAMMPS will run less efficiently, due to the large volume of communication needed to acquire ghost atoms around a processor's irregular-shaped sub-domain. For extreme values of tilt, LAMMPS may also lose atoms and generate an error.

The *fixedpoint* keyword specifies the fixed point for barostat volume changes. By default, it is the center of the box. Whatever point is chosen will not move during the simulation. For example, if the lower periodic boundaries pass through (0,0,0), and this point is provided to *fixedpoint*, then the lower periodic boundaries will remain at (0,0,0), while the upper periodic boundaries will move twice as far. In all cases, the particle trajectories are unaffected by the chosen value, except for a time-dependent constant translation of positions.

---

**Note:** Using a barostat coupled to tilt dimensions *xy*, *xz*, *yz* can sometimes result in arbitrarily large values of the tilt dimensions, i.e. a dramatically deformed simulation box. LAMMPS allows the tilt factors to grow a small amount beyond the normal limit of half the box length (0.6 times the box length), and then performs a box “flip” to an equivalent periodic cell. See the discussion of the *flip* keyword above, to allow this bound to be exceeded, if desired.

---

The flip operation is described in more detail in the page for *fix deform*. Both the barostat dynamics and the atom trajectories are unaffected by this operation. However, if a tilt factor is incremented by a large amount (1.5 times the box length) on a single timestep, LAMMPS can not accommodate this event and will terminate the simulation with an error. This error typically indicates that there is something badly wrong with how the simulation was constructed, such as specifying values of *Pstart* that are too far from the current stress value, or specifying a timestep that is too large.

Triclinic barostatting should be used with care. This also is true for other barostat styles, although they tend to be more forgiving of insults. In particular, it is important to recognize that equilibrium liquids can not support a shear stress and that equilibrium solids can not support shear stresses that exceed the yield stress.

One exception to this rule is if the first dimension in the tilt factor (x for xy) is non-periodic. In that case, the limits on the tilt factor are not enforced, since flipping the box in that dimension does not change the atom positions due to non-periodicity. In this mode, if you tilt the system to extreme angles, the simulation will simply become inefficient due to the highly skewed simulation box.

---

**Note:** Unlike the *fix temp/berendsen* command which performs thermostating but NO time integration, this *fix* performs thermostating/barostatting AND time integration. Thus you should not use any other time integration *fix*, such as *fix nve* on atoms to which this *fix* is applied. Likewise, *fix npt/cauchy* should not normally be used on atoms that also have their temperature controlled by another *fix* - e.g. by *fix langevin* or *fix temp/rescale* commands.

---

See the *Howto thermostat* and *Howto barostat* doc pages for a discussion of different ways to compute temperature and perform thermostating and barostatting.

---

This *fix* compute a temperature and pressure each timestep. To do this, the *fix* creates its own computes of style “temp” and “pressure”, as if one of these sets of commands had been issued:

```
compute fix-ID_temp all temp
compute fix-ID_press all pressure fix-ID_temp
```

The group for both the new temperature and pressure compute is “all” since pressure is computed for the entire system. See the *compute temp* and *compute pressure* commands for details. Note that the IDs of the new computes are the *fix-ID* + underscore + “temp” or *fix-ID* + underscore + “press”.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of these *fix*’s temperature or pressure via the *compute\_modify* command. Or you can print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this *fix*.

Like other *fixes* that perform thermostating, this *fix* can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this *fix* that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

This *fix* can be used with either the *verlet* or *respa integrators*. When using this *fix* with *respa*, LAMMPS uses an integrator constructed according to the following factorization of the Liouville propagator (for two rRESPA levels):

$$\begin{aligned} \exp(iL\Delta t) = & \hat{E} \exp\left(iL_{T\text{-baro}} \frac{\Delta t}{2}\right) \exp\left(iL_{T\text{-part}} \frac{\Delta t}{2}\right) \exp\left(iL_{\epsilon,2} \frac{\Delta t}{2}\right) \exp\left(iL_2^{(2)} \frac{\Delta t}{2}\right) \\ & \times \left[ \exp\left(iL_2^{(1)} \frac{\Delta t}{2n}\right) \exp\left(iL_{\epsilon,1} \frac{\Delta t}{2n}\right) \exp\left(iL_1 \frac{\Delta t}{n}\right) \exp\left(iL_{\epsilon,1} \frac{\Delta t}{2n}\right) \exp\left(iL_2^{(1)} \frac{\Delta t}{2n}\right) \right]^n \\ & \times \exp\left(iL_2^{(2)} \frac{\Delta t}{2}\right) \exp\left(iL_{\epsilon,2} \frac{\Delta t}{2}\right) \exp\left(iL_{T\text{-part}} \frac{\Delta t}{2}\right) \exp\left(iL_{T\text{-baro}} \frac{\Delta t}{2}\right) \\ & + \mathcal{O}(\Delta t^3) \end{aligned}$$



This factorization differs somewhat from that of Tuckerman et al, in that the barostat is only updated at the outermost rRESPA level, whereas Tuckerman's factorization requires splitting the pressure into pieces corresponding to the forces computed at each rRESPA level. In theory, the latter method will exhibit better numerical stability. In practice, because Pdamp is normally chosen to be a large multiple of the outermost rRESPA timestep, the barostat dynamics are not the limiting factor for numerical stability. Both factorizations are time-reversible and can be shown to preserve the phase space measure of the underlying non-Hamiltonian equations of motion.

---

**Note:** Under NPT dynamics, for a system with zero initial total linear momentum, the total momentum fluctuates close to zero. It may occasionally undergo brief excursions to non-negligible values, before returning close to zero. Over long simulations, this has the effect of causing the center-of-mass to undergo a slow random walk. This can be mitigated by resetting the momentum at infrequent intervals using the *fix momentum* command.

---

#### 17.113.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of all the thermostat and barostat variables to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* and *press* options are supported by this fix. You can use them to assign a *compute* you have defined to this fix which will be used in its thermostating or barostating procedure, as described above. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

---

**Note:** If both the *temp* and *press* keywords are used in a single thermo\_modify command (or in two separate commands), then the order in which the keywords are specified is important. Note that a *pressure compute* defines its own temperature compute as an argument when it is specified. The *temp* keyword will override this (for the pressure compute being used by fix npt), but only if the *temp* keyword comes after the *press* keyword. If the *temp* keyword comes before the *press* keyword, then the new pressure compute specified by the *press* keyword will be unaffected by the *temp* setting.

---

The cumulative energy change in the system imposed by this fix, due to thermostating and/or barostating, is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

This fix also computes a global vector of quantities, which can be accessed by various *output commands*. The vector values are “intensive”.

The vector stores internal Nose/Hoover thermostat and barostat variables. The number and meaning of the vector values depends on which fix is used and the settings for keywords *tchain* and *pchain*, which specify the number of Nose/Hoover chains for the thermostat and barostat. If no thermostating is done, then *tchain* is 0. If no barostating is done, then *pchain* is 0. In the following list, “ndof” is 0, 1, 3, or 6, and is the number of degrees of freedom in the barostat. Its value is 0 if no barostat is used, else its value is 6 if any off-diagonal stress tensor component is barostatted, else its value is 1 if *couple xyz* is used or *couple xy* for a 2d simulation, otherwise its value is 3.

The order of values in the global vector and their meaning is as follows. The notation means there are tchain values for eta, followed by tchain for eta\_dot, followed by ndof for omega, etc:

- eta[tchain] = particle thermostat displacements (unitless)
- eta\_dot[tchain] = particle thermostat velocities (1/time units)

- `omega[ndof]` = barostat displacements (unitless)
- `omega_dot[ndof]` = barostat velocities (1/time units)
- `etap[pchain]` = barostat thermostat displacements (unitless)
- `etap_dot[pchain]` = barostat thermostat velocities (1/time units)
- `PE_eta[tchain]` = potential energy of each particle thermostat displacement (energy units)
- `KE_eta_dot[tchain]` = kinetic energy of each particle thermostat velocity (energy units)
- `PE_omega[ndof]` = potential energy of each barostat displacement (energy units)
- `KE_omega_dot[ndof]` = kinetic energy of each barostat velocity (energy units)
- `PE_etap[pchain]` = potential energy of each barostat thermostat displacement (energy units)
- `KE_etap_dot[pchain]` = kinetic energy of each barostat thermostat velocity (energy units)
- `PE_strain[1]` = scalar strain energy (energy units)

This fix can ramp its external temperature and pressure over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

---

### 17.113.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

*X*, *y*, *z* cannot be barostatted if the associated dimension is not periodic. *Xy*, *xz*, and *yz* can only be barostatted if the simulation domain is triclinic and the second dimension in the keyword (*y* dimension in *xy*) is periodic. *Z*, *xz*, and *yz*, cannot be barostatted for 2D simulations. The *create\_box*, *read\_data*, and *read\_restart* commands specify whether the simulation box is orthogonal or non-orthogonal (triclinic) and explain the meaning of the *xy,xz,yz* tilt factors.

For the *temp* keyword, the final *Tstop* cannot be 0.0 since it would make the external *T* = 0.0 at some timestep during the simulation which is not allowed in the Nose/Hoover formulation.

The *scaleyz yes* and *scalexz yes* keyword/value pairs can not be used for 2D simulations. *scaleyz yes*, *scalexz yes*, and *scalexy yes* options can only be used if the second dimension in the keyword is periodic, and if the tilt factor is not coupled to the barostat via keywords *tri*, *yz*, *xz*, and *xy*.

The *alpha* keyword modifies the barostat as per Miller et al. (Miller)\_"#nc-Miller" so that the Cauchy stress is controlled. *alpha* is the non-dimensional parameter, typically set to 0.001 or 0.01 that determines how aggressively the algorithm drives the system towards the set Cauchy stresses. Larger values of *alpha* will modify the system more quickly, but can lead to instabilities. Smaller values will lead to longer convergence time. Since *alpha* also influences how much the stress fluctuations deviate from the equilibrium fluctuations, it should be set as small as possible.

A *continue* value of *yes* indicates that the fix is subsequent to a previous run with the *npt/cauchy* fix, and the intention is to continue from the converged stress state at the end of the previous run. This may be required, for example, when implementing a multi-step loading/unloading sequence over several fixes.

Setting *alpha* to zero is not permitted. To “turn off” the cauchystat control and thus restore the equilibrium stress fluctuations, two subsequent fixes should be used. In the first, fix *npt/cauchy* is used and the simulation box equilibrates to the correct shape for the desired stresses. In the second, fix *npt* is used instead which uses the original Parrinello-Rahman algorithm, but now with the corrected simulation box shape from using fix *npt/cauchy*.

This fix can be used with dynamic groups as defined by the *group* command. Likewise it can be used with groups to which atoms are added or deleted over time, e.g. a deposition simulation. However, the conservation properties of the

thermostat and barostat are defined for systems with a static set of atoms. You may observe odd behavior if the atoms in a group vary dramatically over time or the atom count becomes very small.

### 17.113.6 Related commands

*fix nve, fix\_modify, run\_style*

### 17.113.7 Default

The keyword defaults are tchain = 3, pchain = 3, mtk = yes, tloop = ploop = 1, nreset = 0, drag = 0.0, dilate = all, couple = none, cauchy = no, scaleyz = scalexz = scalexy = yes if periodic in second dimension and not coupled to barostat, otherwise no.

**(Martyna)** Martyna, Tobias and Klein, J Chem Phys, 101, 4177 (1994).

**(Parrinello)** Parrinello and Rahman, J Appl Phys, 52, 7182 (1981).

**(Tuckerman)** Tuckerman, Alejandre, Lopez-Rendon, Jochim, and Martyna, J Phys A: Math Gen, 39, 5629 (2006).

**(Shinoda)** Shinoda, Shiga, and Mikami, Phys Rev B, 69, 134103 (2004).

**(Miller)** Miller, Tadmor, Gibson, Bernstein and Pavia, J Chem Phys, 144, 184107 (2016).

## 17.114 fix npt/sphere command

Accelerator Variants: *npt/sphere/omp*

### 17.114.1 Syntax

```
fix ID group-ID npt/sphere keyword value ...
```

- ID, group-ID are documented in *fix* command npt/sphere = style name of this fix command zero or more keyword/value pairs may be appended
- keyword = *disc*

*disc* value = none = treat particles as 2d discs, not spheres

- NOTE: additional thermostat and barostat and dipole related keyword/value pairs from the *fix npt* command can be appended

### 17.114.2 Examples

```
fix 1 all npt/sphere temp 300.0 300.0 100.0 iso 0.0 0.0 1000.0
fix 2 all npt/sphere temp 300.0 300.0 100.0 x 5.0 5.0 1000.0
fix 2 all npt/sphere temp 300.0 300.0 100.0 x 5.0 5.0 1000.0 disc
fix 2 all npt/sphere temp 300.0 300.0 100.0 x 5.0 5.0 1000.0 drag 0.2
fix 2 all npt/sphere temp 300.0 300.0 100.0 x 5.0 5.0 1000.0 update dipole
fix 2 water npt/sphere temp 300.0 300.0 100.0 aniso 0.0 0.0 1000.0 dilate partial
```

### 17.114.3 Description

Perform constant NPT integration to update position, velocity, and angular velocity each timestep for finite-size spherical particles in the group using a Nose/Hoover temperature thermostat and Nose/Hoover pressure barostat. P is pressure; T is temperature. This creates a system trajectory consistent with the isothermal-isobaric ensemble.

This fix differs from the *fix npt* command, which assumes point particles and only updates their position and velocity.

The thermostat is applied to both the translational and rotational degrees of freedom for the spherical particles, assuming a compute is used which calculates a temperature that includes the rotational degrees of freedom (see below). The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

If the *disc* keyword is used, then each particle is treated as a 2d disc (circle) instead of as a sphere. This is only possible for 2d simulations, as defined by the *dimension* keyword. The only difference between discs and spheres in this context is their moment of inertia, as used in the time integration.

Additional parameters affecting the thermostat and barostat are specified by keywords and values documented with the *fix npt* command. See, for example, discussion of the *temp*, *iso*, *aniso*, and *dilate* keywords.

The particles in the fix group are the only ones whose velocities and positions are updated by the velocity/position update portion of the NPT integration.

Regardless of what particles are in the fix group, a global pressure is computed for all particles. Similarly, when the size of the simulation box is changed, all particles are re-scaled to new positions, unless the keyword *dilate* is specified with a value of *partial*, in which case only the particles in the fix group are re-scaled. The latter can be useful for leaving the coordinates of particles in a solid substrate unchanged and controlling the pressure of a surrounding fluid.

---

This fix computes a temperature and pressure each timestep. To do this, the fix creates its own computes of style “temp/sphere” and “pressure”, as if these commands had been issued:

```
compute fix-ID_temp all temp/sphere
compute fix-ID_press all pressure fix-ID_temp
```

See the *compute temp/sphere* and *compute pressure* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix\_ID + underscore + “press”, and the group for the new computes is “all” since pressure is computed for the entire system.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of this fix’s temperature or pressure via the *compute\_modify* command or print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 17.114.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover thermostat and barostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* and *press* options are supported by this fix. You can use them to assign a *compute* you have defined to this fix which will be used in its thermostating or barostating procedure. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix npt* command.

This fix can ramp its target temperature and pressure over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

#### 17.114.5 Restrictions

This fix requires that atoms store torque and angular velocity (omega) and a radius as defined by the *atom\_style sphere* command.

All particles in the group must be finite-size spheres. They cannot be point particles.

Use of the *disc* keyword is only allowed for 2d simulations, as defined by the *dimension* keyword.

#### 17.114.6 Related commands

*fix npt*, *fix nve\_sphere*,  
*fix nvt\_sphere*, *fix npt\_asphere*, *fix\_modify*

#### 17.114.7 Default

none

## 17.115 fix numdiff command

### 17.115.1 Syntax

```
fix ID group-ID numdiff Nevery delta
```

- ID, group-ID are documented in *fix* command
- numdiff = style name of this fix command
- Nevery = calculate force by finite difference every this many timesteps
- delta = size of atom displacements (distance units)

### 17.115.2 Examples

```
fix 1 all numdiff 10 1e-6  
fix 1 movegroup numdiff 100 0.01
```

### 17.115.3 Description

Calculate forces through finite difference calculations of energy versus position. These forces can be compared to analytic forces computed by pair styles, bond styles, etc. This can be useful for debugging or other purposes.

The group specified with the command means only atoms within the group have their averages computed. Results are set to 0.0 for atoms not in the group.

This fix performs a loop over all atoms in the group. For each atom and each component of force it adds *delta* to the position, and computes the new energy of the entire system. It then subtracts *delta* from the original position and again computes the new energy of the system. It then restores the original position. That component of force is calculated as the difference in energy divided by two times *delta*.

---

**Note:** It is important to choose a suitable value for delta, the magnitude of atom displacements that are used to generate finite difference approximations to the exact forces. For typical systems, a value in the range of 1 part in 1e4 to 1e5 of the typical separation distance between atoms in the liquid or solid state will be sufficient. However, the best value will depend on a multitude of factors including the stiffness of the interatomic potential, the thermodynamic state of the material being probed, and so on. The only way to be sure that you have made a good choice is to do a sensitivity study on a representative atomic configuration, sweeping over a wide range of values of delta. If delta is too small, the output forces will vary erratically due to truncation effects. If delta is increased beyond a certain point, the output forces will start to vary smoothly with delta, due to growing contributions from higher order derivatives. In between these two limits, the numerical force values should be largely independent of delta.

---

**Note:** The cost of each energy evaluation is essentially the cost of an MD timestep. Thus invoking this fix once for a 3d system has a cost of 6N timesteps, where N is the total number of atoms in the system. So this fix can be very expensive to use for large systems. One expedient alternative is to define the fix for a group containing only a few atoms.

---

The *Nevery* argument specifies on what timesteps the force will be used calculated by finite difference.

The *delta* argument specifies the size of the displacement each atom will undergo.

---

#### 17.115.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix produces a per-atom array which can be accessed by various *output commands*, which stores the components of the force on each atom as calculated by finite difference. The per-atom values can only be accessed on timesteps that are multiples of *Nevery* since that is when the finite difference forces are calculated. See the examples in *examples/numdiff* directory to see how this fix can be used to directly compare with the analytic forces computed by LAMMPS.

The array values calculated by this compute will be in force *units*.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is invoked during *energy minimization*.

#### 17.115.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 17.115.6 Related commands

*dynamical\_matrix*, *fix numdiff/virial*,

#### 17.115.7 Default

none

### 17.116 fix numdiff/virial command

#### 17.116.1 Syntax

```
fix ID group-ID numdiff/virial Nevery delta
```

- ID, group-ID are documented in *fix* command
- numdiff/virial = style name of this fix command
- Nevery = calculate virial by finite difference every this many timesteps
- delta = magnitude of strain fields (dimensionless)

## 17.116.2 Examples

```
fix 1 all numdiff/stress 10 1e-6
```

---

## 17.116.3 Description

Calculate the virial stress tensor through a finite difference calculation of energy versus strain. These values can be compared to the analytic virial tensor computed by pair styles, bond styles, etc. This can be useful for debugging or other purposes. The specified group must be “all”.

This fix applies linear strain fields of magnitude *delta* to all the atoms relative to a point at the center of the box. The strain fields are in six different directions, corresponding to the six Cartesian components of the stress tensor defined by LAMMPS. For each direction it applies the strain field in both the positive and negative senses, and the new energy of the entire system is calculated after each. The difference in these two energies divided by two times *delta*, approximates the corresponding component of the virial stress tensor, after applying a suitable unit conversion.

---

**Note:** It is important to choose a suitable value for delta, the magnitude of strains that are used to generate finite difference approximations to the exact virial stress. For typical systems, a value in the range of 1 part in 1e5 to 1e6 will be sufficient. However, the best value will depend on a multitude of factors including the stiffness of the interatomic potential, the thermodynamic state of the material being probed, and so on. The only way to be sure that you have made a good choice is to do a sensitivity study on a representative atomic configuration, sweeping over a wide range of values of delta. If delta is too small, the output values will vary erratically due to truncation effects. If delta is increased beyond a certain point, the output values will start to vary smoothly with delta, due to growing contributions from higher order derivatives. In between these two limits, the numerical virial values should be largely independent of delta.

---

The *Nevery* argument specifies on what timesteps the force will be used calculated by finite difference.

The *delta* argument specifies the size of the displacement each atom will undergo.

---

## 17.116.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix produces a global vector which can be accessed by various *output commands*, which stores the components of the virial stress tensor as calculated by finite difference. The global vector can only be accessed on timesteps that are multiples of *Nevery* since that is when the finite difference virial is calculated. See the examples in *examples/numdiff* directory to see how this fix can be used to directly compare with the analytic virial stress tensor computed by LAMMPS.

The order of the virial stress tensor components is *xx*, *yy*, *zz*, *yz*, *xz*, and *xy*, consistent with Voigt notation. Note that the vector produced by *compute pressure* uses a different ordering, with *yz* and *xy* swapped.

The vector values calculated by this compute are “intensive”. The vector values will be in pressure *units*.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is invoked during *energy minimization*.



### 17.116.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.116.6 Related commands

*fix numdiff*, *compute pressure*

### 17.116.7 Default

none

## 17.117 fix nve command

Accelerator Variants: *nve/gpu*, *nve/intel*, *nve/kk*, *nve/omp*

### 17.117.1 Syntax

```
fix ID group-ID nve
```

- ID, group-ID are documented in *fix* command
- nve = style name of this fix command

### 17.117.2 Examples

```
fix 1 all nve
```

### 17.117.3 Description

Perform plain time integration to update position and velocity for atoms in the group each timestep. This creates a system trajectory consistent with the microcanonical ensemble (NVE) provided there are (full) periodic boundary conditions and no other “manipulations” of the system (e.g. fixes that modify forces or velocities).

This fix invokes the velocity form of the Stoermer-Verlet time integration algorithm (velocity-Verlet). Other time integration options can be invoked using the *run\_style* command.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 17.117.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.117.5 Restrictions

none

#### 17.117.6 Related commands

*fix nvt*, *fix npt*, *run\_style*

#### 17.117.7 Default

none

### 17.118 fix nve/asphere command

Accelerator Variants: *nve/asphere/gpu*, *nve/asphere/intel*

#### 17.118.1 Syntax

```
fix ID group-ID nve/asphere
```

- ID, group-ID are documented in *fix* command
- nve/asphere = style name of this fix command

#### 17.118.2 Examples

```
fix 1 all nve/asphere
```

### 17.118.3 Description

Perform constant NVE integration to update position, velocity, orientation, and angular velocity for aspherical particles in the group each timestep. V is volume; E is energy. This creates a system trajectory consistent with the microcanonical ensemble.

This fix differs from the *fix nve* command, which assumes point particles and only updates their position and velocity.

### 17.118.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 17.118.5 Restrictions

This fix is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular momentum and a quaternion as defined by the *atom\_style ellipsoid* command.

All particles in the group must be finite-size. They cannot be point particles, but they can be aspherical or spherical as defined by their shape attribute.

### 17.118.6 Related commands

*fix nve*, *fix nve/sphere*

### 17.118.7 Default

none

## 17.119 fix nve/asphere/noforce command

### 17.119.1 Syntax

```
fix ID group-ID nve/asphere/noforce
```

- ID, group-ID are documented in *fix* command
- nve/asphere/noforce = style name of this fix command

### 17.119.2 Examples

```
fix 1 all nve/asphere/noforce
```

### 17.119.3 Description

Perform updates of position and orientation, but not velocity or angular momentum for atoms in the group each timestep. In other words, the force and torque on the atoms is ignored and their velocity and angular momentum are not updated. The atom velocities and angular momenta are used to update their positions and orientation.

This is useful as an implicit time integrator for Fast Lubrication Dynamics, since the velocity and angular momentum are updated by the *pair\_style lubricateU* command.

---

### 17.119.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.119.5 Restrictions

This fix is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular momentum and a quaternion as defined by the *atom\_style ellipsoid* command.

All particles in the group must be finite-size. They cannot be point particles, but they can be aspherical or spherical as defined by their shape attribute.

## 17.119.6 Related commands

*fix nve/noforce*, *fix nve/asphere*

## 17.119.7 Default

none

# 17.120 fix nve/awpmd command

## 17.120.1 Syntax

```
fix ID group-ID nve/awpmd
```

- ID, group-ID are documented in *fix* command
- nve/awpmd = style name of this fix command

## 17.120.2 Examples

```
fix 1 all nve/awpmd
```

## 17.120.3 Description

Perform constant NVE integration to update position and velocity for nuclei and electrons in the group for the *Antisymmetrized Wave Packet Molecular Dynamics* model. V is volume; E is energy. This creates a system trajectory consistent with the microcanonical ensemble.

The operation of this fix is exactly like that described by the *fix nve* command, except that the width and width-velocity of the electron wave functions are also updated.

## 17.120.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.120.5 Restrictions

This fix is part of the AWPMD package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.120.6 Related commands

*fix nve*

### 17.120.7 Default

none

## 17.121 fix nve/body command

### 17.121.1 Syntax

```
fix ID group-ID nve/body
```

- ID, group-ID are documented in *fix* command
- nve/body = style name of this fix command

### 17.121.2 Examples

```
fix 1 all nve/body
```

### 17.121.3 Description

Perform constant NVE integration to update position, velocity, orientation, and angular velocity for body particles in the group each timestep. V is volume; E is energy. This creates a system trajectory consistent with the microcanonical ensemble. See the *Howto body* page for more details on using body particles.

This fix differs from the *fix nve* command, which assumes point particles and only updates their position and velocity.

### 17.121.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.121.5 Restrictions

This fix is part of the BODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular momentum and a quaternion as defined by the *atom\_style body* command.

All particles in the group must be body particles. They cannot be point particles.

### 17.121.6 Related commands

*fix nve, fix nve/sphere, fix nve/asphere*

### 17.121.7 Default

none

## 17.122 fix nve/bpm/sphere command

### 17.122.1 Syntax

```
fix ID group-ID nve/bpm/sphere
```

- ID, group-ID are documented in *fix* command
- nve/bpm/sphere = style name of this fix command
- zero or more keyword/value pairs may be appended
- keyword = *disc*  
*disc* value = none = treat particles as 2d discs, not spheres

### 17.122.2 Examples

```
fix 1 all nve/bpm/sphere
fix 1 all nve/bpm/sphere disc
```

### 17.122.3 Description

Perform constant NVE integration to update position, velocity, angular velocity, and quaternion orientation for finite-size spherical particles in the group each timestep. V is volume; E is energy. This creates a system trajectory consistent with the microcanonical ensemble.

This fix differs from the *fix nve* command, which assumes point particles and only updates their position and velocity. It also differs from the *fix nve/sphere* command which assumes finite-size spheroid particles which do not store a quaternion. It thus does not update a particle's orientation or quaternion.

If the *disc* keyword is used, then each particle is treated as a 2d disc (circle) instead of as a sphere. This is only possible for 2d simulations, as defined by the *dimension* keyword. The only difference between discs and spheres in this context is their moment of inertia, as used in the time integration.

#### 17.122.4 Restart, `fix_modify`, `output`, `run start/stop`, `minimize info`

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.122.5 Restrictions

This fix requires that atoms store torque, angular velocity ( $\omega$ ), a radius, and a quaternion as defined by the *atom\_style bpm/sphere* command.

All particles in the group must be finite-size spheres with quaternions. They cannot be point particles.

Use of the *disc* keyword is only allowed for 2d simulations, as defined by the *dimension* keyword.

#### 17.122.6 Related commands

*fix nve*, *fix nve/sphere*

#### 17.122.7 Default

none

### 17.123 `fix nve/dot` command

#### 17.123.1 Syntax

```
fix ID group-ID nve/dot
```

- ID, group-ID are documented in *fix* command
- nve/dot = style name of this fix command

#### 17.123.2 Examples

```
fix 1 all nve/dot
```



### 17.123.3 Description

Apply a rigid-body integrator as described in (*Davidchack*) to a group of atoms, but without Langevin dynamics. This command performs Molecular dynamics (MD) via a velocity-Verlet algorithm and an evolution operator that rotates the quaternion degrees of freedom, similar to the scheme outlined in (*Miller*).

This command is the equivalent of the *fix nve/dotc/langevin* without damping and noise and can be used to determine the stability range in a NVE ensemble prior to using the Langevin-type DOTC-integrator (see also *fix nve/dotc/langevin*). The command is equivalent to the *fix nve*. The particles are always considered to have a finite size.

An example input file can be found in `/examples/PACKAGES/cgdna/examples/duplex1/`. Further details of the implementation and stability of the integrator are contained in (*Henrich*). The preprint version of the article can be found [here](#).

### 17.123.4 Restrictions

These pair styles can only be used if LAMMPS was built with the *CG-DNA* package and the MOLECULE and ASPHERE package. See the *Build package* page for more info.

### 17.123.5 Related commands

*fix nve/dotc/langevin*, *fix nve*

### 17.123.6 Default

none

**(Davidchack)** R.L Davidchack, T.E. Ouldridge, and M.V. Tretyakov. J. Chem. Phys. 142, 144114 (2015).

**(Miller)** T. F. Miller III, M. Eleftheriou, P. Pattnaik, A. Ndirango, G. J. Martyna, J. Chem. Phys., 116, 8649-8659 (2002).

**(Henrich)** O. Henrich, Y. A. Gutierrez-Fosado, T. Curk, T. E. Ouldridge, Eur. Phys. J. E 41, 57 (2018).

## 17.124 *fix nve/dotc/langevin* command

### 17.124.1 Syntax

```
fix ID group-ID nve/dotc/langevin Tstart Tstop damp seed keyword value
```

- ID, group-ID are documented in *fix* command
- nve/dotc/langevin = style name of this fix command
- Tstart, Tstop = desired temperature at start/end of run (temperature units)
- damp = damping parameter (time units)
- seed = random number seed to use for white noise (positive integer)
- keyword = *angmom*

`angmom` value = factor  
 factor = do thermostat rotational degrees of freedom via the angular momentum and  
 ↪ apply numeric scale factor as discussed below

## 17.124.2 Examples

```
fix 1 all nve/dotc/langevin 1.0 1.0 0.03 457145 angmom 10
fix 1 all nve/dotc/langevin 0.1 0.1 78.9375 457145 angmom 10
```

## 17.124.3 Description

Apply a rigid-body Langevin-type integrator of the kind “Langevin C” as described in (*Davidchack*) to a group of atoms, which models an interaction with an implicit background solvent. This command performs Brownian dynamics (BD) via a technique that splits the integration into a deterministic Hamiltonian part and the Ornstein-Uhlenbeck process for noise and damping. The quaternion degrees of freedom are updated though an evolution operator which performs a rotation in quaternion space, preserves the quaternion norm and is akin to (*Miller*).

In terms of syntax this command has been closely modelled on the *fix langevin* and its *angmom* option. But it combines the *fix nve* and the *fix langevin* in one single command. The main feature is improved stability over the standard integrator, permitting slightly larger timestep sizes.

---

**Note:** Unlike the *fix langevin* this command performs also time integration of the translational and quaternion degrees of freedom.

---

The total force on each atom will have the form:

$$F = F_c + F_f + F_r$$

$$F_f = - \frac{m}{\text{damp}} v$$

$$F_r \propto \sqrt{\frac{k_B T m}{dt \text{ damp}}}$$

$F_c$  is the conservative force computed via the usual inter-particle interactions (*pair\_style*, *bond\_style*, etc). The  $F_f$  and  $F_r$  terms are implicitly taken into account by this *fix* on a per-particle basis.

$F_f$  is a frictional drag or viscous damping term proportional to the particle’s velocity. The proportionality constant for each atom is computed as  $\frac{m}{\text{damp}}$ , where  $m$  is the mass of the particle and *damp* is the damping factor specified by the user.

$F_r$  is a force due to solvent atoms at a temperature  $T$  randomly bumping into the particle. As derived from the fluctuation/dissipation theorem, its magnitude as shown above is proportional to  $\sqrt{\frac{k_B T m}{dt \text{ damp}}}$ , where  $k_B$  is the Boltzmann constant,  $T$  is the desired temperature,  $m$  is the mass of the particle,  $dt$  is the timestep size, and *damp* is the damping factor. Random numbers are used to randomize the direction and magnitude of this force as described in (*Dunweg*), where a uniform random number is used (instead of a Gaussian random number) for speed.

---

*Tstart* and *Tstop* have to be constant values, i.e. they cannot be variables. If used together with the oxDNA force field for coarse-grained simulation of DNA please note that  $T = 0.1$  in oxDNA units corresponds to  $T = 300$  K.

The *damp* parameter is specified in time units and determines how rapidly the temperature is relaxed. For example, a value of 0.03 means to relax the temperature in a timespan of (roughly) 0.03 time units  $\tau$  (see the *units* command). The *damp* factor can be thought of as inversely related to the viscosity of the solvent, i.e. a small relaxation time implies a

high-viscosity solvent and vice versa. See the discussion about gamma and viscosity in the documentation for the *fix viscous* command for more details. Note that the value 78.9375 in the second example above corresponds to a diffusion constant, which is about an order of magnitude larger than realistic ones. This has been used to sample configurations faster in Brownian dynamics simulations.

The random # *seed* must be a positive integer. A Marsaglia random number generator is used. Each processor uses the input seed to generate its own unique seed and its own stream of random numbers. Thus the dynamics of the system will not be identical on two runs on different numbers of processors.

The keyword/value option has to be used in the following way:

This *fix* has to be used together with the *angmom* keyword. The particles are always considered to have a finite size. The keyword *angmom* enables thermostating of the rotational degrees of freedom in addition to the usual translational degrees of freedom.

The scale factor after the *angmom* keyword gives the ratio of the rotational to the translational friction coefficient.

An example input file can be found in `examples/PACKAGES/cgdna/examples/duplex2/`. Further details of the implementation and stability of the integrators are contained in ([Henrich](#)). The preprint version of the article can be found [here](#).

---

#### 17.124.4 Restrictions

These pair styles can only be used if LAMMPS was built with the *CG-DNA* package and the MOLECULE and ASPHERE package. See the [Build package](#) page for more info.

#### 17.124.5 Related commands

*fix nve*, *fix langevin*, *fix nve/dot*, *bond\_style oxdna/fene*, *bond\_style oxdna2/fene*, *pair\_style oxdna/excv*, *pair\_style oxdna2/excv*

#### 17.124.6 Default

none

---

**(Davidchack)** R.L. Davidchack, T.E. Ouldridge, M.V. Tretyakov. *J. Chem. Phys.* 142, 144114 (2015).

**(Miller)** T. F. Miller III, M. Eleftheriou, P. Pattnaik, A. Ndirango, G. J. Martyna, *J. Chem. Phys.*, 116, 8649-8659 (2002).

**(Dunweg)** B. Dunweg, W. Paul, *Int. J. Mod. Phys. C*, 2, 817-27 (1991).

**(Henrich)** O. Henrich, Y. A. Gutierrez-Fosado, T. Curk, T. E. Ouldridge, *Eur. Phys. J. E* 41, 57 (2018).

## 17.125 fix nve/eff command

### 17.125.1 Syntax

```
fix ID group-ID nve/eff
```

- ID, group-ID are documented in *fix* command
- nve/eff = style name of this fix command

### 17.125.2 Examples

```
fix 1 all nve/eff
```

### 17.125.3 Description

Perform constant NVE integration to update position and velocity for nuclei and electrons in the group for the *electron force field* model. V is volume; E is energy. This creates a system trajectory consistent with the microcanonical ensemble.

The operation of this fix is exactly like that described by the *fix nve* command, except that the radius and radial velocity of electrons are also updated.

### 17.125.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.125.5 Restrictions

This fix is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.125.6 Related commands

*fix nve*, *fix nvt/eff*, *fix npt/eff*

### 17.125.7 Default

none

## 17.126 fix nve/limit command

### 17.126.1 Syntax

```
fix ID group-ID nve/limit xmax
```

- ID, group-ID are documented in *fix* command
- nve = style name of this fix command
- xmax = maximum distance an atom can move in one timestep (distance units)

### 17.126.2 Examples

```
fix 1 all nve/limit 0.1
```

### 17.126.3 Description

Perform constant NVE updates of position and velocity for atoms in the group each timestep. A limit is imposed on the maximum distance an atom can move in one timestep. This is useful when starting a simulation with a configuration containing highly overlapped atoms. Normally this would generate huge forces which would blow atoms out of the simulation box, causing LAMMPS to stop with an error.

Using this fix can overcome that problem. Forces on atoms must still be computable (which typically means 2 atoms must have a separation distance  $> 0.0$ ). But large velocities generated by large forces are reset to a value that corresponds to a displacement of length  $xmax$  in a single timestep.  $Xmax$  is specified in distance units; see the *units* command for details. The value of  $xmax$  should be consistent with the neighbor skin distance and the frequency of neighbor list re-building, so that pairwise interactions are not missed on successive timesteps as atoms move. See the *neighbor* and *neigh\_modify* commands for details.

Note that if a velocity reset occurs the integrator will not conserve energy. On steps where no velocity resets occur, this integrator is exactly like the *fix nve* command. Since forces are unaltered, pressures computed by thermodynamic output will still be very large for overlapped configurations.

---

**Note:** You should not use *fix shake* in conjunction with this fix. That is because fix shake applies constraint forces based on the predicted positions of atoms after the next timestep. It has no way of knowing the timestep may change due to this fix, which will cause the constraint forces to be invalid. A better strategy is to turn off fix shake when performing initial dynamics that need this fix, then turn fix shake on when doing normal dynamics with a fixed-size timestep.

---

### 17.126.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the count of how many updates of atom's velocity/position were limited by the maximum distance criterion. This should be roughly the number of atoms so affected, except that updates occur at both the beginning and end of a timestep in a velocity Verlet timestepping algorithm. This is a cumulative quantity for the current run, but is re-initialized to zero each time a run is performed. The scalar value calculated by this fix is "extensive".

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.126.5 Restrictions

none

## 17.126.6 Related commands

*fix nve*, *fix nve/force*, *pair\_style soft*

## 17.126.7 Default

none

# 17.127 fix nve/line command

## 17.127.1 Syntax

```
fix ID group-ID nve/line
```

- ID, group-ID are documented in *fix* command
- nve/line = style name of this fix command

## 17.127.2 Examples

```
fix 1 all nve/line
```

## 17.127.3 Description

Perform constant NVE integration to update position, velocity, orientation, and angular velocity for line segment particles in the group each timestep. V is volume; E is energy. This creates a system trajectory consistent with the microcanonical ensemble. See *Howto spherical* page for an overview of using line segment particles.

This fix differs from the *fix nve* command, which assumes point particles and only updates their position and velocity.

## 17.127.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.127.5 Restrictions

This fix is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that particles be line segments as defined by the *atom\_style line* command.

### 17.127.6 Related commands

*fix nve*, *fix nve/asphere*

### 17.127.7 Default

none

## 17.128 fix nve/manifold/rattle command

### 17.128.1 Syntax

```
fix ID group-ID nve/manifold/rattle tol maxit manifold manifold-args keyword value ...
```

- ID, group-ID are documented in *fix* command
- nve/manifold/rattle = style name of this fix command
- tol = tolerance to which Newton iteration must converge
- maxit = maximum number of iterations to perform
- manifold = name of the manifold
- manifold-args = parameters for the manifold
- one or more keyword/value pairs may be appended

keyword = *every*

every values = N

N = print info about iteration every N steps. N = 0 means no output

### 17.128.2 Examples

```
fix 1 all nve/manifold/rattle 1e-4 10 sphere 5.0
fix step all nve/manifold/rattle 1e-8 100 ellipsoid 2.5 2.5 5.0 every 25
```

### 17.128.3 Description

Perform constant NVE integration to update position and velocity for atoms constrained to a curved surface (manifold) in the group each timestep. The constraint is handled by RATTLE (*Andersen*) written out for the special case of single-particle constraints as explained in (*Paquay*). V is volume; E is energy. This way, the dynamics of particles constrained to curved surfaces can be studied. If combined with *fix langevin*, this generates Brownian motion of particles constrained to a curved surface. For a list of currently supported manifolds and their parameters, see the *Howto manifold* doc page.

Note that the particles must initially be close to the manifold in question. If not, RATTLE will not be able to iterate until the constraint is satisfied, and an error is generated. For simple manifolds this can be achieved with *region* and *create\_atoms* commands, but for more complex surfaces it might be more useful to write a script.

The manifold args may be equal-style variables, like so:

```
variable R equal "ramp(5.0,3.0)"
fix shrink_sphere all nve/manifold/rattle 1e-4 10 sphere v_R
```

In this case, the manifold parameter will change in time according to the variable. This is not a problem for the time integrator as long as the change of the manifold is slow with respect to the dynamics of the particles. Note that if the manifold has to exert work on the particles because of these changes, the total energy might not be conserved.

---

### 17.128.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

---

### 17.128.5 Restrictions

This fix is part of the MANIFOLD package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

---

### 17.128.6 Related commands

*fix nvt/manifold/rattle*, *fix manifoldforce*

### 17.128.7 Default

every = 0, tchain = 3

---

(**Andersen**) Andersen, J. Comp. Phys. 52, 24, (1983).

(**Paquay**) Paquay and Kusters, Biophys. J., 110, 6, (2016). preprint available at [arXiv:1411.3019](https://arxiv.org/abs/1411.3019).



## 17.129 fix nve/noforce command

### 17.129.1 Syntax

```
fix ID group-ID nve
```

- ID, group-ID are documented in *fix* command
- nve/noforce = style name of this fix command

### 17.129.2 Examples

```
fix 3 wall nve/noforce
```

### 17.129.3 Description

Perform updates of position, but not velocity for atoms in the group each timestep. In other words, the force on the atoms is ignored and their velocity is not updated. The atom velocities are used to update their positions.

This can be useful for wall atoms, when you set their velocities, and want the wall to move (or stay stationary) in a prescribed fashion.

This can also be accomplished via the *fix setforce* command, but with *fix nve/noforce*, the forces on the wall atoms are unchanged, and can thus be printed by the *dump* command or queried with an equal-style *variable* that uses the *fcf()* group function to compute the total force on the group of atoms.

### 17.129.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.129.5 Restrictions

none

### 17.129.6 Related commands

*fix nve*

## 17.129.7 Default

none

## 17.130 fix nve/sphere command

Accelerator Variants: *nve/sphere/omp*, *nve/sphere/kk*

### 17.130.1 Syntax

```
fix ID group-ID nve/sphere
```

- ID, group-ID are documented in *fix* command
- nve/sphere = style name of this fix command
- zero or more keyword/value pairs may be appended
- keyword = *update* or *disc*

*update* value = *dipole* or *dipole/dlm*

*dipole* = update orientation of dipole moment during integration

*dipole/dlm* = use DLM integrator to update dipole orientation

*disc* value = none = treat particles as 2d discs, not spheres

### 17.130.2 Examples

```
fix 1 all nve/sphere
fix 1 all nve/sphere update dipole
fix 1 all nve/sphere disc
fix 1 all nve/sphere update dipole/dlm
```

### 17.130.3 Description

Perform constant NVE integration to update position, velocity, and angular velocity for finite-size spherical particles in the group each timestep. V is volume; E is energy. This creates a system trajectory consistent with the microcanonical ensemble.

This fix differs from the *fix nve* command, which assumes point particles and only updates their position and velocity.

If the *update* keyword is used with the *dipole* value, then the orientation of the dipole moment of each particle is also updated during the time integration. This option should be used for models where a dipole moment is assigned to finite-size particles, e.g. spheroids via use of the *atom\_style hybrid sphere dipole* command.

The default dipole orientation integrator can be changed to the Dullweber-Leimkuhler-McLachlan integration scheme (*Dullweber*) when using *update* with the value *dipole/dlm*. This integrator is symplectic and time-reversible, giving better energy conservation and allows slightly longer timesteps at only a small additional computational cost.

If the *disc* keyword is used, then each particle is treated as a 2d disc (circle) instead of as a sphere. This is only possible for 2d simulations, as defined by the *dimension* keyword. The only difference between discs and spheres in this context is their moment of inertia, as used in the time integration.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 17.130.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.130.5 Restrictions

This fix requires that atoms store torque and angular velocity (*omega*) and a radius as defined by the *atom\_style sphere* command. If the *dipole* keyword is used, then they must also store a dipole moment as defined by the *atom\_style dipole* command.

All particles in the group must be finite-size spheres. They cannot be point particles.

Use of the *disc* keyword is only allowed for 2d simulations, as defined by the *dimension* keyword.

### 17.130.6 Related commands

*fix nve*, *fix nve/asphere*

### 17.130.7 Default

none

---

**(Dullweber)** Dullweber, Leimkuhler and McLachlan, J Chem Phys, 107, 5840 (1997).

## 17.131 fix nve/spin command

### 17.131.1 Syntax

<code>fix ID group-ID nve/spin keyword values</code>
--

- ID, group-ID are documented in *fix* command
- nve/spin = style name of this fix command

- keyword = *lattice*

*lattice* value = *moving* or *frozen*

*moving* = integrate both spin and atomic degrees of freedom

*frozen* = integrate spins on a fixed lattice

### 17.131.2 Examples

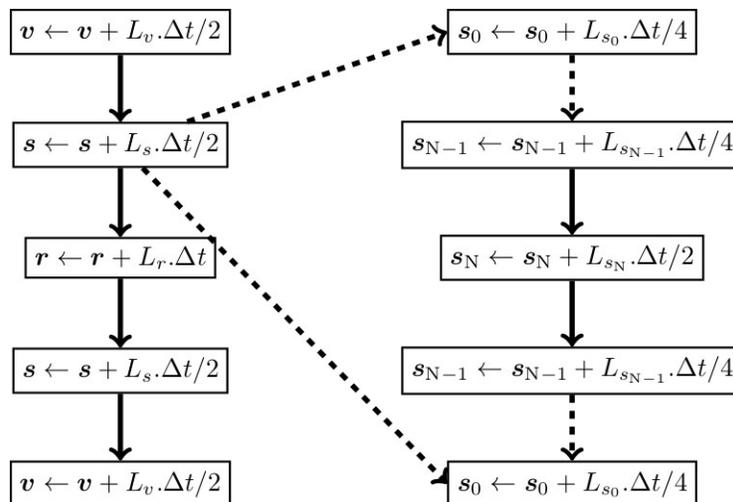
```
fix 3 all nve/spin lattice moving
fix 1 all nve/spin lattice frozen
```

### 17.131.3 Description

Perform a symplectic integration for the spin or spin-lattice system.

The *lattice* keyword defines if the spins are integrated on a lattice of fixed atoms (*lattice* = *frozen*), or if atoms are moving (*lattice* = *moving*). The first case corresponds to a spin dynamics calculation, and the second to a spin-lattice calculation. By default a spin-lattice integration is performed (*lattice* = *moving*).

The *nve/spin* fix applies a Suzuki-Trotter decomposition to the equations of motion of the spin lattice system, following the scheme:



according to the implementation reported in (*Omelyan*).

A sectoring method enables this scheme for parallel calculations. The implementation of this sectoring algorithm is reported in (*Tranchida*).

### 17.131.4 Restrictions

This fix style can only be used if LAMMPS was built with the SPIN package. See the *Build package* page for more info.

To use the spin algorithm, it is necessary to define a map with the atom\_modify command. Typically, by adding the command:

```
atom_modify map array
```

before you create the simulation box. Note that the keyword “hash” instead of “array” is also valid.

### 17.131.5 Related commands

*atom\_style spin, fix nve*

### 17.131.6 Default

The option default is lattice = moving.

---

**(Omelyan)** Omelyan, Mryglod, and Folk. Phys. Rev. Lett. 86(5), 898. (2001).

**(Tranchida)** Tranchida, Plimpton, Thibaudau and Thompson, Journal of Computational Physics, 372, 406-425, (2018).

## 17.132 fix nve/tri command

### 17.132.1 Syntax

```
fix ID group-ID nve/tri
```

- ID, group-ID are documented in *fix* command
- nve/tri = style name of this fix command

### 17.132.2 Examples

```
fix 1 all nve/tri
```

### 17.132.3 Description

Perform constant NVE integration to update position, velocity, orientation, and angular momentum for triangular particles in the group each timestep. V is volume; E is energy. This creates a system trajectory consistent with the microcanonical ensemble. See the *Howto spherical* page for an overview of using triangular particles.

This fix differs from the *fix nve* command, which assumes point particles and only updates their position and velocity.

### 17.132.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.132.5 Restrictions

This fix is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that particles be triangles as defined by the *atom\_style tri* command.

### 17.132.6 Related commands

*fix nve*, *fix nve/asphere*

### 17.132.7 Default

none

## 17.133 fix nvk command

### 17.133.1 Syntax

```
fix ID group-ID nvk
```

- ID, group-ID are documented in *fix* command
- nvk = style name of this fix command

### 17.133.2 Examples

```
fix 1 all nvk
```

### 17.133.3 Description

Perform constant kinetic energy integration using the Gaussian thermostat to update position and velocity for atoms in the group each timestep. V is volume; K is kinetic energy. This creates a system trajectory consistent with the isokinetic ensemble.

The equations of motion used are those of Minary et al in (*Minary*), a variant of those initially given by Zhang in (*Zhang*).

The kinetic energy will be held constant at its value given when *fix nvk* is initiated. If a different kinetic energy is desired, the *velocity* command should be used to change the kinetic energy prior to this fix.

### 17.133.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.133.5 Restrictions

The Gaussian thermostat only works when it is applied to all atoms in the simulation box. Therefore, the group must be set to all.

This fix has not yet been implemented to work with the RESPA integrator.

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.133.6 Related commands

none

### 17.133.7 Default

none

---

**(Minary)** Minary, Martyna, and Tuckerman, J Chem Phys, 18, 2510 (2003).

**(Zhang)** Zhang, J Chem Phys, 106, 6102 (1997).

## 17.134 fix nvt/asphere command

Accelerator Variants: *nvt/asphere/omp*

### 17.134.1 Syntax

```
fix ID group-ID nvt/asphere keyword value ...
```

- ID, group-ID are documented in *fix* command
- nvt/asphere = style name of this fix command
- additional thermostat related keyword/value pairs from the *fix nvt* command can be appended

## 17.134.2 Examples

```
fix 1 all nvt/asphere temp 300.0 300.0 100.0
fix 1 all nvt/asphere temp 300.0 300.0 100.0 drag 0.2
```

## 17.134.3 Description

Perform constant NVT integration to update position, velocity, orientation, and angular velocity each timestep for aspherical or ellipsoidal particles in the group using a Nose/Hoover temperature thermostat. V is volume; T is temperature. This creates a system trajectory consistent with the canonical ensemble.

This fix differs from the *fix nvt* command, which assumes point particles and only updates their position and velocity.

The thermostat is applied to both the translational and rotational degrees of freedom for the aspherical particles, assuming a compute is used which calculates a temperature that includes the rotational degrees of freedom (see below). The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

Additional parameters affecting the thermostat are specified by keywords and values documented with the *fix nvt* command. See, for example, discussion of the *temp* and *drag* keywords.

This fix computes a temperature each timestep. To do this, the fix creates its own compute of style “temp/asphere”, as if this command had been issued:

```
compute fix-ID_temp group-ID temp/asphere
```

See the *compute temp/asphere* command for details. Note that the ID of the new compute is the fix-ID + underscore + “temp”, and the group for the new compute is the same as the fix group.

Note that this is NOT the compute used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp*. This means you can change the attributes of this fix’s temperature (e.g. its degrees-of-freedom) via the *compute\_modify* command or print this temperature during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.



### 17.134.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover thermostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a *compute* you have defined to this fix which will be used in its thermostating procedure.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix nvt* command.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

### 17.134.5 Restrictions

This fix is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular momentum and a quaternion as defined by the *atom\_style ellipsoid* command.

All particles in the group must be finite-size. They cannot be point particles, but they can be aspherical or spherical as defined by their shape attribute.

### 17.134.6 Related commands

*fix nvt*, *fix nve\_asphere*, *fix npt\_asphere*, *fix\_modify*

### 17.134.7 Default

none

## 17.135 fix nvt/body command

### 17.135.1 Syntax

```
fix ID group-ID nvt/body keyword value ...
```

- ID, group-ID are documented in *fix* command
- nvt/body = style name of this fix command
- additional thermostat related keyword/value pairs from the *fix nvt* command can be appended

## 17.135.2 Examples

```
fix 1 all nvt/body temp 300.0 300.0 100.0
fix 1 all nvt/body temp 300.0 300.0 100.0 drag 0.2
```

## 17.135.3 Description

Perform constant NVT integration to update position, velocity, orientation, and angular velocity each timestep for body particles in the group using a Nose/Hoover temperature thermostat. V is volume; T is temperature. This creates a system trajectory consistent with the canonical ensemble.

This fix differs from the *fix nvt* command, which assumes point particles and only updates their position and velocity.

The thermostat is applied to both the translational and rotational degrees of freedom for the body particles, assuming a compute is used which calculates a temperature that includes the rotational degrees of freedom (see below). The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

Additional parameters affecting the thermostat are specified by keywords and values documented with the *fix nvt* command. See, for example, discussion of the *temp* and *drag* keywords.

This fix computes a temperature each timestep. To do this, the fix creates its own compute of style “temp/body”, as if this command had been issued:

```
compute fix-ID_temp group-ID temp/body
```

See the *compute temp/body* command for details. Note that the ID of the new compute is the fix-ID + underscore + “temp”, and the group for the new compute is the same as the fix group.

Note that this is NOT the compute used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp*. This means you can change the attributes of this fix’s temperature (e.g. its degrees-of-freedom) via the *compute\_modify* command or print this temperature during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

### 17.135.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover thermostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a *compute* you have defined to this fix which will be used in its thermostating procedure.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix nvt* command.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

### 17.135.5 Restrictions

This fix is part of the BODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular momentum and a quaternion as defined by the *atom\_style body* command.

### 17.135.6 Related commands

*fix nvt, fix nve\_body, fix npt\_body, fix\_modify*

### 17.135.7 Default

none

## 17.136 fix nvt/manifold/rattle command

### 17.136.1 Syntax

```
fix ID group-ID nvt/manifold/rattle tol maxit manifold manifold-args keyword value ...
```

- ID, group-ID are documented in *fix* command
- nvt/manifold/rattle = style name of this fix command
- tol = tolerance to which Newton iteration must converge
- maxit = maximum number of iterations to perform
- manifold = name of the manifold
- manifold-args = parameters for the manifold
- one or more keyword/value pairs may be appended

keyword = *temp* or *tchain* or *every*  
*temp* values = Tstart Tstop Tdamp  
Tstart, Tstop = external temperature at start/end of run  
Tdamp = temperature damping parameter (time units)  
*tchain* value = N  
N = length of thermostat chain (1 = single thermostat)  
*every* value = N  
N = print info about iteration every N steps. N = 0 means no output

## 17.136.2 Examples

```
fix 1 all nvt/manifold/rattle 1e-4 10 cylinder 3.0 temp 1.0 1.0 10.0
```

---

## 17.136.3 Description

This fix combines the RATTLE-based (*Andersen*) time integrator of *fix nve/manifold/rattle* (*Paquay*) with a Nose-Hoover-chain thermostat to sample the canonical ensemble of particles constrained to a curved surface (manifold). This sampling does suffer from discretization bias of  $O(dt)$ . For a list of currently supported manifolds and their parameters, see the *Howto manifold* doc page.

---

## 17.136.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

---

## 17.136.5 Restrictions

This fix is part of the MANIFOLD package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

---

## 17.136.6 Related commands

*fix nve/manifold/rattle*, *fix manifoldforce* **Default:** every = 0

---

(**Andersen**) Andersen, J. Comp. Phys. 52, 24, (1983).

(**Paquay**) Paquay and Kusters, Biophys. J., 110, 6, (2016). preprint available at [arXiv:1411.3019](https://arxiv.org/abs/1411.3019).

## 17.137 fix nvt/sllod command

Accelerator Variants: *nvt/sllod/intel*, *nvt/sllod/omp*, *nvt/sllod/kk*

### 17.137.1 Syntax

```
fix ID group-ID nvt/sllod keyword value ...
```

- ID, group-ID are documented in *fix* command
- nvt/sllod = style name of this fix command
- additional thermostat related keyword/value pairs from the *fix nvt* command can be appended

### 17.137.2 Examples

```
fix 1 all nvt/sllod temp 300.0 300.0 100.0
fix 1 all nvt/sllod temp 300.0 300.0 100.0 drag 0.2
```

### 17.137.3 Description

Perform constant NVT integration to update positions and velocities each timestep for atoms in the group using a Nose/Hoover temperature thermostat. V is volume; T is temperature. This creates a system trajectory consistent with the canonical ensemble.

This thermostat is used for a simulation box that is changing size and/or shape, for example in a non-equilibrium MD (NEMD) simulation. The size/shape change is induced by use of the *fix deform* command, so each point in the simulation box can be thought of as having a “streaming” velocity. This position-dependent streaming velocity is subtracted from each atom’s actual velocity to yield a thermal velocity which is used for temperature computation and thermostatting. For example, if the box is being sheared in x, relative to y, then points at the bottom of the box (low y) have a small x velocity, while points at the top of the box (hi y) have a large x velocity. These velocities do not contribute to the thermal “temperature” of the atom.

---

**Note:** *Fix deform* has an option for remapping either atom coordinates or velocities to the changing simulation box. To use *fix nvt/sllod*, *fix deform* should NOT remap atom positions, because *fix nvt/sllod* adjusts the atom positions and velocities to create a velocity profile that matches the changing box size/shape. *Fix deform* SHOULD remap atom velocities when atoms cross periodic boundaries since that is consistent with maintaining the velocity profile created by *fix nvt/sllod*. LAMMPS will give an error if this setting is not consistent.

---

The SLLOD equations of motion, originally proposed by Hoover and Ladd (see (*Evans and Morriss*)), were proven to be equivalent to Newton’s equations of motion for shear flow by (*Evans and Morriss*). They were later shown to generate the desired velocity gradient and the correct production of work by stresses for all forms of homogeneous flow by (*Daivis and Todd*). The LAMMPS implementation corresponds to the p-SLLOD variant of SLLOD. (*Edwards*). The equations of motion are coupled to a Nose/Hoover chain thermostat in a velocity Verlet formulation, closely following the implementation used for the *fix nvt* command.

---

**Note:** A recent (2017) book by (*Daivis and Todd*) discusses use of the SLLOD method and non-equilibrium MD (NEMD) thermostatting generally, for both simple and complex fluids, e.g. molecular systems. The latter can be tricky to do correctly.

---

Additional parameters affecting the thermostat are specified by keywords and values documented with the *fix nvt* command. See, for example, discussion of the *temp* and *drag* keywords.

This fix computes a temperature each timestep. To do this, the fix creates its own compute of style “temp/deform”, as if this command had been issued:

```
compute fix-ID_temp group-ID temp/deform
```

See the *compute temp/deform* command for details. Note that the ID of the new compute is the fix-ID + underscore + “temp”, and the group for the new compute is the same as the fix group.

Note that this is NOT the compute used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp*. This means you can change the attributes of this fix’s temperature (e.g. its degrees-of-freedom) via the *compute\_modify* command or print this temperature during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 17.137.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover thermostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a *compute* you have defined to this fix which will be used in its thermostating procedure.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix nvt* command.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

## 17.137.5 Restrictions

This fix works best without Nose-Hoover chain thermostats, i.e. using `tchain = 1`. Setting `tchain` to larger values can result in poor equilibration.

## 17.137.6 Related commands

*fix nve*, *fix nvt*, *fix temp/rescale*, *fix langevin*, *fix\_modify*, *compute temp/deform*

## 17.137.7 Default

Same as *fix nvt*, except `tchain = 1`.

---

**(Evans and Morriss)** Evans and Morriss, Phys Rev A, 30, 1528 (1984).

**(Daivis and Todd)** Daivis and Todd, J Chem Phys, 124, 194103 (2006).

**(Edwards)** Edwards, Baig, and Keffer, J Chem Phys 124, 194104 (2006).

**(Daivis and Todd)** Daivis and Todd, Nonequilibrium Molecular Dynamics (book), Cambridge University Press, <https://doi.org/10.1017/9781139017848>, (2017).

## 17.138 fix nvt/sllod/eff command

### 17.138.1 Syntax

```
fix ID group-ID nvt/sllod/eff keyword value ...
```

- ID, group-ID are documented in *fix* command
- nvt/sllod/eff = style name of this fix command
- additional thermostat related keyword/value pairs from the *fix nvt/eff* command can be appended

### 17.138.2 Examples

```
fix 1 all nvt/sllod/eff temp 300.0 300.0 0.1
fix 1 all nvt/sllod/eff temp 300.0 300.0 0.1 drag 0.2
```

### 17.138.3 Description

Perform constant NVT integration to update positions and velocities each timestep for nuclei and electrons in the group for the *electron force field* model, using a Nose/Hoover temperature thermostat. V is volume; T is temperature. This creates a system trajectory consistent with the canonical ensemble.

The operation of this fix is exactly like that described by the *fix nvt/sllod* command, except that the radius and radial velocity of electrons are also updated and thermostatted. Likewise the temperature calculated by the fix, using the compute it creates (as discussed in the *fix nvt*, *npt*, and *nph* doc page), is performed with a *compute temp/deform/eff* command that includes the eFF contribution to the temperature from the electron radial velocity.

### 17.138.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover thermostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a *compute* you have defined to this fix which will be used in its thermostating procedure.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix nvt/eff* command.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

### 17.138.5 Restrictions

This fix is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix works best without Nose-Hoover chain thermostats, i.e. using *tchain = 1*. Setting *tchain* to larger values can result in poor equilibration.

### 17.138.6 Related commands

*fix nve/eff*, *fix nvt/eff*, *fix langevin/eff*, *fix nvt/sllod*, *fix\_modify*, *compute temp/deform/eff*

### 17.138.7 Default

Same as *fix nvt/eff*, except *tchain = 1*.

---

(Tuckerman) Tuckerman, Mundy, Balasubramanian, Klein, J Chem Phys, 106, 5615 (1997).

## 17.139 fix nvt/sphere command

Accelerator Variants: *nvt/sphere/omp*

### 17.139.1 Syntax

```
fix ID group-ID nvt/sphere keyword value ...
```

- ID, group-ID are documented in *fix* command
- nvt/sphere = style name of this fix command
- zero or more keyword/value pairs may be appended
- keyword = *disc*



`disc value = none = treat particles as 2d discs, not spheres`

- NOTE: additional thermostat and dipole related keyword/value pairs from the `fix nvt` command can be appended

## 17.139.2 Examples

```
fix 1 all nvt/sphere temp 300.0 300.0 100.0
fix 1 all nvt/sphere temp 300.0 300.0 100.0 disc
fix 1 all nvt/sphere temp 300.0 300.0 100.0 drag 0.2
fix 1 all nvt/sphere temp 300.0 300.0 100.0 update dipole
```

## 17.139.3 Description

Perform constant NVT integration to update position, velocity, and angular velocity each timestep for finite-size spherical particles in the group using a Nose/Hoover temperature thermostat. V is volume; T is temperature. This creates a system trajectory consistent with the canonical ensemble.

This fix differs from the `fix nvt` command, which assumes point particles and only updates their position and velocity.

The thermostat is applied to both the translational and rotational degrees of freedom for the spherical particles, assuming a compute is used which calculates a temperature that includes the rotational degrees of freedom (see below). The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

If the `disc` keyword is used, then each particle is treated as a 2d disc (circle) instead of as a sphere. This is only possible for 2d simulations, as defined by the `dimension` keyword. The only difference between discs and spheres in this context is their moment of inertia, as used in the time integration.

Additional parameters affecting the thermostat are specified by keywords and values documented with the `fix nvt` command. See, for example, discussion of the `temp` and `drag` keywords.

This fix computes a temperature each timestep. To do this, the fix creates its own compute of style “temp/sphere”, as if this command had been issued:

```
compute fix-ID_temp group-ID temp/sphere
```

See the `compute temp/sphere` command for details. Note that the ID of the new compute is the fix-ID + underscore + “temp”, and the group for the new compute is the same as the fix group.

Note that this is NOT the compute used by thermodynamic output (see the `thermo_style` command) with ID = `thermo_temp`. This means you can change the attributes of this fix’s temperature (e.g. its degrees-of-freedom) via the `compute_modify` command or print this temperature during thermodynamic output via the `thermo_style custom` command using the appropriate compute-ID. It also means that changing attributes of `thermo_temp` will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with `compute commands` that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the `fix_modify` command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual `compute temp commands` to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

#### 17.139.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the Nose/Hoover thermostat to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a *compute* you have defined to this fix which will be used in its thermostating procedure.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes the same global scalar and global vector of quantities as does the *fix nvt* command.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

#### 17.139.5 Restrictions

This fix requires that atoms store torque and angular velocity (omega) and a radius as defined by the *atom\_style sphere* command.

All particles in the group must be finite-size spheres. They cannot be point particles.

Use of the *disc* keyword is only allowed for 2d simulations, as defined by the *dimension* keyword.

#### 17.139.6 Related commands

*fix nvt*, *fix nve\_sphere*,  
*fix nvt\_asphere*, *fix npt\_sphere*, *fix\_modify*

## 17.139.7 Default

none

## 17.140 fix oneway command

### 17.140.1 Syntax

```
fix ID group-ID oneway N region-ID direction
```

- ID, group-ID are documented in *fix* command
- oneway = style name of this fix command
- N = apply this fix every this many timesteps
- region-ID = ID of region where fix is active
- direction = x or -x or y or -y or z or -z = coordinate and direction of the oneway constraint

### 17.140.2 Examples

```
fix 1 ions oneway 10 semi -x
fix 2 all oneway 1 left -z
fix 3 all oneway 1 right z
```

### 17.140.3 Description

Enforce that particles in the group and in a given region can only move in one direction. This is done by reversing a particle's velocity component, if it has the wrong sign in the specified dimension. The effect is that the particle moves in one direction only.

This can be used, for example, as a simple model of a semi-permeable membrane, or as an implementation of Maxwell's demon.

### 17.140.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.140.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.140.6 Related commands

*fix wall/reflect* command

### 17.140.7 Default

none

## 17.141 fix orient/fcc command

## 17.142 fix orient/bcc command

### 17.142.1 Syntax

```
fix ID group-ID orient/fcc nstats dir alat dE cutlo cuthi file0 file1
fix ID group-ID orient/bcc nstats dir alat dE cutlo cuthi file0 file1
```

- ID, group-ID are documented in *fix* command
- nstats = print stats every this many steps, 0 = never
- dir = 0/1 for which crystal is used as reference
- alat = fcc/bcc cubic lattice constant (distance units)
- dE = energy added to each atom (energy units)
- cutlo,cuthi = values between 0.0 and 1.0, cutlo < cuthi
- file0,file1 = files that specify orientation of each grain

### 17.142.2 Examples

```
fix gb all orient/fcc 0 1 4.032008 0.001 0.25 0.75 xi.vec chi.vec
fix gb all orient/bcc 0 1 2.882 0.001 0.25 0.75 ngb.left ngb.right
```

### 17.142.3 Description

The fix applies an orientation-dependent force to atoms near a planar grain boundary which can be used to induce grain boundary migration (in the direction perpendicular to the grain boundary plane). The motivation and explanation of this force and its application are described in (*Janssens*). The adaptation to bcc crystals is described in (*Wicaksono1*). The computed force is only applied to atoms in the fix group.

The basic idea is that atoms in one grain (on one side of the boundary) have a potential energy  $dE$  added to them. Atoms in the other grain have 0.0 potential energy added. Atoms near the boundary (whose neighbor environment is intermediate between the two grain orientations) have an energy between 0.0 and  $dE$  added. This creates an effective driving force to reduce the potential energy of atoms near the boundary by pushing them towards one of the grain orientations. For  $dir = 1$  and  $dE > 0$ , the boundary will thus move so that the grain described by file0 grows and the grain described by file1 shrinks. Thus this fix is designed for simulations of two-grain systems, either with one grain boundary and free surfaces parallel to the boundary, or a system with periodic boundary conditions and two equal and opposite grain boundaries. In either case, the entire system can displace during the simulation, and such motion should be accounted for in measuring the grain boundary velocity.

The potential energy added to atom I is given by these formulas

$$\xi_i = \sum_{j=1}^{12} |\mathbf{r}_j - \mathbf{r}_j^I| \quad (1)$$

$$\xi_{IJ} = \sum_{j=1}^{12} |\mathbf{r}_j^J - \mathbf{r}_j^I| \quad (2)$$

$$\xi_{low} = cutlo \xi_{IJ} \quad (3)$$

$$\xi_{high} = cuthi \xi_{IJ} \quad (4)$$

$$\omega_i = \frac{\pi}{2} \frac{\xi_i - \xi_{low}}{\xi_{high} - \xi_{low}} \quad (5)$$

$$\begin{aligned} u_i &= 0 & \text{for } \xi_i < \xi_{low} \\ &= dE \frac{1 - \cos(2\omega_i)}{2} & \text{for } \xi_{low} < \xi_i < \xi_{high} \\ &= dE & \text{for } \xi_{high} < \xi_i \end{aligned} \quad (6)$$

which are fully explained in (*Janssens*). For fcc crystals this order parameter  $\xi_i$  for atom I in equation (1) is a sum over the 12 nearest neighbors of atom I. For bcc crystals it is the corresponding sum of the 8 nearest neighbors.  $\mathbf{r}_j$  is the vector from atom I to its neighbor J, and  $\mathbf{R}_j^I$  is a vector in the reference (perfect) crystal. That is, if  $dir = 0/1$ , then  $\mathbf{R}_j^I$  is a vector to an atom coord from file 0/1. Equation (2) gives the expected value of the order parameter  $\xi_{IJ}$  in the other grain.  $hi$  and  $lo$  cutoffs are defined in equations (3) and (4), using the input parameters *cutlo* and *cuthi* as thresholds to avoid adding grain boundary energy when the deviation in the order parameter from 0 or 1 is small (e.g. due to thermal fluctuations in a perfect crystal). The added potential energy  $U_i$  for atom I is given in equation (6) where it is interpolated between 0 and  $dE$  using the two threshold  $\xi_i$  values and the  $\omega_i$  value of equation (5).

The derivative of this energy expression gives the force on each atom which thus depends on the orientation of its neighbors relative to the 2 grain orientations. Only atoms near the grain boundary feel a net force which tends to drive them to one of the two grain orientations.

In equation (1), the reference vector used for each neighbor is the reference vector closest to the actual neighbor position. This means it is possible two different neighbors will use the same reference vector. In such cases, the atom in question is far from a perfect orientation and will likely receive the full  $dE$  addition, so the effect of duplicate reference vector usage is small.

The *dir* parameter determines which grain wants to grow at the expense of the other. A value of 0 means the first grain will shrink; a value of 1 means it will grow. This assumes that *dE* is positive. The reverse will be true if *dE* is negative.

The *alat* parameter is the cubic lattice constant for the fcc or bcc material and is only used to compute a cutoff distance of  $1.57 * alat / \sqrt{2}$  for finding the 12 or 8 nearest neighbors of each atom (which should be valid for an fcc or bcc crystal). A longer/shorter cutoff can be imposed by adjusting *alat*. If a particular atom has less than 12 or 8 neighbors within the cutoff, the order parameter of equation (1) is effectively multiplied by 12 or 8 divided by the actual number of neighbors within the cutoff.

The *dE* parameter is the maximum amount of additional energy added to each atom in the grain which wants to shrink.

The *cutlo* and *cuthi* parameters are used to reduce the force added to bulk atoms in each grain far away from the boundary. An atom in the bulk surrounded by neighbors at the ideal grain orientation would compute an order parameter of 0 or 1 and have no force added. However, thermal vibrations in the solid will cause the order parameters to be greater than 0 or less than 1. The cutoff parameters mask this effect, allowing forces to only be added to atoms with order-parameters between the cutoff values.

*File0* and *file1* are filenames for the two grains which each contain 6 vectors (6 lines with 3 values per line) which specify the grain orientations. Each vector is a displacement from a central atom (0,0,0) to a nearest neighbor atom in an fcc lattice at the proper orientation. The vector lengths should all be identical since an fcc lattice has a coordination number of 12. Only 6 are listed due to symmetry, so the list must include one from each pair of equal-and-opposite neighbors. A pair of orientation files for a Sigma=5 tilt boundary are shown below. A tutorial that can help for writing the orientation files is given in ([Wicaksono2](#))

#### 17.142.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy of atom interactions with the grain boundary driving force to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify respa* option is supported by these fixes. This allows to set at which level of the *r-RESPA* integrator a fix is adding its forces. Default is the outermost level.

This fix calculates a global scalar which can be accessed by various *output commands*. The scalar is the potential energy change due to this fix. The scalar value calculated by this fix is “extensive”.

This fix also calculates a per-atom array which can be accessed by various *output commands*. The array stores the order parameter Xi and normalized order parameter (0 to 1) for each atom. The per-atom values can be accessed on any timestep.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

This fix is not invoked during *energy minimization*.

#### 17.142.5 Restrictions

These fixes are part of the ORIENT package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

These fixes should only be used with fcc or bcc lattices.

## 17.142.6 Related commands

*fix\_modify*

## 17.142.7 Default

none

(**Janssens**) Janssens, Olmsted, Holm, Foiles, Plimpton, Derlet, Nature Materials, 5, 124-127 (2006).

(**Wicaksono1**) Wicaksono, Sinclair, Militzer, Computational Materials Science, 117, 397-405 (2016).

(**Wicaksono2**) Wicaksono, figshare, <https://doi.org/10.6084/m9.figshare.1488628.v1> (2015).

For illustration purposes, here are example files that specify a Sigma=5 <100> tilt boundary. This is for a lattice constant of 3.5706 Angs.

file0:

0.798410432046075	1.7853000000000000	1.596820864092150
-0.798410432046075	1.7853000000000000	-1.596820864092150
2.395231296138225	0.0000000000000000	0.798410432046075
0.798410432046075	0.0000000000000000	-2.395231296138225
1.596820864092150	1.7853000000000000	-0.798410432046075
1.596820864092150	-1.7853000000000000	-0.798410432046075

file1:

-0.798410432046075	1.7853000000000000	1.596820864092150
0.798410432046075	1.7853000000000000	-1.596820864092150
0.798410432046075	0.0000000000000000	2.395231296138225
2.395231296138225	0.0000000000000000	-0.798410432046075
1.596820864092150	1.7853000000000000	0.798410432046075
1.596820864092150	-1.7853000000000000	0.798410432046075

## 17.143 fix orient/eco command

```
fix ID group-ID orient/eco u0 eta cutoff orientationsFile
```

- ID, group-ID are documented in fix command
- u0 = energy added to each atom (energy units)
- eta = cutoff value (usually 0.25)
- cutoff = cutoff radius for orientation parameter calculation
- orientationsFile = file that specifies orientation of each grain

## 17.143.1 Examples

```
fix gb all orient/eco 0.08 0.25 3.524 sigma5.ori
```

## 17.143.2 Description

The fix applies a synthetic driving force to a grain boundary which can be used for the investigation of grain boundary motion. The affiliation of atoms to either of the two grains forming the grain boundary is determined from an orientation-dependent order parameter as described in (*Ulomek*). The potential energy of atoms is either increased by an amount of  $0.5*u0$  or  $-0.5*u0$  according to the orientation of the surrounding crystal. This creates a potential energy gradient which pushes atoms near the grain boundary to orient according to the energetically favorable grain orientation. This fix is designed for applications in bicrystal system with one grain boundary and open ends, or two opposite grain boundaries in a periodic system. In either case, the entire system can experience a displacement during the simulation which needs to be accounted for in the evaluation of the grain boundary velocity. While the basic method is described in (*Ulomek*), the implementation follows the efficient implementation from (*Schrott & Mohles*). The synthetic potential energy added to an atom  $j$  is given by the following formulas

$$w(|\vec{r}_{jk}|) = w_{jk} = \begin{cases} \frac{|\vec{r}_{jk}|^4}{r_{\text{cut}}^4} - 2\frac{|\vec{r}_{jk}|^2}{r_{\text{cut}}^2} + 1, & |\vec{r}_{jk}| < r_{\text{cut}} \\ 0, & |\vec{r}_{jk}| \geq r_{\text{cut}} \end{cases}$$

$$\chi_j = \frac{1}{N} \sum_{l=1}^3 \left[ |\psi_l^{\text{I}}(\vec{r}_j)|^2 - |\psi_l^{\text{II}}(\vec{r}_j)|^2 \right]$$

$$\psi_l^{\text{X}}(\vec{r}_j) = \sum_{k \in \Gamma_j} w_{jk} \exp(i\vec{r}_{jk} \cdot \vec{q}_l^{\text{X}})$$

$$u(\chi_j) = \frac{u_0}{2} \begin{cases} 1, & \chi_j \geq \eta \\ \sin\left(\frac{\pi\chi_j}{2\eta}\right), & -\eta < \chi_j < \eta \\ -1, & \chi_j \leq -\eta \end{cases}$$

which are fully explained in (*Ulomek*) and (*Schrott & Mohles*).

The force on each atom is the negative gradient of the synthetic potential energy. It depends on the surrounding of this atom. An atom far from the grain boundary does not experience a synthetic force as its surrounding is that of an oriented single crystal and thermal fluctuations are masked by the parameter *eta*. Near the grain boundary however, the gradient is nonzero and synthetic force terms are computed. The orientationsFile specifies the perfect oriented crystal basis vectors for the two adjoining crystals. The first three lines (line=row vector) for the energetically penalized and the last three lines for the energetically favored grain assuming  $u0$  is positive. For negative  $u0$ , this is reversed. With the *cutoff* parameter, the size of the region around each atom which is used in the order parameter computation is defined. The cutoff must be smaller than the interaction range of the MD potential. It should at least include the nearest neighbor shell. For high temperatures or low angle grain boundaries, it might be beneficial to increase the cutoff in order to get a more precise identification of the atoms surrounding. However, computation time will increase as more atoms are considered in the order parameter and force computation. It is also worth noting that the cutoff radius must not exceed the communication distance for ghost atoms in LAMMPS. With orientationsFile, the 6 oriented crystal basis vectors is specified. Each line of the input file contains the three components of a primitive lattice vector oriented according to the grain orientation in the simulation box. The first (last) three lines correspond to the primitive lattice vectors of the first (second) grain. An example for a  $\Sigma(001)$  mis-orientation is given at the end.

If no synthetic energy difference between the grains is created,  $u0 = 0$ , the force computation is omitted. In this case, still, the order parameter of the driving force is computed and can be used to track the grain boundary motion throughout the simulation.



### 17.143.3 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy of atom interactions with the grain boundary driving force to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

This fix calculates a per-atom array with 2 columns, which can be accessed by indices 1-1 by any command that uses per-atom values from a fix as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The first column is the order parameter for each atom; the second is the thermal masking value for each atom. Both are described above.

No parameter of this fix can be used with the start/stop keywords of the run command. This fix is not invoked during energy minimization.

### 17.143.4 Restrictions

This fix is part of the ORIENT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.143.5 Related commands

*fix\_modify*

*fix\_orient*

### 17.143.6 Default

none

---

(**Ulomek**) Ulomek, Brien, Foiles, Mohles, Modelling Simul. Mater. Sci. Eng. 23 (2015) 025007

(**Schratt & Mohles**) Schratt, Mohles. Comp. Mat. Sci. 182 (2020) 109774

---

For illustration purposes, here is an example file that specifies a  $\Sigma = 5\langle 001 \rangle$  tilt grain boundary. This is for a lattice constant of 3.52 Angstrom:

```
sigma5.ori:
1.671685  0.557228  1.76212
0.557228 -1.671685  1.76212
2.228913 -1.114456  0.000000
0.557228  1.671685  1.76212
1.671685 -0.557228  1.76212
2.228913  1.114456  0.000000
```

## 17.144 fix pafi command

### 17.144.1 Syntax

```
fix ID group-ID pafi compute-ID Temp Tdamp seed keyword values...
```

- ID, group-ID are documented in *fix* command
- pafi = style name of this fix command
- compute-ID = ID of a *compute property/atom* that holds data used by this fix
- Temp = desired temperature (temperature units)
- Tdamp = damping parameter (time units)
- seed = random number seed to use for white noise (positive integer)
- keyword = *overdamped* or *com*
  - overdamped* value = *yes* or *no* or 1 or 0
    - yes* or 1 = Brownian (overdamped) integration in hyperplane
    - no* or 0 = Langevin integration in hyperplane
  - com* value = *yes* or *no* or 1 or 0
    - yes* or 1 = zero linear momentum, fixing center or mass (recommended)
    - no* or 0 = do not zero linear momentum, allowing center of mass drift

### 17.144.2 Examples

```
compute pa all property/atom d_nx d_ny d_nz d_dnx d_dny d_dnz d_ddnx d_ddny d_ddnz
run 0 post no
fix hp all pafi pa 500.0 0.01 434 overdamped yes
```

### 17.144.3 Description

Perform Brownian or Langevin integration whilst constraining the system to lie in some hyperplane, which is expected to be the tangent plane to some reference pathway in a solid state system. The instantaneous value of a modified force projection is also calculated, whose time integral can be shown to be equal to the true free energy gradient along the minimum free energy path local to the reference pathway. A detailed discussion of the projection technique can be found in (*Swinburne*).

This fix can be used with LAMMPS as demonstrated in `examples/PACKAGES/pafi`, though it is primarily intended to be coupled with the PAFI C++ code, developed at <https://github.com/tomswinburne/pafi>, which distributes multiple LAMMPS workers in parallel to compute and collate hyperplane-constrained averages, allowing the calculation of free energy barriers and pathways.

A *compute property/atom* must be provided with 9 fields per atom coordinate, which in order are the x,y,z coordinates of a configuration on the reference path, the x,y,z coordinates of the path tangent (derivative of path position with path coordinate) and the x,y,z coordinates of the change in tangent (derivative of path tangent with path coordinate).

A 4-element vector is also calculated by this fix. The 4 components are the modified projected force, its square, the expected projection of the minimum free energy path tangent on the reference path tangent and the minimum image distance between the current configuration and the reference configuration, projected along the path tangent. This latter value should be essentially zero.

**Note:** When `com=yes/1`, which is recommended, the provided tangent vector must also have zero center of mass. This can be achieved by subtracting from each coordinate of the path tangent the average `x,y,z` value. The PAFI C++ code (see above) can generate these paths for use in LAMMPS.

---

**Note:** When `overdamped=yes/1`, the `Tdamp` parameter should be around 5-10 times smaller than that used in typical Langevin integration. See *fix langevin* for typical values.

---

#### 17.144.4 Restart, `fix_modify`, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the `fix_modify` options are relevant to this fix.

This fix produces a global vector each timestep which can be accessed by various *output commands*.

#### 17.144.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 17.144.6 Default

The option defaults are `com = yes`, `overdamped = no`

---

(Swinburne) Swinburne and Marinica, Physical Review Letters, 120, 1 (2018)

### 17.145 `fix pair` command

#### 17.145.1 Syntax

```
fix ID group-ID pair N pstyle name flag ...
```

- ID, group-ID are documented in *fix* command
- pair = style name of this fix command
- N = invoke this fix once every N timesteps
- pstyle = name of pair style to extract info from (e.g. eam)
- one or more name/flag pairs can be listed
- name = name of quantity the pair style allows extraction of
- flag = 1 if pair style needs to be triggered to produce data for name, 0 if not

## 17.145.2 Examples

```
fix request all pair 100 eam rho 0
fix request all pair 100 amoeba uind 0 uinp 0
```

## 17.145.3 Description

New in version 15Sep2022.

Extract per-atom quantities from a pair style and store them in this fix so they can be accessed by other LAMMPS commands, e.g. by a *dump* command or by another *fix*, *compute*, or *variable* command.

These are example use cases:

- extract per-atom density from *pair\_style eam* to a dump file
- extract induced dipoles from *pair\_style amoeba* to a dump file
- extract accuracy metrics from a machine-learned potential to trigger output when a condition is met (see the *dump\_modify skip* command)

The *N* argument determines how often the fix is invoked.

The *pstyle* argument is the name of the pair style. It can be a sub-style used in a *pair\_style hybrid* command.

One or more *name/flag* pairs of arguments follow. Each *name* is a per-atom quantity which the pair style must recognize as an extraction request. See the doc pages for individual *pair\_styles* to see what fix pair requests (if any) they support.

The *flag* setting determines whether this fix will also trigger the pair style to compute the named quantity so it can be extracted. If the quantity is always computed by the pair style, no trigger is needed; specify *flag* = 0. If the quantity is not always computed (e.g. because it is expensive to calculate), then specify *flag* = 1. This will trigger the quantity to be calculated only on timesteps it is needed. Again, see the doc pages for individual *pair\_styles* to determine which fix pair requests (if any) need to be triggered with a *flag* = 1 setting.

The per-atom data extracted from the pair style is stored by this fix as either a per-atom vector or array. If there is only one *name* argument specified and the pair style computes a single value for each atom, then this fix stores it as a per-atom vector. Otherwise a per-atom array is created, with its data in the order of the *name* arguments.

For example, *pair\_style amoeba* allows extraction of two named quantities: “uind” and “uinp”, both of which are 3-vectors for each atom, i.e. dipole moments. In the example below a 6-column per-atom array will be created. Columns 1-3 will store the “uind” values; columns 4-6 will store the “uinp” values.

```
pair_style amoeba
fix ex all pair amoeba 10 uind 0 uinp 0
```

## 17.145.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

As explained above, this fix produces a per-atom vector or array which can be accessed by various *output commands*. If an array is produced, the number of columns is the sum of the number of per-atom quantities produced by each *name* argument requested from the pair style.

## 17.145.5 Restrictions

none

## 17.145.6 Related commands

*compute pair*

## 17.145.7 Default

none

# 17.146 fix phonon command

## 17.146.1 Syntax

```
fix ID group-ID phonon N Noutput Nwait map_file prefix keyword values ...
```

- ID, group-ID are documented in *fix* command
- phonon = style name of this fix command
- N = measure the Green's function every this many timesteps
- Noutput = output the dynamical matrix every this many measurements
- Nwait = wait this many timesteps before measuring
- map\_file = *file* or *GAMMA*

*file* is the file that contains the mapping info between atom ID and the lattice ↵  
↵indices.

*GAMMA* flags to treat the whole simulation box as a unit cell, so that the mapping info can be generated internally. In this case, dynamical matrix at only the ↵  
↵gamma-point will/can be evaluated.

- prefix = prefix for output files
- one or none keyword/value pairs may be appended
- keyword = *sysdim* or *nasr*

*sysdim* value = d

d = dimension of the system, usually the same as the MD model dimension

*nasr* value = n

n = number of iterations to enforce the acoustic sum rule

## 17.146.2 Examples

```
fix 1 all phonon 20 5000 2000000 map.in LJ1D sysdim 1
fix 1 all phonon 20 5000 2000000 map.in EAM3D
fix 1 all phonon 10 5000 5000000 GAMMA EAMOD nasr 100
```

## 17.146.3 Description

Calculate the dynamical matrix from molecular dynamics simulations based on fluctuation-dissipation theory for a group of atoms.

Consider a crystal with  $N$  unit cells in three dimensions labeled  $l = (l_1, l_2, l_3)$  where  $l_i$  are integers. Each unit cell is defined by three linearly independent vectors  $\mathbf{a}_1, \mathbf{a}_2, \mathbf{a}_3$  forming a parallelepiped, containing  $K$  basis atoms labeled  $k$ .

Based on fluctuation-dissipation theory, the force constant coefficients of the system in reciprocal space are given by (Campana, Kong)

$$\Phi_{k\alpha, k'\beta}(\mathbf{q}) = k_B T \mathbf{G}_{k\alpha, k'\beta}^{-1}(\mathbf{q})$$

where  $\mathbf{G}$  is the Green's functions coefficients given by

$$\mathbf{G}_{k\alpha, k'\beta}(\mathbf{q}) = \langle \mathbf{u}_{k\alpha}(\mathbf{q}) \bullet \mathbf{u}_{k'\beta}^*(\mathbf{q}) \rangle$$

where  $\langle \dots \rangle$  denotes the ensemble average, and

$$\mathbf{u}_{k\alpha}(\mathbf{q}) = \sum_l \mathbf{u}_{lk\alpha} \exp(i\mathbf{q}\mathbf{r}_l)$$

is the  $\alpha$  component of the atomic displacement for the  $k$  th atom in the unit cell in reciprocal space at  $\mathbf{q}$ . In practice, the Green's functions coefficients can also be measured according to the following formula,

$$\mathbf{G}_{k\alpha, k'\beta}(\mathbf{q}) = \langle \mathbf{R}_{k\alpha}(\mathbf{q}) \bullet \mathbf{R}_{k'\beta}^*(\mathbf{q}) \rangle - \langle \mathbf{R} \rangle_{k\alpha}(\mathbf{q}) \bullet \langle \mathbf{R} \rangle_{k'\beta}^*(\mathbf{q})$$

where  $\mathbf{R}$  is the instantaneous positions of atoms, and  $\langle \mathbf{R} \rangle$  is the averaged atomic positions. It gives essentially the same results as the displacement method and is easier to implement in an MD code.

Once the force constant matrix is known, the dynamical matrix  $\mathbf{D}$  can then be obtained by

$$\mathbf{D}_{k\alpha, k'\beta}(\mathbf{q}) = (m_k m_{k'})^{-\frac{1}{2}} \Phi_{k\alpha, k'\beta}(\mathbf{q})$$

whose eigenvalues are exactly the phonon frequencies at  $\mathbf{q}$ .

This fix uses positions of atoms in the specified group and calculates two-point correlations. To achieve this, the positions of the atoms are examined every *Nevery* steps and are Fourier-transformed into reciprocal space, where the averaging process and correlation computation is then done. After every *Noutput* measurements, the matrix  $\mathbf{G}(\mathbf{q})$  is calculated and inverted to obtain the elastic stiffness coefficients. The dynamical matrices are then constructed and written to *prefix.bin.timestep* files in binary format and to the file *prefix.log* for each wave-vector  $\mathbf{q}$ .

A detailed description of this method can be found in (Kong2011).

The *sysdim* keyword is optional. If specified with a value smaller than the dimensionality of the LAMMPS simulation, its value is used for the dynamical matrix calculation. For example, using LAMMPS to model a 2D or 3D system, the phonon dispersion of a 1D atomic chain can be computed using *sysdim* = 1.

The *nasr* keyword is optional. An iterative procedure is employed to enforce the acoustic sum rule on  $\Phi$  at  $\Gamma$ , and the number provided by keyword *nasr* gives the total number of iterations. For a system whose unit cell has only one atom, *nasr* = 1 is sufficient; for other systems, *nasr* = 10 is typically sufficient.

The *map\_file* contains the mapping information between the lattice indices and the atom IDs, which tells the code which atom sits at which lattice point; the lattice indices start from 0. An auxiliary code, *latgen*, can be employed to generate the compatible map file for various crystals.

In case one simulates a non-periodic system, where the whole simulation box is treated as a unit cell, one can set *map\_file* as *GAMMA*, so that the mapping info will be generated internally and a file is not needed. In this case, the dynamical matrix at only the gamma-point will/can be evaluated. Please keep in mind that *fix-phonon* is designed for crystals, it will be inefficient and even degrade the performance of *lammmps* in case the unit cell is too large.

The calculated dynamical matrix elements are written out in *energy/distance<sup>2</sup>/mass* units. The coordinates for *q* points in the log file is in the units of the basis vectors of the corresponding reciprocal lattice.

#### 17.146.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify temp* option is supported by this fix. You can use it to change the temperature compute from *thermo\_temp* to the one that reflects the true temperature of atoms in the group.

No global scalar or vector or per-atom quantities are stored by this fix for access by various *output commands*.

Instead, this fix outputs its initialization information (including mapping information) and the calculated dynamical matrices to the file *prefix.log*, with the specified *prefix*. The dynamical matrices are also written to files *prefix.bin.timestep* in binary format. These can be read by the post-processing tool in *tools/phonon* to compute the phonon density of states and/or phonon dispersion curves.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

This fix is not invoked during *energy minimization*.

#### 17.146.5 Restrictions

This fix assumes a crystalline system with periodical lattice. The temperature of the system should not exceed the melting temperature to keep the system in its solid state.

This fix is part of the PHONON package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires LAMMPS be built with an FFT library. See the *Build settings* page for details.

#### 17.146.6 Related commands

*compute msd, dynamical\_matrix*

#### 17.146.7 Default

The option defaults are *sysdim* = the same dimension as specified by the *dimension* command, and *nasr* = 20.

---

**(Campana)** C. Campana and M. H. Muser, *Practical Green's function approach to the simulation of elastic semi-infinite solids*, Phys. Rev. B [74], 075420 (2006)

**(Kong)** L.T. Kong, G. Bartels, C. Campana, C. Denniston, and Martin H. Muser, *Implementation of Green's function molecular dynamics: An extension to LAMMPS*, Computer Physics Communications [180](6):1004-1010 (2009).

L.T. Kong, C. Denniston, and Martin H. Muser, *An improved version of the Green's function molecular dynamics method*, *Computer Physics Communications* [182](2):540-541 (2011).

(Kong2011) L.T. Kong, *Phonon dispersion measured directly from molecular dynamics simulations*, *Computer Physics Communications* [182](10):2201-2207, (2011).

## 17.147 fix pimd command

### 17.147.1 Syntax

```
fix ID group-ID pimd keyword value ...
```

- ID, group-ID are documented in *fix* command
- pimd = style name of this fix command
- zero or more keyword/value pairs may be appended
- keyword = *method* or *fmass* or *sp* or *temp* or *nhc*  
*method* value = *pimd* or *nmpimd* or *cmd*  
*fmass* value = scaling factor on mass  
*sp* value = scaling factor on Planck constant  
*temp* value = temperature (temperarate units)  
*nhc* value = Nc = number of chains in Nose-Hoover thermostat

### 17.147.2 Examples

```
fix 1 all pimd method nmpimd fmass 1.0 sp 2.0 temp 300.0 nhc 4
```

### 17.147.3 Description

This command performs quantum molecular dynamics simulations based on the Feynman path integral to include effects of tunneling and zero-point motion. In this formalism, the isomorphism of a quantum partition function for the original system to a classical partition function for a ring-polymer system is exploited, to efficiently sample configurations from the canonical ensemble (*Feynman*). The classical partition function and its components are given by the following equations:

$$Z = \int d\mathbf{q}d\mathbf{p} \cdot \exp[-\beta H_{eff}]$$

$$H_{eff} = \left( \sum_{i=1}^P \frac{p_i^2}{2m_i} \right) + V_{eff}$$

$$V_{eff} = \sum_{i=1}^P \left[ \frac{mP}{2\beta^2 \hbar^2} (q_i - q_{i+1})^2 + \frac{1}{P} V(q_i) \right]$$

The interested user is referred to any of the numerous references on this methodology, but briefly, each quantum particle in a path integral simulation is represented by a ring-polymer of P quasi-beads, labeled from 1 to P. During the simulation, each quasi-bead interacts with beads on the other ring-polymers with the same imaginary time index (the second term in the effective potential above). The quasi-beads also interact with the two neighboring quasi-beads through the spring potential in imaginary-time space (first term in effective potential). To sample the canonical ensemble, a Nose-Hoover massive chain thermostat is applied (*Tuckerman*). With the massive chain algorithm, a chain of NH thermostats



is coupled to each degree of freedom for each quasi-bead. The keyword *temp* sets the target temperature for the system and the keyword *nhc* sets the number  $N_c$  of thermostats in each chain. For example, for a simulation of  $N$  particles with  $P$  beads in each ring-polymer, the total number of NH thermostats would be  $3 \times N \times P \times N_c$ .

---

**Note:** This fix implements a complete velocity-verlet integrator combined with NH massive chain thermostat, so no other time integration fix should be used.

---

The *method* keyword determines what style of PIMD is performed. A value of *pimd* is standard PIMD. A value of *nmpimd* is for normal-mode PIMD. A value of *cmd* is for centroid molecular dynamics (CMD). The difference between the styles is as follows.

In standard PIMD, the value used for a bead's fictitious mass is arbitrary. A common choice is to use  $M_i = m/P$ , which results in the mass of the entire ring-polymer being equal to the real quantum particle. But it can be difficult to efficiently integrate the equations of motion for the stiff harmonic interactions in the ring polymers.

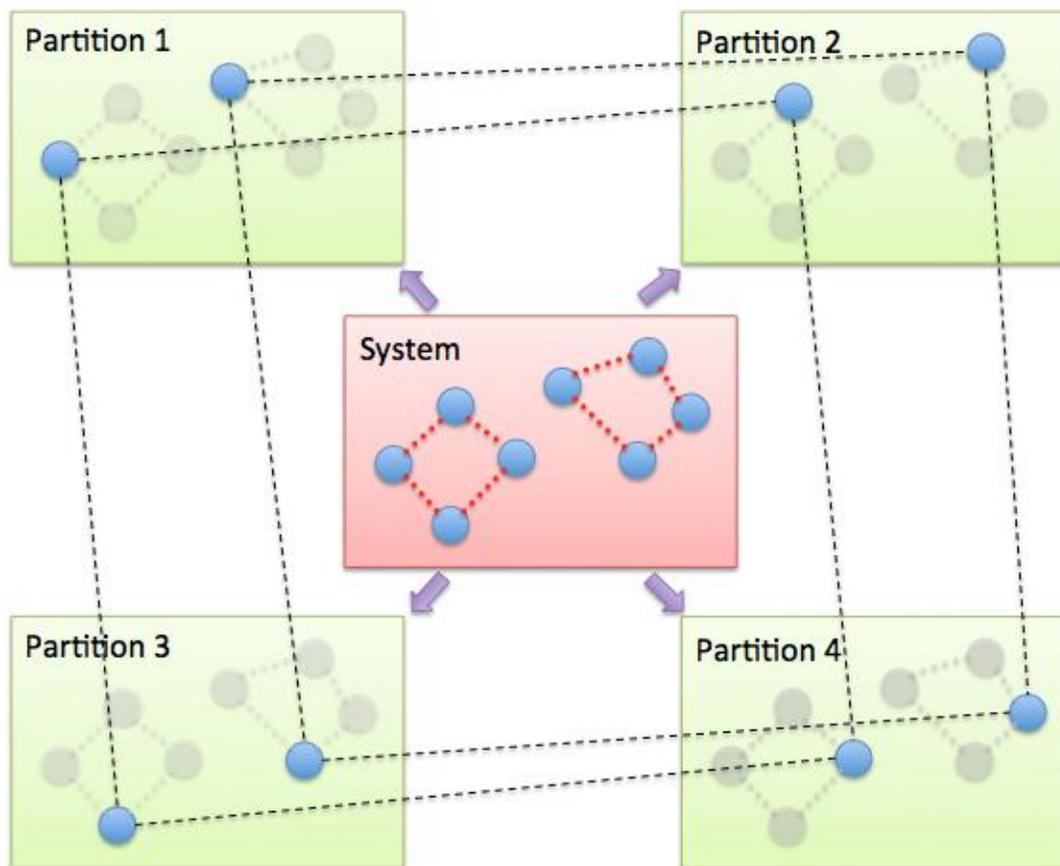
A useful way to resolve this issue is to integrate the equations of motion in a normal mode representation, using Normal Mode Path-Integral Molecular Dynamics (NMPIMD) (*Caol*). In NMPIMD, the NH chains are attached to each normal mode of the ring-polymer and the fictitious mass of each mode is chosen as  $M_k =$  the eigenvalue of the  $K$ th normal mode for  $k > 0$ . The  $k = 0$  mode, referred to as the zero-frequency mode or centroid, corresponds to overall translation of the ring-polymer and is assigned the mass of the real particle.

Motion of the centroid can be effectively uncoupled from the other normal modes by scaling the fictitious masses to achieve a partial adiabatic separation. This is called a Centroid Molecular Dynamics (CMD) approximation (*Caol2*). The time-evolution (and resulting dynamics) of the quantum particles can be used to obtain centroid time correlation functions, which can be further used to obtain the true quantum correlation function for the original system. The CMD method also uses normal modes to evolve the system, except only the  $k > 0$  modes are thermostatted, not the centroid degrees of freedom.

The keyword *fmass* sets a further scaling factor for the fictitious masses of beads, which can be used for the Partial Adiabatic CMD (*Hone*), or to be set as  $P$ , which results in the fictitious masses to be equal to the real particle masses.

The keyword *sp* is a scaling factor on Planck's constant, which can be useful for debugging or other purposes. The default value of 1.0 is appropriate for most situations.

The PIMD algorithm in LAMMPS is implemented as a hyper-parallel scheme as described in (*Calhoun*). In LAMMPS this is done by using *multi-replica feature* in LAMMPS, where each quasi-particle system is stored and simulated on a separate partition of processors. The following diagram illustrates this approach. The original system with 2 ring polymers is shown in red. Since each ring has 4 quasi-beads (imaginary time slices), there are 4 replicas of the system, each running on one of the 4 partitions of processors. Each replica (shown in green) owns one quasi-bead in each ring.



To run a PIMD simulation with  $M$  quasi-beads in each ring polymer using  $N$  MPI tasks for each partition's domain-decomposition, you would use  $P = M \times N$  processors (cores) and run the simulation as follows:

```
mpirun -np P lmp_mpi -partition MxN -in script
```

Note that in the LAMMPS input script for a multi-partition simulation, it is often very useful to define a *uloop-style variable* such as

```
variable ibead uloop M pad
```

where  $M$  is the number of quasi-beads (partitions) used in the calculation. The `uloop` variable can then be used to manage I/O related tasks for each of the partitions, e.g.

```
dump dcd all dcd 10 system_${ibead}.dcd
restart 1000 system_${ibead}.restart1 system_${ibead}.restart2
read_restart system_${ibead}.restart2
```

## 17.147.4 Restrictions

This fix is part of the REPLICIA package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

A PIMD simulation can be initialized with a single data file read via the [read\\_data](#) command. However, this means all quasi-beads in a ring polymer will have identical positions and velocities, resulting in identical trajectories for all quasi-beads. To avoid this, users can simply initialize velocities with different random number seeds assigned to each partition, as defined by the uloop variable, e.g.

```
velocity all create 300.0 1234${ibead} rot yes dist gaussian
```

## 17.147.5 Default

The keyword defaults are method = pimd, fmass = 1.0, sp = 1.0, temp = 300.0, and nhc = 2.

**(Feynman)** R. Feynman and A. Hibbs, Chapter 7, Quantum Mechanics and Path Integrals, McGraw-Hill, New York (1965).

**(Tuckerman)** M. Tuckerman and B. Berne, J Chem Phys, 99, 2796 (1993).

**(Cao1)** J. Cao and B. Berne, J Chem Phys, 99, 2902 (1993).

**(Cao2)** J. Cao and G. Voth, J Chem Phys, 100, 5093 (1994).

**(Hone)** T. Hone, P. Rossky, G. Voth, J Chem Phys, 124, 154103 (2006).

**(Calhoun)** A. Calhoun, M. Pavese, G. Voth, Chem Phys Letters, 262, 415 (1996).

## 17.148 fix plane force command

### 17.148.1 Syntax

```
fix ID group-ID plane force x y z
```

- ID, group-ID are documented in [fix](#) command
- plane force = style name of this fix command
- x y z = 3-vector that is normal to the plane

### 17.148.2 Examples

```
fix hold boundary plane force 1.0 0.0 0.0
```

### 17.148.3 Description

Adjust the forces on each atom in the group so that only the components of force in the plane specified by the normal vector (x,y,z) remain. This is done by subtracting out the component of force perpendicular to the plane.

If the initial velocity of the atom is 0.0 (or in the plane), then it should continue to move in the plane thereafter.

### 17.148.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

### 17.148.5 Restrictions

none

### 17.148.6 Related commands

*fix lineforce*

### 17.148.7 Default

none

## 17.149 fix plumed command

### 17.149.1 Syntax

```
fix ID group-ID plumed keyword value ...
```

- ID, group-ID are documented in *fix* command
- plumed = style name of this fix command
- keyword = *plumedfile* or *outfile*

*plumedfile* arg = name of PLUMED input file to use (default: NULL)

*outfile* arg = name of file on which to write the PLUMED log (default: NULL)

## 17.149.2 Examples

```
fix pl all plumed all plumed plumedfile plumed.dat outfile p.log
```

## 17.149.3 Description

This fix instructs LAMMPS to call the **PLUMED** library, which allows one to perform various forms of trajectory analysis on the fly and to also use methods such as umbrella sampling and metadynamics to enhance the sampling of phase space.

The documentation included here only describes the fix plumed command itself. This command is LAMMPS specific, whereas most of the functionality implemented in PLUMED will work with a range of MD codes, and when PLUMED is used as a stand alone code for analysis. The full [documentation for PLUMED](#) is available online and included in the PLUMED source code. The PLUMED library development is hosted at <https://github.com/plumed/plumed2> A detailed discussion of the code can be found in (*Tribello*).

There is an example input for using this package with LAMMPS in the examples/PACKAGES/plumed directory.

The command to make LAMMPS call PLUMED during a run requires two keyword value pairs pointing to the PLUMED input file and an output file for the PLUMED log. The user must specify these arguments every time PLUMED is to be used. Furthermore, the fix plumed command should appear in the LAMMPS input file **after** relevant input parameters (e.g. the timestep) have been set.

The *group-ID* entry is ignored. LAMMPS will always pass all the atoms to PLUMED and there can only be one instance of the plumed fix at a time. The way the plumed fix is implemented ensures that the minimum amount of information required is communicated. Furthermore, PLUMED supports multiple, completely independent collective variables, multiple independent biases and multiple independent forms of analysis. There is thus really no restriction in functionality by only allowing only one plumed fix in the LAMMPS input.

The *plumedfile* keyword allows the user to specify the name of the PLUMED input file. Instructions as to what should be included in a plumed input file can be found in the [documentation for PLUMED](#)

The *outfile* keyword allows the user to specify the name of a file in which to output the PLUMED log. This log file normally just repeats the information that is contained in the input file to confirm it was correctly read and parsed. The names of the files in which the results are stored from the various analysis options performed by PLUMED will be specified by the user in the PLUMED input file.

## 17.149.4 Restart, fix\_modify, output, run start/stop, minimize info

When performing a restart of a calculation that involves PLUMED you must include a RESTART command in the PLUMED input file as detailed in the [PLUMED documentation](#). When the restart command is found in the PLUMED input PLUMED will append to the files that were generated in the run that was performed previously. No part of the PLUMED restart data is included in the LAMMPS restart files. Furthermore, any history dependent bias potentials that were accumulated in previous calculations will be read in when the RESTART command is included in the PLUMED input.

The *fix\_modify energy* option is supported by this fix to add the energy change from the biasing force added by PLUMED to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy yes*.

The *fix\_modify virial* option is supported by this fix to add the contribution from the biasing force to the global pressure of the system via the *compute pressure* command. This can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial yes*.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the PLUMED energy mentioned above. The scalar value calculated by this fix is “extensive”.

Note that other quantities of interest can be output by commands that are native to PLUMED.

### 17.149.5 Restrictions

This fix is part of the PLUMED package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

There can only be one fix plumed command active at a time.

### 17.149.6 Related commands

*fix smd fix colvars*

### 17.149.7 Default

The default options are plumedfile = NULL and outfile = NULL

---

(Tribello) G.A. Tribello, M. Bonomi, D. Branduardi, C. Camilloni and G. Bussi, *Comp. Phys. Comm* 185, 604 (2014)

## 17.150 fix poems command

### 17.150.1 Syntax

```
fix ID group-ID poems keyword values
```

- ID, group-ID are documented in *fix* command
- poems = style name of this fix command
- keyword = *group* or *file* or *molecule*

*group* values = list of group IDs

*molecule* values = none

*file* values = filename

### 17.150.2 Examples

```
fix 3 fluid poems group clump1 clump2 clump3
fix 3 fluid poems file cluster.list
```

### 17.150.3 Description

Treats one or more sets of atoms as coupled rigid bodies. This means that each timestep the total force and torque on each rigid body is computed and the coordinates and velocities of the atoms are updated so that the collection of bodies move as a coupled set. This can be useful for treating a large biomolecule as a collection of connected, coarse-grained particles.

The coupling, associated motion constraints, and time integration is performed by the software package *Parallelizable Open source Efficient Multibody Software (POEMS)* which computes the constrained rigid-body motion of articulated (jointed) multibody systems (*Anderson*). POEMS was written and is distributed by Prof Kurt Anderson, his graduate student Rudranarayan Mukherjee, and other members of his group at Rensselaer Polytechnic Institute (RPI). Rudranarayan developed the LAMMPS/POEMS interface. For copyright information on POEMS and other details, please refer to the documents in the poems directory distributed with LAMMPS.

This fix updates the positions and velocities of the rigid atoms with a constant-energy time integration, so you should not update the same atoms via other fixes (e.g. nve, nvt, npt, temp/rescale, langevin).

Each body must have a non-degenerate inertia tensor, which means it must contain at least 3 non-collinear atoms. Which atoms are in which bodies can be defined via several options.

For option *group*, each of the listed groups is treated as a rigid body. Note that only atoms that are also in the fix group are included in each rigid body.

For option *molecule*, each set of atoms in the group with a different molecule ID is treated as a rigid body.

For option *file*, sets of atoms are read from the specified file and each set is treated as a rigid body. Each line of the file specifies a rigid body in the following format:

```
ID type atom1-ID atom2-ID atom3-ID ...
```

ID as an integer from 1 to M (the number of rigid bodies). Type is any integer; it is not used by the fix poems command. The remaining arguments are IDs of atoms in the rigid body, each typically from 1 to N (the number of atoms in the system). Only atoms that are also in the fix group are included in each rigid body. Blank lines and lines that begin with '#' are skipped.

A connection between a pair of rigid bodies is inferred if one atom is common to both bodies. The POEMS solver treats that atom as a spherical joint with 3 degrees of freedom. Currently, a collection of bodies can only be connected by joints as a linear chain. The entire collection of rigid bodies can represent one or more chains. Other connection topologies (tree, ring) are not allowed, but will be added later. Note that if no joints exist, it is more efficient to use the *fix rigid* command to simulate the system.

When the poems fix is defined, it will print out statistics on the total # of clusters, bodies, joints, atoms involved. A cluster in this context means a set of rigid bodies connected by joints.

For computational efficiency, you should turn off pairwise and bond interactions within each rigid body, as they no longer contribute to the motion. The "neigh\_modify exclude" and "delete\_bonds" commands can be used to do this if each rigid body is a group.

For computational efficiency, you should only define one fix poems which includes all the desired rigid bodies. LAMMPS will allow multiple poems fixes to be defined, but it is more expensive.

The degrees-of-freedom removed by coupled rigid bodies are accounted for in temperature and pressure computations. Similarly, the rigid body contribution to the pressure virial is also accounted for. The latter is only correct if forces within the bodies have been turned off, and there is only a single fix poems defined.

### 17.150.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify virial* option is supported by this fix to add the contribution due to the added forces and torques on atoms to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial yes*.

The *fix\_modify bodyforces* option is supported by this fix style to set whether per-body forces and torques are computed early or late in a timestep, i.e. at the post-force stage or at the final-integrate stage, respectively.

No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.150.5 Restrictions

This fix is part of the *POEMS* package. It is only enabled if LAMMPS was built with that package, which also requires the *POEMS* library be built and linked with LAMMPS. See the *Build package* page for more info.

### 17.150.6 Related commands

*fix rigid*, *delete\_bonds*, *neigh\_modify* exclude

### 17.150.7 Default

none

---

(Anderson) Anderson, Mukherjee, Critchley, Ziegler, and Lipton “POEMS: Parallelizable Open-source Efficient Multibody Software “, *Engineering With Computers* (2006). ([link to paper](#))

## 17.151 fix polarize/bem/gmres command

## 17.152 fix polarize/bem/icc command

## 17.153 fix polarize/functional command

### 17.153.1 Syntax

```
fix ID group-ID style nevery tolerance ...
```

- ID, group-ID are documented in *fix* command
- style = *polarize/bem/gmres* or *polarize/bem/icc* or *polarize/functional*
- Nevery = this fixed is invoked every this many timesteps
- tolerance = the relative tolerance for the iterative solver to stop



## 17.153.2 Examples

```
fix 2 interface polarize/bem/gmres 5 0.0001
fix 1 interface polarize/bem/icc 1 0.0001
fix 3 interface polarize/functional 1 0.001
```

Used in input scripts:

```
examples/PACKAGES/dielectric/in.confined
examples/PACKAGES/dielectric/in.nopbc
```

## 17.153.3 Description

These fixes compute induced charges at the interface between two impermeable media with different dielectric constants. The interfaces need to be discretized into vertices, each representing a boundary element. The vertices are treated as if they were regular atoms or particles. *atom\_style dielectric* should be used since it defines the additional properties of each interface particle such as interface normal vectors, element areas, and local dielectric mismatch. These fixes also require the use of *pair\_style* and *kpace\_style* with the *dielectric* suffix. At every time step, given a configuration of the physical charges in the system (such as atoms and charged particles) these fixes compute and update the charge of the interface particles. The interfaces are allowed to move during the simulation with appropriate time integrators (for example, with *fix\_rigid*).

Consider an interface between two media: one with dielectric constant of 78 (water), the other of 4 (silica). The interface is discretized into 2000 boundary elements, each represented by an interface particle. Suppose that each interface particle has a normal unit vector pointing from the silica medium to water. The dielectric difference along the normal vector is then  $78 - 4 = 74$ , the mean dielectric value is  $(78 + 4) / 2 = 41$ . Each boundary element also has its area and the local mean curvature (which is used by these fixes for computing a correction term in the local electric field). To model charged interfaces, the interface particle will have a non-zero charge value, coming from its area and surface charge density.

For non-interface particles such as atoms and charged particles, the interface normal vectors, element area, and dielectric mismatch are irrelevant. Their local dielectric value is used to rescale their actual charge when computing the Coulombic interactions. For instance, for a cation carrying a charge of +2 (in charge unit) in an implicit solvent with dielectric constant of 40 would have actual charge of +2, and a local dielectric constant value of 40. It is assumed that the particles cannot pass through the interface during the simulation so that its local dielectric constant value does not change.

There are some example scripts for using these fixes with LAMMPS in the `examples/PACKAGES/dielectric` directory. The README file therein contains specific details on the system setup. Note that the example data files show the additional fields (columns) needed for *atom\_style dielectric* beyond the conventional fields *id*, *mol*, *type*, *q*, *x*, *y*, and *z*.

For fix *polarize/bem/gmres* and fix *polarize/bem/icc* the induced charges of the atoms in the specified group, which are the vertices on the interface, are computed using the equation:

$$\sigma_b(\mathbf{s}) = \frac{1 - \bar{\epsilon}}{\bar{\epsilon}} \sigma_f(\mathbf{s}) - \epsilon_0 \frac{\Delta\epsilon}{\bar{\epsilon}} \mathbf{E}(\mathbf{s}) \cdot \mathbf{n}(\mathbf{s})$$

- $\sigma_b$  is the induced charge density at the interface vertex  $\mathbf{s}$ .
- $\bar{\epsilon}$  is the mean dielectric constant at the interface vertex:  $\bar{\epsilon} = (\epsilon_1 + \epsilon_2)/2$ .
- $\Delta\epsilon$  is the dielectric constant difference at the interface vertex:  $\Delta\epsilon = \epsilon_1 - \epsilon_2$
- $\sigma_f$  is the free charge density at the interface vertex
- $\mathbf{E}(\mathbf{s})$  is the electrical field at the vertex

- $\mathbf{n}(s)$  is the unit normal vector at the vertex pointing from medium with  $\epsilon_2$  to that with  $\epsilon_1$

Fix *polarize/bem/gmres* employs the Generalized Minimum Residual (GMRES) as described in (*Barros*) to solve  $\sigma_b$ .

Fix *polarize/bem/icc* employs the successive over-relaxation algorithm as described in (*Tyagi*) to solve  $\sigma_b$ .

The iterative solvers would terminate either when the maximum relative change in the induced charges in consecutive iterations is below the set tolerance, or when the number of iterations reaches *iter\_max* (see below).

Fix *polarize/functional* employs the energy functional variation approach as described in (*Jadhao*) to solve  $\sigma_b$ .

More details on the implementation of these fixes and their recommended use are described in (*NguyenTD*).

#### 17.153.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify* command provides certain options to control the induced charge solver and the initial values of the interface elements:

```
itr_max arg
  arg = maximum number of iterations for convergence
dielectrics ediff emean epsilon area charge
  ediff = dielectric difference
  emean = dielectric mean
  epsilon = local dielectric value
  area = element area
  charge = real interface charge
```

*polarize/bem/gmres* or *polarize/bem/icc* compute a global 2-element vector which can be accessed by various *output commands*. The first element is the number of iterations when the solver terminates (of which the upper bound is set by *iter\_max*). The second element is the RMS error.

#### 17.153.5 Restrictions

These fixes are part of the DIELECTRIC package. It is only enabled if LAMMPS was built with that package, which requires that also the KSPACE package is installed. See the *Build package* page for more info.

Note that the *polarize/bem/gmres* and *polarize/bem/icc* fixes only support *units lj, real, metal, si* and *nano* at the moment.

#### 17.153.6 Related commands

*pair\_coeff*, *fix polarize*, *read\_data*, *pair\_style lj/cut/coul/long/dielectric*, *kpace\_style ppm/dielectric*, *compute efield/atom*

## 17.153.7 Default

`iter_max = 20`

(**Barros**) Barros, Sinkovits, Luijten, J. Chem. Phys, 140, 064903 (2014)

(**Tyagi**) Tyagi, Suzen, Sega, Barbosa, Kantorovich, Holm, J Chem Phys, 132, 154112 (2010)

(**Jadhao**) Jadhao, Solis, Olvera de la Cruz, J Chem Phys, 138, 054119 (2013)

(**NguyenTD**) Nguyen, Li, Bagchi, Solis, Olvera de la Cruz, Comput Phys Commun 241, 80-19 (2019)

## 17.154 fix pour command

### 17.154.1 Syntax

```
fix ID group-ID pour N type seed keyword values ...
```

- ID, group-ID are documented in *fix* command
- pour = style name of this fix command
- N = # of particles to insert
- type = atom type to assign to inserted particles (offset for molecule insertion)
- seed = random # seed (positive integer)
- one or more keyword/value pairs may be appended to args
- keyword = *region* or *diam* or *vol* or *rate* or *dens* or *vel* or *mol* or *rigid* or *shake* or *ignore*

*region* value = region-ID

region-ID = ID of region to use as insertion volume

*diam* values = dstyle args

dstyle = *one* or *range* or *poly*

*one* args = D

D = single diameter for inserted particles (distance units)

*range* args = Dlo Dhi

Dlo,Dhi = range of diameters for inserted particles (distance units)

*poly* args = Npoly D1 P1 D2 P2 ...

Npoly = # of (D,P) pairs

D1,D2,... = diameter for subset of inserted particles (distance units)

P1,P2,... = percentage of inserted particles with this diameter (0-1)

*id* values = idflag

idflag = *max* or *next* = how to choose IDs for inserted particles and molecules

*vol* values = fraction Nattempt

fraction = desired volume fraction for filling insertion volume

Nattempt = max # of insertion attempts per particle

*rate* value = V

V = z velocity (3d) or y velocity (2d) at which

insertion volume moves (velocity units)

*dens* values = Rholo Rhohi

Rholo,Rhohi = range of densities for inserted particles (mass/volume units)

*vel* values (3d) = vxlo vxhi vylo vyhi vz

*vel* values (2d) = vxlo vxhi vy

`vxlo,vxhi` = range of x velocities for inserted particles (velocity units)  
`vylo,vyhi` = range of y velocities for inserted particles (velocity units)  
`vz` = z velocity (3d) assigned to inserted particles (velocity units)  
`vy` = y velocity (2d) assigned to inserted particles (velocity units)  
`mol` value = template-ID  
    template-ID = ID of molecule template specified in a separate `molecule` command  
`molfrac` values = f1 f2 ... fN  
    f1 to fN = relative probability of creating each of N molecules in template-ID  
`rigid` value = fix-ID  
    fix-ID = ID of `fix rigid/small` command  
`shake` value = fix-ID  
    fix-ID = ID of `fix shake` command  
`ignore` value = none  
    skip any line or triangle particles when detecting possible overlaps with inserted particles

## 17.154.2 Examples

```
fix 3 all pour 1000 2 29494 region myblock
fix 2 all pour 10000 1 19985583 region disk vol 0.33 100 rate 1.0 diam range 0.9 1.1
fix 2 all pour 10000 1 19985583 region disk diam poly 2 0.7 0.4 1.5 0.6
fix ins all pour 500 1 4767548 vol 0.8 10 region slab mol object rigid myRigid
```

## 17.154.3 Description

Insert finite-size particles or molecules into the simulation box every few timesteps within a specified region until N particles or molecules have been inserted. This is typically used to model the pouring of granular particles into a container under the influence of gravity. For the remainder of this doc page, a single inserted atom or molecule is referred to as a “particle”.

If inserted particles are individual atoms, they are assigned the specified atom type. If they are molecules, the type of each atom in the inserted molecule is specified in the file read by the `molecule` command, and those values are added to the specified atom type. E.g. if the file specifies atom types 1,2,3, and those are the atom types you want for inserted molecules, then specify `type = 0`. If you specify `type = 2`, the in the inserted molecule will have atom types 3,4,5.

All atoms in the inserted particle are assigned to two groups: the default group “all” and the group specified in the fix pour command (which can also be “all”).

This command must use the `region` keyword to define an insertion volume. The specified region must have been previously defined with a `region` command. It must be of type `block` or a z-axis `cylinder` and must be defined with side = `in`. The cylinder style of region can only be used with 3d simulations.

Individual atoms are inserted, unless the `mol` keyword is used. It specifies a `template-ID` previously defined using the `molecule` command, which reads a file that defines the molecule. The coordinates, atom types, center-of-mass, moments of inertia, etc, as well as any bond/angle/etc and special neighbor information for the molecule can be specified in the molecule file. See the `molecule` command for details. The only settings required to be in this file are the coordinates and types of atoms in the molecule.

If the molecule template contains more than one molecule, the relative probability of depositing each molecule can be specified by the `molfrac` keyword. N relative probabilities, each from 0.0 to 1.0, are specified, where N is the number of molecules in the template. Each time a molecule is inserted, a random number is used to sample from the list of relative probabilities. The N values must sum to 1.0.

If you wish to insert molecules via the `mol` keyword, that will be treated as rigid bodies, use the `rigid` keyword, specifying as its value the ID of a separate `fix rigid/small` command which also appears in your input script.

---

**Note:** If you wish the new rigid molecules (and other rigid molecules) to be thermostatted correctly via *fix rigid/small/nvt* or *fix rigid/small/npt*, then you need to use the “fix\_modify dynamic/dof yes” command for the rigid fix. This is to inform that fix that the molecule count will vary dynamically.

---

If you wish to insert molecules via the *mol* keyword, that will have their bonds or angles constrained via SHAKE, use the *shake* keyword, specifying as its value the ID of a separate *fix shake* command which also appears in your input script.

Each timestep particles are inserted, they are placed randomly inside the insertion volume so as to mimic a stream of poured particles. If they are molecules they are also oriented randomly. Each atom in the particle is tested for overlaps with existing particles, including effects due to periodic boundary conditions if applicable. If an overlap is detected, another random insertion attempt is made; see the *vol* keyword discussion below. The larger the volume of the insertion region, the more particles that can be inserted at any one timestep. Particles are inserted again after enough time has elapsed that the previously inserted particles fall out of the insertion volume under the influence of gravity. Insertions continue every so many timesteps until the desired # of particles has been inserted.

---

**Note:** If you are monitoring the temperature of a system where the particle count is changing due to adding particles, you typically should use the *compute\_modify dynamic yes* command for the temperature compute you are using.

---

All other keywords are optional with defaults as shown below.

The *diam* option is only used when inserting atoms and specifies the diameters of inserted particles. There are 3 styles: *one*, *range*, or *poly*. For *one*, all particles will have diameter *D*. For *range*, the diameter of each particle will be chosen randomly and uniformly between the specified *Dlo* and *Dhi* bounds. For *poly*, a series of *Npoly* diameters is specified. For each diameter a percentage value from 0.0 to 1.0 is also specified. The *Npoly* percentages must sum to 1.0. For the example shown above with “diam 2 0.7 0.4 1.5 0.6”, all inserted particles will have a diameter of 0.7 or 1.5. 40% of the particles will be small; 60% will be large.

Note that for molecule insertion, the diameters of individual atoms in the molecule can be specified in the file read by the *molecule* command. If not specified, the diameter of each atom in the molecule has a default diameter of 1.0.

The *id* option has two settings which are used to determine the atom or molecule IDs to assign to inserted particles/molecules. In both cases a check is done of the current system to find the maximum current atom and molecule ID of any existing particle. Newly inserted particles and molecules are assigned IDs that increment those max values. For the *max* setting, which is the default, this check is done at every insertion step, which allows for particles to leave the system, and their IDs to potentially be re-used. For the *next* setting this check is done only once when the fix is specified, which can be more efficient if you are sure particles will not be added in some other way.

The *vol* option specifies what volume fraction of the insertion volume will be filled with particles. For particles with a size specified by the *diam range* keyword, they are assumed to all be of maximum diameter *Dhi* for purposes of computing their contribution to the volume fraction.

The higher the volume fraction value, the more particles are inserted each timestep. Since inserted particles cannot overlap, the maximum volume fraction should be no higher than about 0.6. Each timestep particles are inserted, LAMMPS will make up to a total of *M* tries to insert the new particles without overlaps, where  $M = \# \text{ of inserted particles} * N_{\text{attempt}}$ . If LAMMPS is unsuccessful at completing all insertions, it prints a warning.

The *dens* and *vel* options enable inserted particles to have a range of densities or xy velocities. The specific values for a particular inserted particle will be chosen randomly and uniformly between the specified bounds. Internally, the density value for a particle is converted to a mass, based on the radius (volume) of the particle. The *vz* or *vy* value for option *vel* assigns a z-velocity (3d) or y-velocity (2d) to each inserted particle.

The *rate* option moves the insertion volume in the z direction (3d) or y direction (2d). This enables pouring particles from a successively higher height over time.

The *ignore* option is useful when running a simulation that used line segment (2d) or triangle (3d) particles, typically to define boundaries for spherical granular particles to interact with. See the *atom\_style line or tri* command for details. Lines and triangles store their size, and if the size is large it may overlap (in a spherical sense) with the insertion region, even if the line/triangle is oriented such that there is no actual overlap. This can prevent particles from being inserted. The *ignore* keyword causes the overlap check to skip any line or triangle particles. Obviously you should only use it if there is in fact no overlap of the line or triangle particles with the insertion region.

---

#### 17.154.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. This means you must be careful when restarting a pouring simulation, when the restart file was written in the middle of the pouring operation. Specifically, you should use a new *fix pour* command in the input script for the restarted simulation that continues the operation. You will need to adjust the arguments of the original *fix pour* command to do this.

Also note that because the state of the random number generator is not saved in restart files, you cannot do “exact” restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior if you adjust the *fix pour* parameters appropriately.

None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.154.5 Restrictions

This fix is part of the GRANULAR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

For 3d simulations, a gravity fix in the -z direction must be defined for use in conjunction with this fix. For 2d simulations, gravity must be defined in the -y direction.

The specified insertion region cannot be a “dynamic” region, as defined by the *region* command.

#### 17.154.6 Related commands

*fix deposit*, *fix gravity*, *region*

#### 17.154.7 Default

Insertions are performed for individual particles, i.e. no *mol* setting is defined. If the *mol* keyword is used, the default for *molfrac* is an equal probabilities for all molecules in the template. Additional option defaults are *diam* = one 1.0, *dens* = 1.0 1.0, *vol* = 0.25 50, *rate* = 0.0, *vel* = 0.0 0.0 0.0 0.0 0.0 (for 3d), *vel* = 0.0 0.0 0.0 (for 2d), and *id* = max.

## 17.155 fix precession/spin command

### 17.155.1 Syntax

```
fix ID group precession/spin style args
```

- ID, group are documented in *fix* command
- precession/spin = style name of this fix command
- style = *zeeman* or *anisotropy* or *cubic* or *stt*

*zeeman* args = H x y z

H = intensity of the magnetic field (in Tesla)

x y z = vector direction of the field

*anisotropy* args = K x y z

K = intensity of the magnetic anisotropy (in eV)

x y z = vector direction of the anisotropy

*cubic* args = K1 K2c n1x n1y n1z n2x n2y n2z n3x n3y n3z

K1 and K2c = intensity of the magnetic anisotropy (in eV)

n1x to n3z = three direction vectors of the cubic anisotropy

*stt* args = J x y z

J = intensity of the spin-transfer torque field

x y z = vector direction of the field

### 17.155.2 Examples

```
fix 1 all precession/spin zeeman 0.1 0.0 0.0 1.0
fix 1 3 precession/spin anisotropy 0.001 0.0 0.0 1.0
fix 1 iron precession/spin cubic 0.001 0.0005 1.0 0.0 0.0 0.0 1.0 0.0 0.0 0.0 1.0
fix 1 all precession/spin zeeman 0.1 0.0 0.0 1.0 anisotropy 0.001 0.0 0.0 1.0
```

### 17.155.3 Description

This fix applies a precession torque to each magnetic spin in the group.

Style *zeeman* is used for the simulation of the interaction between the magnetic spins in the defined group and an external magnetic field:

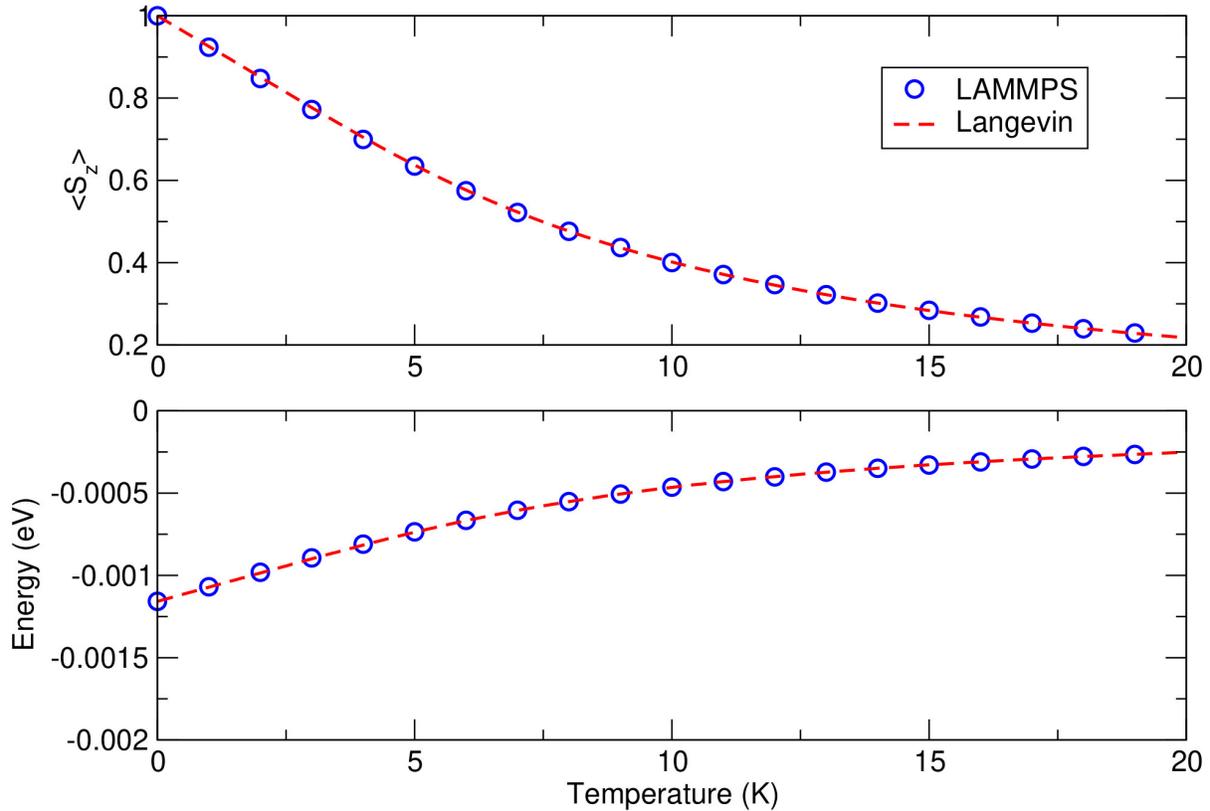
$$H_{Zeeman} = -g \sum_{i=0}^N \mu_i \vec{s}_i \cdot \vec{B}_{ext}$$

with:

- $\vec{B}_{ext}$  the external magnetic field (in T)
- $g$  the Lande factor (hard-coded as  $g = 2.0$ )
- $\vec{s}_i$  the unitary vector describing the orientation of spin  $i$
- $\mu_i$  the atomic moment of spin  $i$  given as a multiple of the Bohr magneton  $\mu_B$  (for example,  $\mu_i \approx 2.2$  in bulk iron).

The field value in Tesla is multiplied by the gyromagnetic ratio,  $g \cdot \mu_B / \hbar$ , converting it into a precession frequency in rad.THz (in metal units and with  $\mu_B = 5.788 \cdot 10^{-5}$  eV/T).

As a comparison, the figure below displays the simulation of a single spin (of norm  $\mu_i = 1.0$ ) submitted to an external magnetic field of  $|B_{ext}| = 10.0$  Tesla (and oriented along the z axis). The upper plot shows the average magnetization along the external magnetic field axis and the lower plot the Zeeman energy, both as a function of temperature. The reference result is provided by the plot of the Langevin function for the same parameters.



The temperature effects are accounted for by connecting the spin  $i$  to a thermal bath using a Langevin thermostat (see [fix langevin/spin](#) for the definition of this thermostat).

Style *anisotropy* is used to simulate an easy axis or an easy plane for the magnetic spins in the defined group:

$$H_{aniso} = - \sum_{i=1}^N K_{an}(\mathbf{r}_i) (\vec{s}_i \cdot \vec{n}_i)^2$$

with  $n$  defining the direction of the anisotropy, and  $K$  (in eV) its intensity. If  $K > 0$ , an easy axis is defined, and if  $K < 0$ , an easy plane is defined.

Style *cubic* is used to simulate a cubic anisotropy, with three possible easy axis for the magnetic spins in the defined group:

$$H_{cubic} = - \sum_{i=1}^N K_1 \left[ (\vec{s}_i \cdot \vec{n}_1)^2 (\vec{s}_i \cdot \vec{n}_2)^2 + (\vec{s}_i \cdot \vec{n}_2)^2 (\vec{s}_i \cdot \vec{n}_3)^2 + (\vec{s}_i \cdot \vec{n}_1)^2 (\vec{s}_i \cdot \vec{n}_3)^2 \right] + K_2^{(c)} (\vec{s}_i \cdot \vec{n}_1)^2 (\vec{s}_i \cdot \vec{n}_2)^2 (\vec{s}_i \cdot \vec{n}_3)^2$$

with  $K_1$  and  $K_{2c}$  (in eV) the intensity coefficients and  $\vec{n}_1$ ,  $\vec{n}_2$  and  $\vec{n}_3$  defining the three anisotropic directions defined by the command (from  $n1x$  to  $n3z$ ). For  $\vec{n}_1 = (100)$ ,  $\vec{n}_2 = (010)$ , and  $\vec{n}_3 = (001)$ ,  $K_1 < 0$  defines an iron type anisotropy (easy axis along the (001)-type cube edges), and  $K_1 > 0$  defines a nickel type anisotropy (easy axis along the (111)-type cube diagonals).  $K_2^c > 0$  also defines easy axis along the (111)-type cube diagonals. See chapter 2 of ([Skomski](#)) for more details on cubic anisotropies.



Style *stt* is used to simulate the interaction between the spins and a spin-transfer torque. See equation (7) of (*Chirac*) for more details about the implemented spin-transfer torque term.

In all cases, the choice of  $(xyz)$  only imposes the vector directions for the forces. Only the direction of the vector is important; its length is ignored (the entered vectors are normalized).

Those styles can be combined within one single command line.

---

**Note:** The norm of all vectors defined with the precession/spin command have to be non-zero. For example, defining “fix 1 all precession/spin zeeman 0.1 0.0 0.0 0.0” would result in an error message. Since those vector components are used to compute the inverse of the field (or anisotropy) vector norm, setting a zero-vector would result in a division by zero.

---

#### 17.155.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the energy associated with the spin precession torque to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the potential energy (in energy units) discussed in the previous paragraph. The scalar value is an “extensive” quantity.

No information about this fix is written to *binary restart files*.

#### 17.155.5 Restrictions

The *precession/spin* style is part of the SPIN package. This style is only enabled if LAMMPS was built with this package, and if the atom\_style “spin” was declared. See the *Build package* page for more info.

#### 17.155.6 Related commands

*atom\_style spin*

#### 17.155.7 Default

none

---

(**Skomski**) Skomski, R. (2008). Simple models of magnetism. Oxford University Press.

(**Chirac**) Chirac, Theophile, et al. Ultrafast antiferromagnetic switching in NiO induced by spin transfer torques. Physical Review B 102.13 (2020): 134415.

## 17.156 fix press/berendsen command

### 17.156.1 Syntax

```
fix ID group-ID press/berendsen keyword value ...
```

- ID, group-ID are documented in *fix* command
- press/berendsen = style name of this fix command

one or more keyword value pairs may be appended

keyword = *iso* or *aniso* or *x* or *y* or *z* or *couple* or *dilate* or *modulus*

*iso* or *aniso* values = Pstart Pstop Pdamp

Pstart,Pstop = scalar external pressure at start/end of run (pressure units)

Pdamp = pressure damping parameter (time units)

*x* or *y* or *z* values = Pstart Pstop Pdamp

Pstart,Pstop = external stress tensor component at start/end of run (pressure\_↪units)

Pdamp = stress damping parameter (time units)

*couple* = *none* or *xyz* or *xy* or *yz* or *xz*

*modulus* value = bulk modulus of system (pressure units)

*dilate* value = *all* or *partial*

### 17.156.2 Examples

```
fix 1 all press/berendsen iso 0.0 0.0 1000.0
fix 2 all press/berendsen aniso 0.0 0.0 1000.0 dilate partial
```

### 17.156.3 Description

Reset the pressure of the system by using a Berendsen barostat (*Berendsen*), which rescales the system volume and (optionally) the atoms coordinates within the simulation box every timestep.

Regardless of what atoms are in the fix group, a global pressure is computed for all atoms. Similarly, when the size of the simulation box is changed, all atoms are re-scaled to new positions, unless the keyword *dilate* is specified with a value of *partial*, in which case only the atoms in the fix group are re-scaled. The latter can be useful for leaving the coordinates of atoms in a solid substrate unchanged and controlling the pressure of a surrounding fluid.

---

**Note:** Unlike the *fix npt* or *fix nph* commands which perform Nose/Hoover barostatting AND time integration, this fix does NOT perform time integration. It only modifies the box size and atom coordinates to effect barostatting. Thus you must use a separate time integration fix, like *fix nve* or *fix nvt* to actually update the positions and velocities of atoms. This fix can be used in conjunction with thermostating fixes to control the temperature, such as *fix nvt* or *fix langevin* or *fix temp/berendsen*.

---

See the *Howto barostat* page for a discussion of different ways to perform barostatting.

---

The barostat is specified using one or more of the *iso*, *aniso*, *x*, *y*, *z*, and *couple* keywords. These keywords give you the ability to specify the 3 diagonal components of an external stress tensor, and to couple various of these components together so that the dimensions they represent are varied together during a constant-pressure simulation. Unlike the

*fix npt* and *fix nph* commands, this fix cannot be used with triclinic (non-orthogonal) simulation boxes to control all 6 components of the general pressure tensor.

The target pressures for each of the 3 diagonal components of the stress tensor can be specified independently via the *x*, *y*, *z*, keywords, which correspond to the 3 simulation box dimensions. For each component, the external pressure or tensor component at each timestep is a ramped value during the run from *Pstart* to *Pstop*. If a target pressure is specified for a component, then the corresponding box dimension will change during a simulation. For example, if the *y* keyword is used, the *y*-box length will change. A box dimension will not change if that component is not specified, although you have the option to change that dimension via the *fix deform* command.

For all barostat keywords, the *Pdamp* parameter determines the time scale on which pressure is relaxed. For example, a value of 10.0 means to relax the pressure in a timespan of (roughly) 10 time units (tau or fs or ps - see the *units* command).

---

**Note:** A Berendsen barostat will not work well for arbitrary values of *Pdamp*. If *Pdamp* is too small, the pressure and volume can fluctuate wildly; if it is too large, the pressure will take a very long time to equilibrate. A good choice for many models is a *Pdamp* of around 1000 timesteps. However, note that *Pdamp* is specified in time units, and that timesteps are NOT the same as time units for most *units* settings.

---



---

**Note:** The relaxation time is actually also a function of the bulk modulus of the system (inverse of isothermal compressibility). The bulk modulus has units of pressure and is the amount of pressure that would need to be applied (isotropically) to reduce the volume of the system by a factor of 2 (assuming the bulk modulus was a constant, independent of density, which it's not). The bulk modulus can be set via the keyword *modulus*. The *Pdamp* parameter is effectively multiplied by the bulk modulus, so if the pressure is relaxing faster than expected or desired, increasing the bulk modulus has the same effect as increasing *Pdamp*. The converse is also true. LAMMPS does not attempt to guess a correct value of the bulk modulus; it just uses 10.0 as a default value which gives reasonable relaxation for a Lennard-Jones liquid, but will be way off for other materials and way too small for solids. Thus you should experiment to find appropriate values of *Pdamp* and/or the *modulus* when using this fix.

---

The *couple* keyword allows two or three of the diagonal components of the pressure tensor to be “coupled” together. The value specified with the keyword determines which are coupled. For example, *xz* means the *Pxx* and *Pzz* components of the stress tensor are coupled. *xyz* means all 3 diagonal components are coupled. Coupling means two things: the instantaneous stress will be computed as an average of the corresponding diagonal components, and the coupled box dimensions will be changed together in lockstep, meaning coupled dimensions will be dilated or contracted by the same percentage every timestep. The *Pstart*, *Pstop*, *Pdamp* parameters for any coupled dimensions must be identical. *Couple xyz* can be used for a 2d simulation; the *z* dimension is simply ignored.

---

The *iso* and *aniso* keywords are simply shortcuts that are equivalent to specifying several other keywords together.

The keyword *iso* means couple all 3 diagonal components together when pressure is computed (hydrostatic pressure), and dilate/contract the dimensions together. Using “*iso Pstart Pstop Pdamp*” is the same as specifying these 4 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
couple xyz
```

The keyword *aniso* means *x*, *y*, and *z* dimensions are controlled independently using the *Pxx*, *Pyy*, and *Pzz* components of the stress tensor as the driving forces, and the specified scalar external pressure. Using “*aniso Pstart Pstop Pdamp*” is the same as specifying these 4 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
couple none
```

This fix computes a temperature and pressure each timestep. To do this, the fix creates its own computes of style “temp” and “pressure”, as if these commands had been issued:

```
compute fix-ID_temp group-ID temp
compute fix-ID_press group-ID pressure fix-ID_temp
```

See the *compute temp* and *compute pressure* commands for details. Note that the IDs of the new computes are the fix-ID + underscore + “temp” or fix-ID + underscore + “press”, and the group for the new computes is the same as the fix group.

Note that these are NOT the computes used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp* and *thermo\_press*. This means you can change the attributes of this fix’s temperature or pressure via the *compute\_modify* command or print this temperature or pressure during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* or *thermo\_press* will have no effect on this fix.

#### 17.156.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify temp* and *press* options are supported by this fix. You can use them to assign a *compute* you have defined to this fix which will be used in its temperature and pressure calculations. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

No global or per-atom quantities are stored by this fix for access by various *output commands*.

This fix can ramp its target pressure over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

#### 17.156.5 Restrictions

Any dimension being adjusted by this fix must be periodic.

#### 17.156.6 Related commands

*fix nve, fix nph, fix npt, fix temp/berendsen, fix\_modify*

## 17.156.7 Default

The keyword defaults are dilate = all, modulus = 10.0 in units of pressure for whatever *units* are defined.

(**Berendsen**) Berendsen, Postma, van Gunsteren, DiNola, Haak, J Chem Phys, 81, 3684 (1984).

## 17.157 fix print command

### 17.157.1 Syntax

```
fix ID group-ID print N string keyword value ...
```

- ID, group-ID are documented in *fix* command
- print = style name of this fix command
- N = print every N steps; N can be a variable (see below)
- string = text string to print with optional variable names
- zero or more keyword/value pairs may be appended
- keyword = *file* or *append* or *screen* or *title*

*file* value = filename

*append* value = filename

*screen* value = yes or no

*title* value = string

string = text to print as 1st line of output file

### 17.157.2 Examples

```
fix extra all print 100 "Coords of marker atom = $x $y $z"
fix extra all print 100 "Coords of marker atom = $x $y $z" file coord.txt
```

### 17.157.3 Description

Print a text string every N steps during a simulation run. This can be used for diagnostic purposes or as a debugging tool to monitor some quantity during a run. The text string must be a single argument, so it should be enclosed in single or double quotes if it is more than one word. If it contains variables it must be enclosed in double quotes to insure they are not evaluated when the input script line is read, but will instead be evaluated each time the string is printed.

**Note:** As discussed on the *Commands parse* doc page, the text string can use “immediate” variables, specified as \$(formula) with parenthesis, where the numeric formula has the same syntax as equal-style variables described on the *variable* doc page. This is a convenient way to evaluate a formula immediately without using the variable command to define a named variable and then use that variable in the text string. The formula can include a trailing colon and format string which determines the precision with which the numeric value is output. This is also explained on the *Commands parse* doc page.

Instead of a numeric value, N can be specified as an *equal-style variable*, which should be specified as v\_name, where name is the variable name. In this case, the variable is evaluated at the beginning of a run to determine the **next** timestep at which the string will be written out. On that timestep, the variable will be evaluated again to determine the next timestep, etc. Thus the variable should return timestep values. See the stagger() and logfreq() and stride() math functions for *equal-style variables*, as examples of useful functions to use in this context. For example, the following commands will print output at timesteps 10,20,30,100,200,300,1000,2000,etc:

```
variable      s equal logfreq(10,3,10)
fix extra all print v_s "Coords of marker atom = $x $y $z"
```

The specified group-ID is ignored by this fix.

See the *variable* command for a description of *equal* style variables which are the most useful ones to use with the fix print command, since they are evaluated afresh each timestep that the fix print line is output. Equal-style variables calculate formulas involving mathematical operations, atom properties, group properties, thermodynamic properties, global values calculated by a *compute* or *fix*, or references to other *variables*.

If the *file* or *append* keyword is used, a filename is specified to which the output generated by this fix will be written. If *file* is used, then the filename is overwritten if it already exists. If *append* is used, then the filename is appended to if it already exists, or created if it does not exist.

If the *screen* keyword is used, output by this fix to the screen and logfile can be turned on or off as desired.

The *title* keyword allow specification of the string that will be printed as the first line of the output file, assuming the *file* keyword was used. By default, the title line is as follows:

```
# Fix print output for fix ID
```

where ID is replaced with the fix-ID.

#### 17.157.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.157.5 Restrictions

none

#### 17.157.6 Related commands

*variable, print*

### 17.157.7 Default

The option defaults are no file output, screen = yes, and title string as described above.

## 17.158 fix propel/self command

### 17.158.1 Syntax

```
fix ID group-ID propel/self mode magnitude keyword values
```

- ID, group-ID are documented in *fix* command
- propel/self = style name of this fix command
- mode = *dipole* or *velocity* or *quat*
- magnitude = magnitude of self-propulsion force
- zero or one keyword/value pairs may be appended
- keyword = *qvector*

*qvector* value = direction of force in ellipsoid frame  
*sx*, *sy*, *sz* = components of *qvector*

### 17.158.2 Examples

```
fix active all propel/self dipole 40.0
fix active all propel/self velocity 10.0
fix active all propel/self quat 15.7 qvector 1.0 0.0 0.0
```

### 17.158.3 Description

Add a force to each atom in the group due to a self-propulsion force. The force is given by

$$F_i = f_P e_i$$

where  $i$  is the particle the force is being applied to,  $f_P$  is the magnitude of the force, and  $e_i$  is the vector direction of the force. The specification of  $e_i$  is based on which of the three keywords (*dipole* or *velocity* or *quat*) one selects.

For mode *dipole*,  $e_i$  is just equal to the dipole vectors of the atoms in the group. Therefore, if the dipoles are not unit vectors, the  $e_i$  will not be unit vectors.

---

**Note:** If another command changes the magnitude of the dipole, this force will change accordingly (since  $|e_i|$  will change, which is physically equivalent to re-scaling  $f_P$  while keeping  $|e_i|$  constant), and no warning will be provided by LAMMPS. This is almost never what you want, so ensure you are not changing dipole magnitudes with another LAMMPS fix or pair style. Furthermore, self-propulsion forces (almost) always set  $e_i$  to be a unit vector for all times, so it's best to set all the dipole magnitudes to 1.0 unless you have a good reason not to (see the *set* command on how to do this).

---

For mode *velocity*,  $e_i$  points in the direction of the current velocity (a unit-vector). This can be interpreted as a velocity-dependent friction, as proposed by e.g. (*Erdmann*).

For mode *quat*,  $e_i$  points in the direction of a unit vector, oriented in the coordinate frame of the ellipsoidal particles, which defaults to point along the x-direction. This default behavior can be changed by via the *quatvec* keyword.

The optional *quatvec* keyword specifies the direction of self-propulsion via a unit vector ( $s_x, s_y, s_z$ ). The arguments  $s_x$ ,  $s_y$ , and  $s_z$ , are defined within the coordinate frame of the atom's ellipsoid. For instance, for an ellipsoid with long axis along its x-direction, if one wanted the self-propulsion force to also be along this axis, set  $s_x$  equal to 1 and  $s_y$ ,  $s_z$  both equal to zero. This keyword may only be specified for mode *quat*.

---

**Note:** In using keyword *quatvec*, the three arguments  $s_x$ ,  $s_y$ , and  $s_z$  will be automatically normalized to components of a unit vector internally to avoid users having to explicitly do so themselves. Therefore, in mode *quat*, the vectors  $e_i$  will always be of unit length.

---

Along with adding a force contribution, this fix can also contribute to the virial (pressure) of the system, defined as  $f_P \sum_i \langle e_i \cdot r_i \rangle / (dV)$ , where  $r_i$  is the *unwrapped* coordinate of particle i in the case of periodic boundary conditions. See (*Winkler*) for a discussion of this active pressure contribution.

For modes *dipole* and *quat*, this fix is by default included in pressure computations.

For mode *velocity*, this fix is by default not included in pressure computations.

---

**Note:** In contrast to equilibrium systems, pressure of active systems in general depends on the geometry of the container. The active pressure contribution as calculated in this fix is only valid for certain boundary conditions (spherical walls, rectangular walls, or periodic boundary conditions). For other geometries, the pressure must be measured via explicit calculation of the force per unit area on a wall, and so one must not calculate it using this fix. (Use *fix\_modify* as described below to turn off the virial contribution of this fix). Again, see (*Winkler*) for discussion of why this is the case.

Furthermore, when dealing with active systems, the temperature is no longer well defined. Therefore, one should ensure that the *virial* flag is used in the *compute pressure* command (turning off temperature contributions).

---

#### 17.158.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify virial* option is supported by this fix to add the contribution due to the added forces on atoms to the system's virial as part of *thermodynamic output*. The default is *virial yes* for keywords *dipole* and *quat*. The default is *virial no* for keyword *velocity*.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

#### 17.158.5 Restrictions

With keyword *dipole*, this fix only works when the DIPOLE package is enabled. See the *Build package* page for more info.

This fix is part of the BROWNIAN package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.



## 17.158.6 Related commands

*fix efield* , *fix setforce* , *fix addforce*

## 17.158.7 Default

none

**(Erdmann)** U. Erdmann , W. Ebeling, L. Schimansky-Geier, and F. Schweitzer, Eur. Phys. J. B 15, 105-113, 2000.

**(Winkler)** Winkler, Wysocki, and Gompper, Soft Matter, 11, 6680 (2015).

## 17.159 fix property/atom command

Accelerator Variants: *property/atom/kk*

### 17.159.1 Syntax

```
fix ID group-ID property/atom name1 name2 ... keyword value ...
```

- ID, group-ID are documented in *fix* command
- property/atom = style name of this fix command
- name1,name2,... = *mol* or *q* or *rmas*s or *i\_name* or *d\_name* or *i2\_name* or *d2\_name*
  - mol* = molecule IDs
  - q* = charge
  - rmas*s = per-atom mass
  - i\_name* = new integer vector referenced by name
  - d\_name* = new floating-point vector referenced by name
  - i2\_name* = new integer array referenced by name
    - i2\_name* arg = N = number of columns in the array
  - d2\_name* = new floating-point array referenced by name
    - d2\_name* arg = N = number of columns in the array
- zero or more keyword/value pairs may be appended
- keyword = *ghost*
  - ghost* value = *no* or *yes* for whether ghost atom info is communicated

### 17.159.2 Examples

```
fix 1 all property/atom mol
fix 1 all property/atom i_myflag1 i_myflag2
fix 1 all property/atom d2_sxyz 3 ghost yes
```

### 17.159.3 Description

Create one or more additional per-atom vectors or arrays to store information about atoms and to use during a simulation. The specified *group-ID* is ignored by this fix.

The atom style used for a simulation defines a set of per-atom properties, as explained on the *atom\_style* and *read\_data* doc pages. The latter command defines these properties for each atom in the system when a data file is read. This fix augments the set of per-atom properties with new custom ones. This can be useful in several scenarios.

If the atom style does not define molecule IDs, per-atom charge, or per-atom mass, they can be added using the *mol*, *q* or *rmass* keywords. This could be useful to define “molecules” to use as rigid bodies with the *fix rigid* command, or to carry around an extra flag with atoms (stored as a molecule ID) that can be used by various commands like *compute chunk/atom* to group atoms without having to use the group command (which is limited to a total of 32 groups including all).

Another application is to use the *rmass* flag in order to have per-atom masses instead of per-type masses. This could be used to study isotope effects with partial isotope substitution. *See below* for an example of simulating a mixture of light and heavy water with the TIP4P water potential.

An alternative to using *fix property/atom* for these examples is to use an atom style that does define molecule IDs or charge or per-atom mass (indirectly via diameter and density) or to use a hybrid atom style that combines two or more atom styles to provide the union of all their atom properties. However, this has two practical drawbacks: first, it typically necessitates changing the format of the Atoms section in the data file and second, it may define additional properties that are not needed such as bond lists, which incurs some overhead when there are no bonds.

In the future, we may add additional existing per-atom properties to *fix property/atom*, similar to *mol*, *q* or *rmass*, which “turn-on” specific properties defined by some atom styles, so they can be easily used by atom styles that do not define them.

More generally, the *i\_name* and *d\_name* options allow one or more new custom per-atom vectors to be defined. Likewise the *i2\_name* and *d2\_name* options allow one or more custom per-atom arrays to be defined. The *i2\_name* and *d2\_name* options take an argument *N* which specifies the number of columns in the per-atom array, i.e. the number of attributes associated with each atom.  $N \geq 1$  is required.

Each name must be unique and can use alphanumeric or underscore characters. These vectors and arrays can store whatever values you decide are useful in your simulation. As explained below there are several ways to initialize, access, and output these values, via input script commands, data files, and in new code you add to LAMMPS.

This is effectively a simple way to add per-atom properties to a model without needing to write code for a new *atom style* that defines the properties. Note however that implementing a new atom style allows new atom properties to be more tightly and seamlessly integrated with the rest of the code.

The new atom properties encode values that migrate with atoms to new processors and are written to restart files. If you want the new properties to also be defined for ghost atoms, then use the *ghost* keyword with a value of *yes*. This will invoke extra communication when ghost atoms are created (at every re-neighborings) to insure the new properties are also defined for the ghost atoms.

---

#### Properties on ghost atoms

If you use the *mol*, *q* or *rmass* names, you most likely want to set *ghost yes*, since these properties are stored with ghost atoms if you use an *atom\_style* that defines them. Many LAMMPS operations that use molecule IDs or charge, such as neighbor lists and pair styles, will expect ghost atoms to have these values. LAMMPS will issue a warning if you define those vectors but do not set *ghost yes*.

---

#### Limitations on ghost atom properties

---

The specified properties for ghost atoms are not updated every timestep, but only once every few steps when neighbor lists are re-built. Thus the *ghost* keyword is suitable for static properties, like molecule IDs, but not for dynamic properties that change every step. For the latter, the code you add to LAMMPS to change the properties will also need to communicate their new values to/from ghost atoms, an operation that can be invoked from within a *pair style* or *fix* or *compute* that you write.

This fix is one of a small number that can be defined in an input script before the simulation box is created or atoms are defined. This is so it can be used with the *read\_data* command as described next.

Per-atom properties that are defined by the *atom style* are initialized when atoms are created, e.g. by the *read\_data* or *create\_atoms* commands. The per-atom properties defined by this fix are not. So you need to initialize them explicitly. One way to do this is *read\_data* command, using its *fix* keyword and passing it the fix-ID of this fix.

Thus these commands:

```
fix prop all property/atom mol d_flag
read_data data.txt fix prop NULL Molecules
```

would allow a data file to have a section like this:

```
Molecules
1 4 1.5
2 4 3.0
3 10 1.0
4 10 1.0
5 10 1.0
...
N 763 4.5
```

where N is the number of atoms, the first field on each line is the atom-ID, the next two are a molecule-ID and a floating point value that will be stored in a new property called “flag”. If a per-atom array was specified in the *fix property/atom* command then the *N* values for that array must be specified consecutively for that property on each line. Note that the order of values on each line corresponds to the order of custom names in the *fix property/atom* command.

Note that the the lines of per-atom properties can be listed in any order. Also note that all the per-atom properties specified by the fix ID (*prop* in this case) must be included on each line in the specified data file section (*Molecules* in this case).

Another way of initializing the new properties is via the *set* command. For example, if you wanted molecules defined for every set of 10 atoms, based on their atom-IDs, these commands could be used:

```
fix prop all property/atom mol
variable cluster atom ((id-1)/10)+1
set atom * mol v_cluster
```

The *atom-style variable* will create values for atoms with IDs 31,32,33,...40 that are 4.0,4.1,4.2,...,4.9. When the *set* commands assigns them to the molecule ID for each atom, they will be truncated to an integer value, so atoms 31-40 will all be assigned a molecule ID of 4.

Note that *atomfile-style variables* can also be used in place of atom-style variables, which means in this case that the molecule IDs could be read-in from a separate file and assigned by the *set* command. This allows you to initialize new per-atom properties in a completely general fashion.

For new atom properties specified as *i\_name*, *d\_name*, *i2\_name*, or *d2\_name*, the *dump custom* and *compute property/atom* commands can access their values. This means that the values can be used accessed by fixes like *fix ave/atom*, accessed by other computes like *compute reduce*, or used in *atom-style variables*.

For example, these commands will output both the instantaneous and time-averaged values of two new properties to a custom dump file:

```
fix myprops all property/atom i_flag1 d_flag2
compute 1 all property/atom i_flag1 d_flag2
fix 1 all ave/atom 10 10 100 c_1[1] c_1[2]
dump 1 all custom 100 tmp.dump id x y z i_flag1 d_flag2 f_1[1] f_1[2]
```

---

If you wish to add new *pair styles*, *fixes*, or *computes* that use the per-atom properties defined by this fix, see the *Modify atom* doc page which has details on how the custom properties of this fix can be accessed from added classes.

---

Here is an example of using per-atom masses with TIP4P water to study isotope effects. When setting up simulations with the *TIP4P pair styles* for water, you have to provide exactly one atom type each to identify the water oxygen and hydrogen atoms. Since the atom mass is normally tied to the atom type, this makes it impossible to study multiple isotopes in the same simulation. With *fix property/atom rmass* however, the per-type masses are replaced by per-atom masses. Assuming you have a working input deck for regular TIP4P water, where water oxygen is atom type 1 and water hydrogen is atom type 2, the following lines of input script convert this to using per-atom masses:

```
fix Isotopes all property/atom rmass ghost yes
set type 1 mass 15.9994
set type 2 mass 1.008
```

When writing out the system data with the *write\_data* command, there will be a new section named with the fix-ID (i.e. *Isotopes* in this case). Alternatively, you can take an existing data file and just add this *Isotopes* section with one line per atom containing atom-ID and mass. Either way, the extended data file can be read back with:

```
fix Isotopes all property/atom rmass ghost yes
read_data tip4p-isotopes.data fix Isotopes NULL Isotopes
```

Please note that the first *Isotopes* refers to the fix-ID and the second to the name of the section. The following input script code will now change the first 100 water molecules in this example to heavy water:

```
group hwat id 2:300:3
group hwat id 3:300:3
set group hwat mass 2.0141018
```

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 17.159.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the per-atom values it stores to *binary restart files*, so that the values can be restored when a simulation is restarted. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

**Warning:** When reading data from a restart file, this fix command has to be specified **after** the *read\_restart* command and **exactly** the same was in the input script that created the restart file. LAMMPS will only check whether a fix is of the same style and has the same fix ID and in case of a match will then try to initialize the fix with the data stored in the binary restart file. If the names and associated data types in the new fix property/atom command do not match the old one exactly, data can be corrupted or LAMMPS may crash. If the fix is specified **before** the *read\_restart* command its data will not be restored.

None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.159.5 Restrictions

none

### 17.159.6 Related commands

*read\_data, set, compute property/atom*

### 17.159.7 Default

The default keyword value is ghost = no.

## 17.160 fix python/invoke command

### 17.160.1 Syntax

```
fix ID group-ID python/invoke N callback function_name
```

- ID, group-ID are ignored by this fix
- python/invoke = style name of this fix command
- N = execute every N steps
- callback = *post\_force* or *end\_of\_step*

*post\_force* = callback after force computations on atoms every N time steps

*end\_of\_step* = callback after every N time steps

## 17.160.2 Examples

```
python post_force_callback here """
from lammmps import lammmps

def post_force_callback(lammmps_ptr, vflag):
    lmp = lammmps(ptr=lammmps_ptr)
    # access LAMMPS state using Python interface
    """

python end_of_step_callback here """
def end_of_step_callback(lammmps_ptr):
    lmp = lammmps(ptr=lammmps_ptr)
    # access LAMMPS state using Python interface
    """

fix pf all python/invoke 50 post_force post_force_callback
fix eos all python/invoke 50 end_of_step end_of_step_callback
```

## 17.160.3 Description

This fix allows you to call a Python function during a simulation run. The callback is either executed after forces have been applied to atoms or at the end of every N time steps.

Callback functions must be declared in the global scope of the active Python interpreter. This can either be done by defining it inline using the python command or by importing functions from other Python modules. If LAMMPS is driven using the library interface from Python, functions defined in the driving Python interpreter can also be executed.

Each callback is given a pointer object as first argument. This can be used to initialize an instance of the lammmps Python interface, which gives access to the LAMMPS state from Python.

**Warning:** While you can access the state of LAMMPS via library functions from these callbacks, trying to execute input script commands will in the best case not work or in the worst case result in undefined behavior.

## 17.160.4 Restrictions

This fix is part of the PYTHON package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

Building LAMMPS with the PYTHON package will link LAMMPS with the Python library on your system. Settings to enable this are in the lib/python/Makefile.lammmps file. See the lib/python/README file for information on those settings.

## 17.160.5 Related commands

*python command*

## 17.161 fix python/move command

### 17.161.1 Syntax

```
fix python/move pmodule.CLASS
```

pmodule.CLASS = use class **CLASS** in module/file **pmodule** to compute how to move atoms

### 17.161.2 Examples

```
fix 1 all python/move py_nve.NVE
fix 1 all python/move py_nve.NVE_OPT
```

### 17.161.3 Description

The *python/move* fix style provides a way to define ways how particles are moved during an MD run from python script code, that is loaded from a file into LAMMPS and executed at the various steps where other fixes can be executed. This python script must contain specific python class definitions.

This allows to implement complex position updates and also modified time integration methods. Due to python being an interpreted language, however, the performance of this fix can be moderately to significantly slower than the corresponding C++ code. For specific cases, this performance penalty can be limited through effective use of NumPy.

The python module file has to start with the following code:

```
from __future__ import print_function
import lammmps
import ctypes
import traceback
import numpy as np
#
class LAMMPSFix(object):
    def __init__(self, ptr, group_name="all"):
        self.lmp = lammmps.lammmps(ptr=ptr)
        self.group_name = group_name
#
class LAMMPSFixMove(LAMMPSFix):
    def __init__(self, ptr, group_name="all"):
        super(LAMMPSFixMove, self).__init__(ptr, group_name)
#
    def init(self):
        pass
#
    def initial_integrate(self, vflag):
```

(continues on next page)

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```
    pass
#
def final_integrate(self):
    pass
#
def initial_integrate_respa(self, vflag, ilevel, iloop):
    pass
#
def final_integrate_respa(self, ilevel, iloop):
    pass
#
def reset_dt(self):
    pass
```

Any classes implementing new atom motion functionality have to be derived from the `LAMMPSFixMove` class, overriding the available methods as needed.

Examples for how to do this are in the `examples/python` folder.

---

#### 17.161.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.161.5 Restrictions

This pair style is part of the PYTHON package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 17.161.6 Related commands

*fix nve, fix python/invoke*

#### 17.161.7 Default

none

### 17.162 fix qbmsst command

#### 17.162.1 Syntax

```
fix ID group-ID qbmsst dir shockvel keyword value ...
```

- ID, group-ID are documented in *fix* command



- `qbmsst` = style name of this fix
- `dir` = `x` or `y` or `z`
- `shockvel` = shock velocity (strictly positive, velocity units)
- zero or more keyword/value pairs may be appended
- keyword = `q` or `mu` or `p0` or `v0` or `e0` or `tscale` or `damp` or `seed` or `f_max` or `N_f` or `eta` or `beta` or `T_init`

`q` value = cell mass-like parameter (mass<sup>2</sup>/distance<sup>4</sup> units)

`mu` value = artificial viscosity (mass/distance/time units)

`p0` value = initial pressure in the shock equations (pressure units)

`v0` value = initial simulation cell volume in the shock equations (distance<sup>3</sup> units)

`e0` value = initial total energy (energy units)

`tscale` value = reduction in initial temperature (unitless fraction between 0.0 and 1.0)

`damp` value = damping parameter (time units) inverse of friction  $\gamma$

`seed` value = random number seed (positive integer)

`f_max` value = upper cutoff frequency of the vibration spectrum (1/time units)

`N_f` value = number of frequency bins (positive integer)

`eta` value = coupling constant between the shock system and the quantum thermal bath (positive unitless)

`beta` value = the quantum temperature is updated every `beta` time steps (positive integer)

`T_init` value = quantum temperature for the initial state (temperature units)

## 17.162.2 Examples

```
# (liquid methane modeled with the REAX force field, real units)
fix 1 all qbmsst z 0.122 q 25 mu 0.9 tscale 0.01 damp 200 seed 35082 f_max 0.3 N_f 100
eta 1 beta 400 T_init 110
# (quartz modeled with the BKS force field, metal units)
fix 2 all qbmsst z 72 q 40 tscale 0.05 damp 1 seed 47508 f_max 120.0 N_f 100 eta 1.0
beta 500 T_init 300
```

Two example input scripts are given, including shocked  $\alpha$ -quartz and shocked liquid methane. The input script first equilibrates an initial state with the quantum thermal bath at the target temperature and then applies `fix qbmsst` to simulate shock compression with quantum nuclear correction. The following two figures plot relevant quantities for shocked  $\alpha$ -quartz.

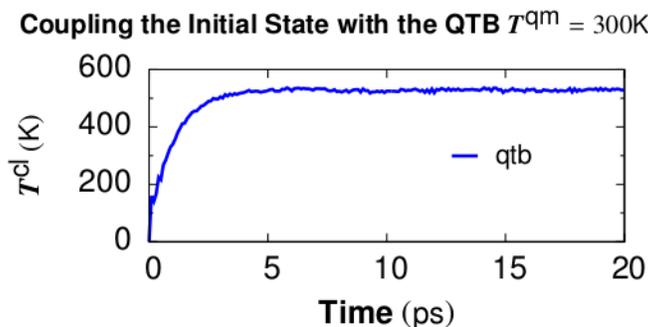


Figure 1. Classical temperature  $T_{cl} = \sum \frac{m_i v_i^2}{3Nk_B}$  vs. time for coupling the  $\alpha$ -quartz initial state with the quantum thermal bath at target quantum temperature  $T^{qm} = 300K$ . The NpH ensemble is used for time integration while QTB provides the colored random force.  $T^{cl}$  converges at the timescale of `damp` which is set to be 1 ps.

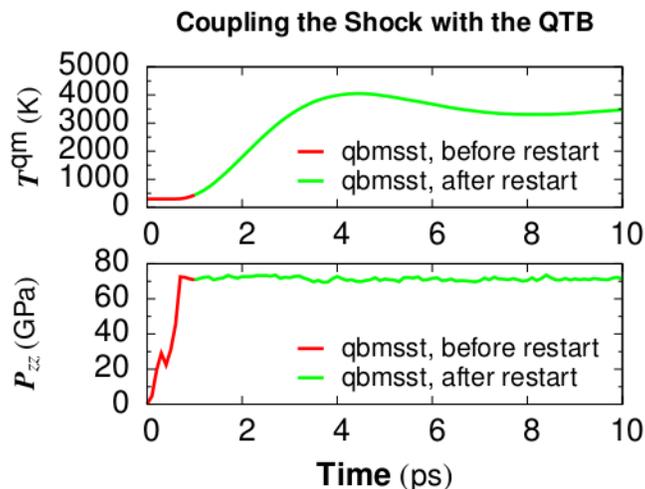


Figure 2. Quantum temperature and pressure vs. time for simulating shocked  $\alpha$ -quartz with *fix qbmsst*. The shock propagates along the *z* direction. Restart of the *fix qbmsst* command is demonstrated in the example input script. Thermodynamic quantities stay continuous before and after the restart.

### 17.162.3 Description

This command performs the Quantum-Bath coupled Multi-Scale Shock Technique (QBMSST) integration. See (*Qi*) for a detailed description of this method. QBMSST provides description of the thermodynamics and kinetics of shock processes while incorporating quantum nuclear effects. The *shockvel* setting determines the steady shock velocity that will be simulated along direction *dir*.

Quantum nuclear effects (*fix qtb*) can be crucial especially when the temperature of the initial state is below the classical limit or there is a great change in the zero point energies between the initial and final states. Theoretical post processing quantum corrections of shock compressed water and methane have been reported as much as 30% of the temperatures (*Goldman*). A self-consistent method that couples the shock to a quantum thermal bath described by a colored noise Langevin thermostat has been developed by Qi et al (*Qi*) and applied to shocked methane. The onset of chemistry is reported to be at a pressure on the shock Hugoniot that is 40% lower than observed with classical molecular dynamics.

It is highly recommended that the system be already in an equilibrium state with a quantum thermal bath at temperature of  $T_{init}$ . The *fix* command *fix qtb* at constant temperature  $T_{init}$  could be used before applying this command to introduce self-consistent quantum nuclear effects into the initial state.

The parameters *q*, *mu*, *e0*, *p0*, *v0* and *tscale* are described in the command *fix msst*. The values of *e0*, *p0*, or *v0* will be calculated on the first step if not specified. The parameter of *damp*, *f\_max*, and *N\_f* are described in the command *fix qtb*.

The *fix qbmsst* command couples the shock system to a quantum thermal bath with a rate that is proportional to the change of the total energy of the shock system,  $E^{tot} - E_0^{tot}$ . Here  $E^{tot}$  consists of both the system energy and a thermal term, see (*Qi*), and  $E_0^{tot} = e0$  is the initial total energy.

The *eta* ( $\eta$ ) parameter is a unitless coupling constant between the shock system and the quantum thermal bath. A small  $\eta$  value cannot adjust the quantum temperature fast enough during the temperature ramping period of shock compression while large  $\eta$  leads to big temperature oscillation. A value of  $\eta$  between 0.3 and 1 is usually appropriate for simulating most systems under shock compression. We observe that different values of  $\eta$  lead to almost the same final thermodynamic state behind the shock, as expected.

The quantum temperature is updated every *beta* ( $\beta$ ) steps with an integration time interval  $\beta$  times longer than the simulation time step. In that case,  $E^{tot}$  is taken as its average over the past  $\beta$  steps. The temperature of the quantum

thermal bath  $T^{qm}$  changes dynamically according to the following equation where  $\Delta_t$  is the MD time step and  $\gamma$  is the friction constant which is equal to the inverse of the *damp* parameter.

$$\frac{dT^{qm}}{dt} = \gamma\eta \sum_{l=1}^{\beta} \frac{E^{tot}(t - l\Delta t) - E_0^{tot}}{3\beta N k_B}$$

The parameter *T\_init* is the initial temperature of the quantum thermal bath and the system before shock loading.

For all pressure styles, the simulation box stays orthorhombic in shape. Parrinello-Rahman boundary conditions (tilted box) are supported by LAMMPS, but are not implemented for QBMSST.

#### 17.162.4 Restart, fix\_modify, output, run start/stop, minimize info

Because the state of the random number generator is not written to *binary restart files*, this fix cannot be restarted “exactly” in an uninterrupted fashion. However, in a statistical sense, a restarted simulation should produce similar behaviors of the system as if it is not interrupted. To achieve such a restart, one should write explicitly the same value for *q*, *mu*, *damp*, *f\_max*, *N\_f*, *eta*, and *beta* and set *tscale* = 0 if the system is compressed during the first run.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

The progress of the QBMSST can be monitored by printing the global scalar and global vector quantities computed by the fix.

As mentioned above, the scalar is the cumulative energy change due to the fix. By monitoring the *thermodynamic econserve* output, this can be used to test if the MD timestep is sufficiently small for accurate integration of the dynamic equations.

The global vector contains five values in the following order. The vector values output by this fix are “intensive”.

[*dhugoniot*, *drayleigh*, *lagrangian\_speed*, *lagrangian\_position*, *quantum\_temperature*]

1. *dhugoniot* is the departure from the Hugoniot (temperature units).
2. *drayleigh* is the departure from the Rayleigh line (pressure units).
3. *lagrangian\_speed* is the laboratory-frame Lagrangian speed (particle velocity) of the computational cell (velocity units).
4. *lagrangian\_position* is the computational cell position in the reference frame moving at the shock speed. This is the distance of the computational cell behind the shock front.
5. *quantum\_temperature* is the temperature of the quantum thermal bath  $T^{qm}$ .

To print these quantities to the log file with descriptive column headers, the following LAMMPS commands are suggested.

```
fix                fix_id all msst z
variable           dhug    equal f_fix_id[1]
variable           dray    equal f_fix_id[2]
variable           lgr_vel equal f_fix_id[3]
variable           lgr_pos equal f_fix_id[4]
variable           T_qm    equal f_fix_id[5]
thermo_style       custom  step temp ke pe lz pzz econserve v_dhug v_dray v_lgr_vel v_lgr_
↳pos v_T_qm f_fix_id
```

It is worth noting that the temp keyword for the *thermo\_style* command prints the instantaneous classical temperature  $T^{cl}$  as described by the *fix qtb* command.

---

### 17.162.5 Restrictions

This fix style is part of the QTB package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

All cell dimensions must be periodic. This fix can not be used with a triclinic cell. The QBMSST fix has been tested only for the group-ID all.

---

### 17.162.6 Related commands

*fix qtb, fix msst*

---

### 17.162.7 Default

The keyword defaults are q = 10, mu = 0, tscale = 0.01, damp = 1, seed = 880302, f\_max = 200.0, N\_f = 100, eta = 1.0, beta = 100, and T\_init=300.0. e0, p0, and v0 are calculated on the first step.

---

(Goldman) Goldman, Reed and Fried, J. Chem. Phys. 131, 204103 (2009)

(Qi) Qi and Reed, J. Phys. Chem. A 116, 10451 (2012).

## 17.163 fix qeq/point command

## 17.164 fix qeq/shielded command

## 17.165 fix qeq/slater command

## 17.166 fix qeq/dynamic command

## 17.167 fix qeq/fire command

### 17.167.1 Syntax

```
fix ID group-ID style Nevery cutoff tolerance maxiter qfile keyword ...
```

- ID, group-ID are documented in *fix* command
- style = *qeq/point* or *qeq/shielded* or *qeq/slater* or *qeq/dynamic* or *qeq/fire*

- Nevery = perform charge equilibration every this many steps
- cutoff = global cutoff for charge-charge interactions (distance unit)
- tolerance = precision to which charges will be equilibrated
- maxiter = maximum iterations to perform charge equilibration
- qfile = a filename with QEq parameters or *coul/streitz* or *reaxff*
- zero or more keyword/value pairs may be appended
- keyword = *alpha* or *qdamp* or *qstep* or *warn*

*alpha* value = Slater type orbital exponent (qeq/slater only)

*qdamp* value = damping factor for damped dynamics charge solver (qeq/dynamic and qeq/fire only)

*qstep* value = time step size for damped dynamics charge solver (qeq/dynamic and qeq/fire only)

*warn* value = do (=yes) or do not (=no) print a warning when the maximum number of iterations is reached

## 17.167.2 Examples

```
fix 1 all qeq/point 1 10 1.0e-6 200 param.qeq1
fix 1 qeq qeq/shielded 1 8 1.0e-6 100 param.qeq2
fix 1 all qeq/slater 5 10 1.0e-6 100 params alpha 0.2
fix 1 qeq qeq/dynamic 1 12 1.0e-3 100 my_qeq
fix 1 all qeq/fire 1 10 1.0e-3 100 my_qeq qdamp 0.2 qstep 0.1
```

## 17.167.3 Description

Perform the charge equilibration (QEq) method as described in (*Rappe and Goddard*) and formulated in (*Nakano*) (also known as the matrix inversion method) and in (*Rick and Stuart*) (also known as the extended Lagrangian method) based on the electronegativity equilization principle.

These fixes can be used with any *pair style* in LAMMPS, so long as per-atom charges are defined. The most typical use-case is in conjunction with a *pair style* that performs charge equilibration periodically (e.g. every timestep), such as the ReaxFF or Streitz-Mintmire potential. But these fixes can also be used with potentials that normally assume per-atom charges are fixed, e.g. a *Buckingham* or *LJ/Coulombic* potential.

Because the charge equilibration calculation is effectively independent of the pair style, these fixes can also be used to perform a one-time assignment of charges to atoms. For example, you could define the QEq fix, perform a zero-timestep run via the *run* command without any pair style defined which would set per-atom charges (based on the current atom configuration), then remove the fix via the *unfix* command before performing further dynamics.

---

**Note:** Computing and using charge values different from published values defined for a fixed-charge potential like Buckingham or CHARMM or AMBER, can have a strong effect on energies and forces, and produces a different model than the published versions.

---



---

**Note:** The *fix qeq/comb* command must still be used to perform charge equilibration with the *COMB potential*. The *fix qeq/reaxff* command can be used to perform charge equilibration with the *ReaxFF force field*, although *fix qeq/shielded*

---

yields the same results as `fix qeq/reaxff` if *Nevery*, *cutoff*, and *tolerance* are the same. Eventually the `fix qeq/reaxff` command will be deprecated.

---

The QEq method minimizes the electrostatic energy of the system (or equalizes the derivative of energy with respect to charge of all the atoms) by adjusting the partial charge on individual atoms based on interactions with their neighbors within *cutoff*. It requires a few parameters in the appropriate units for each atom type which are read from a file specified by *qfile*. The file has the following format

```
1 chi eta gamma zeta qcore
2 chi eta gamma zeta qcore
...
Ntype chi eta gamma zeta qcore
```

There have to be parameters given for every atom type. Wildcard entries are possible using the same type range syntax as for “coeff” commands (i.e., *n\*m*, *n\**, *\*m*, *\**). Later entries will overwrite previous ones. Empty lines or any text following the pound sign (#) are ignored. Each line starts with the atom type followed by five parameters. Only a subset of the parameters is used by each QEq style as described below, thus the others can be set to 0.0 if desired, but all five entries per line are required.

- *chi* = electronegativity in energy units
- *eta* = self-Coulomb potential in energy units
- *gamma* = shielded Coulomb constant defined by *ReaxFF force field* in distance units
- *zeta* = Slater type orbital exponent defined by the *Streitz-Mintmire* potential in reverse distance units
- *qcore* = charge of the nucleus defined by the *Streitz-Mintmire potential* potential in charge units

The `fix qeq` styles will print a warning if the charges are not equilibrated within *tolerance* by *maxiter* steps, unless the *warn* keyword is used with “no” as argument. This latter option may be useful for testing and benchmarking purposes, as it allows to use a fixed number of QEq iterations when *tolerance* is set to a small enough value to always reach the *maxiter* limit. Turning off warnings will avoid the excessive output in that case.

The *qeq/point* style describes partial charges on atoms as point charges. Interaction between a pair of charged particles is  $1/r$ , which is the simplest description of the interaction between charges. Only the *chi* and *eta* parameters from the *qfile* file are used. Note that Coulomb catastrophe can occur if repulsion between the pair of charged particles is too weak. This style solves partial charges on atoms via the matrix inversion method. A tolerance of  $1.0e-6$  is usually a good number.

The *qeq/shielded* style describes partial charges on atoms also as point charges, but uses a shielded Coulomb potential to describe the interaction between a pair of charged particles. Interaction through the shielded Coulomb is given by equation (13) of the *ReaxFF force field* paper. The shielding accounts for charge overlap between charged particles at small separation. This style is the same as *fix qeq/reaxff*, and can be used with *pair\_style reaxff*. Only the *chi*, *eta*, and *gamma* parameters from the *qfile* file are used. When using the string *reaxff* as filename, these parameters are extracted directly from an active *reaxff* pair style. This style solves partial charges on atoms via the matrix inversion method. A tolerance of  $1.0e-6$  is usually a good number.

The *qeq/slater* style describes partial charges on atoms as spherical charge densities centered around atoms via the Slater *1s* orbital, so that the interaction between a pair of charged particles is the product of two Slater *1s* orbitals. The expression for the Slater *1s* orbital is given under equation (6) of the *Streitz-Mintmire* paper. Only the *chi*, *eta*, *zeta*, and *qcore* parameters from the *qfile* file are used. When using the string *coul/streitz* as filename, these parameters are extracted directly from an active *coul/streitz* pair style. This style solves partial charges on atoms via the matrix inversion method. A tolerance of  $1.0e-6$  is usually a good number. Keyword *alpha* can be used to change the Slater type orbital exponent.

The *qeq/dynamic* style describes partial charges on atoms as point charges that interact through  $1/r$ , but the extended Lagrangian method is used to solve partial charges on atoms. Only the *chi* and *eta* parameters from the *qfile* file are used. Note that Coulomb catastrophe can occur if repulsion between the pair of charged particles is too weak. A tolerance of

1.0e-3 is usually a good number. Keyword *qdamp* can be used to change the damping factor, while keyword *qstep* can be used to change the time step size.

The *\*qeq/fire\** style describes the same charge model and charge solver as the *qeq/dynamic* style, but employs a FIRE minimization algorithm to solve for equilibrium charges. Keyword *qdamp* can be used to change the damping factor, while keyword *qstep* can be used to change the time step size.

Note that *qeq/point*, *qeq/shielded*, and *qeq/slater* describe different charge models, whereas the matrix inversion method and the extended Lagrangian method (*qeq/dynamic* and *qeq/fire*) are different solvers.

Note that *qeq/point*, *qeq/dynamic* and *qeq/fire* styles all describe charges as point charges that interact through 1/r relationship, but solve partial charges on atoms using different solvers. These three styles should yield comparable results if the QEq parameters and *Nevery*, *cutoff*, and *tolerance* are the same. Style *qeq/point* is typically faster, *qeq/dynamic* scales better on larger sizes, and *qeq/fire* is faster than *qeq/dynamic*.

---

**Note:** In order to solve the self-consistent equations for electronegativity equalization, LAMMPS imposes the additional constraint that all the charges in the fix group must add up to zero. The initial charge assignments should also satisfy this constraint. LAMMPS will print a warning if that is not the case.

---

---

**Note:** Developing QEq parameters (*chi*, *eta*, *gamma*, *zeta*, and *qcore*) is non-trivial. Charges on atoms are not guaranteed to equilibrate with arbitrary choices of these parameters. We do not develop these QEq parameters. See the *examples/qeq* directory for some examples.

---

#### 17.167.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about these fixes is written to *binary restart files*. No global scalar or vector or per-atom quantities are stored by these fixes for access by various *output commands*. No parameter of these fixes can be used with the *start/stop* keywords of the *run* command.

These fixes are invoked during *energy minimization*.

#### 17.167.5 Restrictions

These fixes are part of the QEQ package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These qeq fixes are not compatible with the GPU and USER-INTEL packages.

These qeq fixes will ignore electric field contributions from *fix efield*.

#### 17.167.6 Related commands

*fix qeq/reaxff*, *fix qeq/comb*

## 17.167.7 Default

warn yes

---

**(Rappe and Goddard)** A. K. Rappe and W. A. Goddard III, J Physical Chemistry, 95, 3358-3363 (1991).

**(Nakano)** A. Nakano, Computer Physics Communications, 104, 59-69 (1997).

**(Rick and Stuart)** S. W. Rick, S. J. Stuart, B. J. Berne, J Chemical Physics 101, 16141 (1994).

**(Streitz-Mintmire)** F. H. Streitz, J. W. Mintmire, Physical Review B, 50, 16, 11996 (1994)

**(ReaxFF)** A. C. T. van Duin, S. Dasgupta, F. Lorant, W. A. Goddard III, J Physical Chemistry, 105, 9396-9049 (2001)

**(QEq/Fire)** T-R. Shan, A. P. Thompson, S. J. Plimpton, in preparation

## 17.168 fix qeq/comb command

Accelerator Variants: *qeq/comb/omp*

### 17.168.1 Syntax

```
fix ID group-ID qeq/comb Nevery precision keyword value ...
```

- ID, group-ID are documented in *fix* command
- qeq/comb = style name of this fix command
- Nevery = perform charge equilibration every this many steps
- precision = convergence criterion for charge equilibration
- zero or more keyword/value pairs may be appended
- keyword = *file*

*file* value = filename

filename = name of file to write QEQ equilibration info to

### 17.168.2 Examples

```
fix 1 surface qeq/comb 10 0.0001
```

### 17.168.3 Description

Perform charge equilibration (QeQ) in conjunction with the COMB (Charge-Optimized Many-Body) potential as described in (*COMB\_1*) and (*COMB\_2*). It performs the charge equilibration portion of the calculation using the so-called QEq method, whereby the charge on each atom is adjusted to minimize the energy of the system. This fix can only be used with the COMB potential; see the *fix qeq/reaxff* command for a QeQ calculation that can be used with any potential.

Only charges on the atoms in the specified group are equilibrated. The fix relies on the pair style (COMB in this case) to calculate the per-atom electronegativity (effective force on the charges). An electronegativity equalization calculation



(or QEq) is performed in an iterative fashion, which in parallel requires communication at each iteration for processors to exchange charge information about nearby atoms with each other. See *Rappe\_and\_Goddard* and *Rick\_and\_Stuart* for details.

During a run, charge equilibration is performed every *Nevery* time steps. Charge equilibration is also always enforced on the first step of each run. The *precision* argument controls the tolerance for the difference in electronegativity for all atoms during charge equilibration. *Precision* is a trade-off between the cost of performing charge equilibration (more iterations) and accuracy.

If the *file* keyword is used, then information about each equilibration calculation is written to the specified file.

---

**Note:** In order to solve the self-consistent equations for electronegativity equalization, LAMMPS imposes the additional constraint that all the charges in the fix group must add up to zero. The initial charge assignments should also satisfy this constraint. LAMMPS will print a warning if that is not the case.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 17.168.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is performing charge equilibration. Default is the outermost level.

This fix produces a per-atom vector which can be accessed by various *output commands*. The vector stores the gradient of the charge on each atom. The per-atom values be accessed on any timestep.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

This fix can be invoked during *energy minimization*.

## 17.168.5 Restrictions

This fix command currently only supports *pair style \*comb\**.

## 17.168.6 Related commands

*pair\_style comb*

## 17.168.7 Default

No file output is performed.

---

(**COMB\_1**) J. Yu, S. B. Sinnott, S. R. Phillpot, Phys Rev B, 75, 085311 (2007),

(**COMB\_2**) T.-R. Shan, B. D. Devine, T. W. Kemper, S. B. Sinnott, S. R. Phillpot, Phys Rev B, 81, 125328 (2010).

(**Rappe\_and\_Goddard**) A. K. Rappe, W. A. Goddard, J Phys Chem 95, 3358 (1991).

(**Rick\_and\_Stuart**) S. W. Rick, S. J. Stuart, B. J. Berne, J Chem Phys 101, 16141 (1994).

## 17.169 fix qeq/reaxff command

Accelerator Variants: *qeq/reaxff/kk*, *qeq/reaxff/omp*

### 17.169.1 Syntax

```
fix ID group-ID qeq/reaxff Nevery cutlo cuthi tolerance params args
```

- ID, group-ID are documented in *fix* command
- qeq/reaxff = style name of this fix command
- Nevery = perform QEq every this many steps
- cutlo,cuthi = lo and hi cutoff for Taper radius
- tolerance = precision to which charges will be equilibrated
- params = reaxff or a filename
- one or more keywords or keyword/value pairs may be appended

keyword = *dual* or *maxiter* or *nowarn*

*dual* = process S and T matrix in parallel (only for qeq/reaxff/omp)

*maxiter* N = limit the number of iterations to N

*nowarn* = do not print a warning message if the maximum number of iterations was reached

### 17.169.2 Examples

```
fix 1 all qeq/reaxff 1 0.0 10.0 1.0e-6 reaxff
fix 1 all qeq/reaxff 1 0.0 10.0 1.0e-6 param.qeq maxiter 500
```

### 17.169.3 Description

Perform the charge equilibration (QEq) method as described in (*Rappe and Goddard*) and formulated in (*Nakano*). It is typically used in conjunction with the ReaxFF force field model as implemented in the *pair\_style reaxff* command, but it can be used with any potential in LAMMPS, so long as it defines and uses charges on each atom. The *fix qeq/comb* command should be used to perform charge equilibration with the *COMB potential*. For more technical details about the charge equilibration performed by *fix qeq/reaxff*, see the (*Aktulga*) paper.

The QEq method minimizes the electrostatic energy of the system by adjusting the partial charge on individual atoms based on interactions with their neighbors. It requires some parameters for each atom type. If the *params* setting above is the word “reaxff”, then these are extracted from the *pair\_style reaxff* command and the ReaxFF force field file it reads in. If a file name is specified for *params*, then the parameters are taken from the specified file and the file must contain one line for each atom type. The latter form must be used when performing QeQ with a non-ReaxFF potential. Each line should be formatted as follows:

```
itype chi eta gamma
```

where *itype* is the atom type from 1 to Ntypes, *chi* denotes the electronegativity in eV, *eta* denotes the self-Coulomb potential in eV, and *gamma* denotes the valence orbital exponent. Note that these 3 quantities are also in the ReaxFF potential file, except that *eta* is defined here as twice the *eta* value in the ReaxFF file. Note that unlike the rest of LAMMPS, the units of this *fix* are hard-coded to be Å, eV, and electronic charge.

The optional *dual* keyword allows to perform the optimization of the S and T matrices in parallel. This is only supported for the *qeq/reaxff/omp* style. Otherwise they are processed separately. The *qeq/reaxff/kk* style always solves the S and T matrices in parallel.

The optional *maxiter* keyword allows changing the max number of iterations in the linear solver. The default value is 200.

The optional *nowarn* keyword silences the warning message printed when the maximum number of iterations was reached. This can be useful for comparing serial and parallel results where having the same fixed number of QEq iterations is desired, which can be achieved by using a very small tolerance and setting *maxiter* to the desired number of iterations.

---

**Note:** In order to solve the self-consistent equations for electronegativity equalization, LAMMPS imposes the additional constraint that all the charges in the *fix* group must add up to zero. The initial charge assignments should also satisfy this constraint. LAMMPS will print a warning if that is not the case.

---

### 17.169.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this *fix* is written to *binary restart files*. This *fix* computes a global scalar (the number of iterations) for access by various *output commands*. No parameter of this *fix* can be used with the *start/stop* keywords of the *run* command.

This *fix* is invoked during *energy minimization*.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 17.169.5 Restrictions

This fix is part of the REAXFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix does not correctly handle interactions involving multiple periodic images of the same atom. Hence, it should not be used for periodic cell dimensions less than 10 angstroms.

This fix may be used in combination with *fix efield* and will apply the external electric field during charge equilibration, but there may be only one fix efield instance used, it may only use a constant electric field, and the electric field vector may only have components in non-periodic directions.

### 17.169.6 Related commands

*pair\_style reaxff*, *fix qeq/shielded*

### 17.169.7 Default

maxiter 200

---

**(Rappe)** Rappe and Goddard III, Journal of Physical Chemistry, 95, 3358-3363 (1991).

**(Nakano)** Nakano, Computer Physics Communications, 104, 59-69 (1997).

**(Aktulga)** Aktulga, Fogarty, Pandit, Grama, Parallel Computing, 38, 245-259 (2012).

## 17.170 fix qmmm command

### 17.170.1 Syntax

```
fix ID group-ID qmmm
```

- ID, group-ID are documented in *fix* command
- qmmm = style name of this fix command

## 17.170.2 Examples

```
fix 1 qmol qmmm
```

## 17.170.3 Description

This fix provides functionality to enable a quantum mechanics/molecular mechanics (QM/MM) coupling of LAMMPS to a quantum mechanical code. The current implementation only supports an ONIOM style mechanical coupling to the [Quantum ESPRESSO](#) plane wave DFT package. Electrostatic coupling is in preparation and the interface has been written in a manner that coupling to other QM codes should be possible without changes to LAMMPS itself.

The interface code for this is in the lib/qmmm directory of the LAMMPS distribution and is being made available at this early stage of development in order to encourage contributions for interfaces to other QM codes. This will allow the LAMMPS side of the implementation to be adapted if necessary before being finalized.

Details about how to use this fix are currently documented in the description of the QM/MM interface code itself in lib/qmmm/README.

## 17.170.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global scalar or vector or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.170.5 Restrictions

This fix is part of the QMMM package. It is only enabled if LAMMPS was built with that package. It also requires building a library provided with LAMMPS. See the [Build package](#) page for more info.

The fix is only functional when LAMMPS is built as a library and linked with a compatible QM program and a QM/MM front end into a QM/MM executable. See the lib/qmmm/README file for details.

## 17.170.6 Related commands

none

## 17.170.7 Default

none

## 17.171 fix qtb command

### 17.171.1 Syntax

```
fix ID group-ID qtb keyword value ...
```

- ID, group-ID are documented in *fix* command
- qtb = style name of this fix
- zero or more keyword/value pairs may be appended
- keyword = *temp* or *damp* or *seed* or *f\_max* or *N\_f*
  - temp* value = target quantum temperature (temperature units)
  - damp* value = damping parameter (time units) inverse of friction *gamma*
  - seed* value = random number seed (positive integer)
  - f\_max* value = upper cutoff frequency of the vibration spectrum (1/time units)
  - N\_f* value = number of frequency bins (positive integer)

### 17.171.2 Examples

```
# (liquid methane modeled with the REAX force field, real units)
fix 1 all nve
fix 1 all qtb temp 110 damp 200 seed 35082 f_max 0.3 N_f 100
# (quartz modeled with the BKS force field, metal units)
fix 2 all nph iso 1.01325 1.01325 1
fix 2 all qtb temp 300 damp 1 seed 47508 f_max 120.0 N_f 100
```

### 17.171.3 Description

This command performs the quantum thermal bath scheme proposed by (*Dammak*) to include self-consistent quantum nuclear effects, when used in conjunction with the *fix nve* or *fix nph* commands.

Classical molecular dynamics simulation does not include any quantum nuclear effect. Quantum treatment of the vibrational modes will introduce zero point energy into the system, alter the energy power spectrum and bias the heat capacity from the classical limit. Missing all the quantum nuclear effects, classical MD cannot model systems at temperatures lower than their classical limits. This effect is especially important for materials with a large population of hydrogen atoms and thus higher classical limits.

The equation of motion implemented by this command follows a Langevin form:

$$m_i a_i = f_i + R_i - m_i \gamma v_i$$

Here  $m_i$ ,  $a_i$ ,  $f_i$ ,  $R_i$ ,  $\gamma$ , and  $v_i$  represent in this order mass, acceleration, force exerted by all other atoms, random force, frictional coefficient (the inverse of damping parameter *damp*), and velocity. The random force  $R_i$  is “colored” so that any vibrational mode with frequency  $\omega$  will have a temperature-sensitive energy  $\theta(\omega, T)$  which resembles the energy expectation for a quantum harmonic oscillator with the same natural frequency:

$$\theta(\omega T) = \frac{\hbar}{2} + \hbar \omega \left[ \exp\left(\frac{\hbar \omega}{k_B T}\right) - 1 \right]^{-1}$$

To efficiently generate the random forces, we employ the method of (*Barrat*), that circumvents the need to generate all random forces for all times before the simulation. The memory requirement of this approach is less demanding

and independent of the simulation duration. Since the total random force  $R_{tot}$  does not necessarily vanish for a finite number of atoms,  $R_i$  is replaced by  $R_i - \frac{R_{tot}}{N_{tot}}$  to avoid collective motion of the system.

The *temp* parameter sets the target quantum temperature. LAMMPS will still have an output temperature in its thermo style. That is the instantaneous classical temperature  $T^{cl}$  derived from the atom velocities at thermal equilibrium. A non-zero  $T^{cl}$  will be present even when the quantum temperature approaches zero. This is associated with zero-point energy at low temperatures.

$$T^{cl} = \sum \frac{m_i v_i^2}{3Nk_B}$$

The *damp* parameter is specified in time units, and it equals the inverse of the frictional coefficient  $\gamma$ .  $\gamma$  should be as small as possible but slightly larger than the timescale of anharmonic coupling in the system which is about 10 ps to 100 ps. When  $\gamma$  is too large, it gives an energy spectrum that differs from the desired Bose-Einstein spectrum. When  $\gamma$  is too small, the quantum thermal bath coupling to the system will be less significant than anharmonic effects, reducing to a classical limit. We find that setting  $\gamma$  between 5 THz and 1 THz could be appropriate depending on the system.

The random number *seed* is a positive integer used to initiate a Marsaglia random number generator. Each processor uses the input seed to generate its own unique seed and its own stream of random numbers. Thus the dynamics of the system will not be identical on two runs on different numbers of processors.

The *f\_max* parameter truncate the noise frequency domain so that vibrational modes with frequencies higher than *f\_max* will not be modulated. If we denote  $\Delta t$  as the time interval for the MD integration, *f\_max* is always reset by the code to make  $\alpha = (int)(2 f_{max} \Delta t)^{-1}$  a positive integer and print out relative information. An appropriate value for the cutoff frequency *f\_max* would be around 2~3  $f_D$ , where  $f_D$  is the Debye frequency.

The *N\_f* parameter is the frequency grid size, the number of points from 0 to *f\_max* in the frequency domain that will be sampled.  $3*2N_f$  per-atom random numbers are required in the random force generation and there could be as many atoms as in the whole simulation that can migrate into every individual processor. A larger *N\_f* provides a more accurate sampling of the spectrum while consumes more memory. With fixed *f\_max* and  $\gamma$ , *N\_f* should be big enough to converge the classical temperature  $T^{cl}$  as a function of target quantum bath temperature. Memory usage per processor could be from 10 to 100 MBytes.

---

**Note:** Unlike the *fix nvt* command which performs Nose/Hoover thermostating AND time integration, this *fix* does NOT perform time integration. It only modifies forces to a colored thermostat. Thus you must use a separate time integration *fix*, like *fix nve* or *fix nph* to actually update the velocities and positions of atoms (as shown in the examples). Likewise, this *fix* should not normally be used with other *fixes* or commands that also specify system temperatures , e.g. *fix nvt* and *fix temp/rescale*.

---

#### 17.171.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this *fix* is written to *binary restart files*. Because the state of the random number generator is not saved in restart files, this means you cannot do “exact” restarts with this *fix*. However, in a statistical sense, a restarted simulation should produce similar behaviors of the system.

This *fix* is not invoked during *energy minimization*.

---

### 17.171.5 Restrictions

This fix style is part of the QTB package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

---

### 17.171.6 Related commands

*fix nve*, *fix nph*, *fix langevin*, *fix qbmsst*

---

### 17.171.7 Default

The keyword defaults are temp = 300, damp = 1, seed = 880302, f\_max=200.0 and N\_f = 100.

---

**(Dammak)** Dammak, Chalopin, Laroche, Hayoun, and Greffet, Phys Rev Lett, 103, 190601 (2009).

**(Barrat)** Barrat and Rodney, J. Stat. Phys, 144, 679 (2011).

## 17.172 fix reaxff/bonds command

Accelerator Variants: *reaxff/bonds/kk*

### 17.172.1 Syntax

```
fix ID group-ID reaxff/bonds Nevery filename
```

- ID, group-ID are documented in *fix* command
- reax/bonds = style name of this fix command
- Nevery = output interval in timesteps
- filename = name of output file

### 17.172.2 Examples

```
fix 1 all reaxff/bonds 100 bonds.reaxff
```



### 17.172.3 Description

Write out the bond information computed by the ReaxFF potential specified by *pair\_style reaxff* in the exact same format as the original stand-alone ReaxFF code of Adri van Duin. The bond information is written to *filename* on timesteps that are multiples of *Nevery*, including timestep 0. For time-averaged chemical species analysis, please see the *fix reaxff/species* command.

The specified group-ID is ignored by this fix.

The format of the output file should be reasonably self-explanatory. The meaning of the column header abbreviations is as follows:

- id = atom id
- type = atom type
- nb = number of bonds
- id\_1 = atom id of first bond
- id\_nb = atom id of Nth bond
- mol = molecule id
- bo\_1 = bond order of first bond
- bo\_nb = bond order of Nth bond
- abo = atom bond order (sum of all bonds)
- nlp = number of lone pairs
- q = atomic charge

If the filename ends with “.gz”, the output file is written in gzipped format. A gzipped dump file will be about 3x smaller than the text version, but will also take longer to write.

---

### 17.172.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 17.172.5 Restrictions

The fix reaxff/bonds command requires that the *pair\_style reaxff* is invoked. This fix is part of the REAXFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

To write gzipped bond files, you must compile LAMMPS with the `-DLAMMPS_GZIP` option.

### 17.172.6 Related commands

*pair\_style reaxff*, *fix reaxff/species*

### 17.172.7 Default

none

## 17.173 fix reaxff/species command

Accelerator Variants: *reaxff/species/kk*

### 17.173.1 Syntax

```
fix ID group-ID reaxff/species Nevery Nrepeat Nfreq filename keyword value ...
```

- ID, group-ID are documented in *fix* command
- reaxff/species = style name of this command
- Nevery = sample bond-order every this many timesteps
- Nrepeat = # of bond-order samples used for calculating averages
- Nfreq = calculate average bond-order every this many timesteps
- filename = name of output file
- zero or more keyword/value pairs may be appended
- keyword = *cutoff* or *element* or *position* or *delete*

*cutoff* value = I J Cutoff

I, J = atom types

Cutoff = Bond-order cutoff value for this pair of atom types

*element* value = Element1, Element2, ...

*position* value = posfreq filepos

posfreq = write position files every this many timestep

filepos = name of position output file

*delete* value = filedel keyword value

filedel = name of delete species output file

keyword = *specieslist* or *masslimit*

*specieslist* value = Nspecies Species1 Species2 ...

Nspecies = number of species in list

*masslimit* value = massmin massmax

massmin = minimum molecular weight of species to delete

massmax = maximum molecular weight of species to delete

## 17.173.2 Examples

```
fix 1 all reaxff/species 10 10 100 species.out
fix 1 all reaxff/species 1 2 20 species.out cutoff 1 1 0.40 cutoff 1 2 0.55
fix 1 all reaxff/species 1 100 100 species.out element Au O H position 1000 AuOH.pos
fix 1 all reaxff/species 1 100 100 species.out delete species.del masslimit 0 50
```

## 17.173.3 Description

Write out the chemical species information computed by the ReaxFF potential specified by *pair\_style reaxff*. Bond-order values (either averaged or instantaneous, depending on value of *Nrepeat*) are used to determine chemical bonds. Every *Nfreq* timesteps, chemical species information is written to *filename* as a two line output. The first line is a header containing labels. The second line consists of the following: timestep, total number of molecules, total number of distinct species, number of molecules of each species. In this context, “species” means a unique molecule. The chemical formula of each species is given in the first line.

**Warning:** In order to compute averaged data, it is required that there are no neighbor list rebuilds for at least  $Nrepeat * Nevery$  steps preceding each *Nfreq* step. For that reason, fix *reaxff/species* may change your neighbor list settings. Reneighboring will occur no more frequently than every  $Nrepeat * Nevery$  timesteps, and will occur less frequently if *Nfreq* is not a multiple of  $Nrepeat * Nevery$ . There will be a warning message showing the new settings. Having a *Nfreq* setting that is larger than what is required for correct computation of the ReaxFF force field interactions, in combination with certain *Nrepeat* and *Nevery* settings, can thus lead to incorrect results. For typical ReaxFF calculations, reneighboring only every 100 steps is already quite a low frequency.

If the filename ends with “.gz”, the output file is written in gzipped format. A gzipped dump file will be about 3x smaller than the text version, but will also take longer to write.

Optional keyword *cutoff* can be assigned to change the minimum bond-order values used in identifying chemical bonds between pairs of atoms. Bond-order cutoffs should be carefully chosen, as bond-order cutoffs that are too small may include too many bonds (which will result in an error), while cutoffs that are too large will result in fragmented molecules. The default cutoff of 0.3 usually gives good results.

The optional keyword *element* can be used to specify the chemical symbol printed for each LAMMPS atom type. The number of symbols must match the number of LAMMPS atom types and each symbol must consist of 1 or 2 alphanumeric characters. Normally, these symbols should be chosen to match the chemical identity of each LAMMPS atom type, as specified using the *reaxff pair\_coeff* command and the ReaxFF force field file.

The optional keyword *position* writes center-of-mass positions of each identified molecules to file *filepos* every *posfreq* timesteps. The first line contains information on timestep, total number of molecules, total number of distinct species, and box dimensions. The second line is a header containing labels. From the third line downward, each molecule writes a line of output containing the following information: molecule ID, number of atoms in this molecule, chemical formula, total charge, and center-of-mass xyz positions of this molecule. The xyz positions are in fractional coordinates relative to the box dimensions.

For the keyword *position*, the *filepos* is the name of the output file. It can contain the wildcard character “\*”. If the “\*” character appears in *filepos*, then one file per snapshot is written at *posfreq* and the “\*” character is replaced with the timestep value. For example, AuO.pos.\* becomes AuO.pos.0, AuO.pos.1000, etc.

The optional keyword *delete* enables the periodic removal of molecules from the system. Criteria for deletion can be either a list of specific chemical formulae or a range of molecular weights. Molecules are deleted every *Nfreq* timesteps, and bond connectivity is determined using the *Nevery* and *Nrepeat* keywords. The *filedel* argument is the name of the output file that records the species that are removed from the system. The *specieslist* keyword permits specific chemical species to be deleted. The *Nspecies* argument specifies how many species are eligible for deletion

and is followed by a list of chemical formulae, whose strings are compared to species identified by this fix. For example, “specieslist 2 CO CO2” deletes molecules that are identified as “CO” and “CO2” in the species output file. When using the *specieslist* keyword, the *filedel* file has the following format: the first line lists the chemical formulae eligible for deletion, and each additional line contains the timestep on which a molecule deletion occurs and the number of each species deleted on that timestep. The *masslimit* keyword permits deletion of molecules with molecular weights between *massmin* and *massmax*. When using the *masslimit* keyword, each line of the *filedel* file contains the timestep on which deletions occurs, followed by how many of each species are deleted (with quantities preceding chemical formulae). The *specieslist* and *masslimit* keywords cannot both be used in the same *reaxff/species* fix.

---

The *Nevery*, *Nrepeat*, and *Nfreq* arguments specify on what timesteps the bond-order values are sampled to get the average bond order. The species analysis is performed using the average bond-order on timesteps that are a multiple of *Nfreq*. The average is over *Nrepeat* bond-order samples, computed in the preceding portion of the simulation every *Nevery* timesteps. *Nfreq* must be a multiple of *Nevery* and *Nevery* must be non-zero even if *Nrepeat* is 1. Also, the timesteps contributing to the average bond-order cannot overlap, i.e.  $Nrepeat * Nevery$  can not exceed *Nfreq*.

For example, if *Nevery*=2, *Nrepeat*=6, and *Nfreq*=100, then bond-order values on timesteps 90,92,94,96,98,100 will be used to compute the average bond-order for the species analysis output on timestep 100.

---

#### 17.173.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes both a global vector of length 2 and a per-atom vector, either of which can be accessed by various *output commands*. The values in the global vector are “intensive”.

The 2 values in the global vector are as follows:

- 1 = total number of molecules
- 2 = total number of distinct species

The per-atom vector stores the molecule ID for each atom as identified by the fix. If an atom is not in a molecule, its ID will be 0. For atoms in the same molecule, the molecule ID for all of them will be the same and will be equal to the smallest atom ID of any atom in the molecule.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 17.173.5 Restrictions

The “fix reaxff/species” requires that *pair\_style reaxff* is used. This fix is part of the REAXFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

To write gzipped species files, you must compile LAMMPS with the `-DLAMMPS_GZIP` option.

### 17.173.6 Related commands

*pair\_style reaxff*, *fix reaxff/bonds*

### 17.173.7 Default

The default values for bond-order cutoffs are 0.3 for all I-J pairs. The default element symbols are C, H, O, N. Position files are not written by default.

## 17.174 fix recenter command

### 17.174.1 Syntax

```
fix ID group-ID recenter x y z keyword value ...
```

- ID, group-ID are documented in *fix* command
- recenter = style name of this fix command
- x,y,z = constrain center-of-mass to these coords (distance units), any coord can also be NULL or INIT (see below)
- zero or more keyword/value pairs may be appended
- keyword = *shift* or *units*

*shift* value = group-ID

group-ID = group of atoms whose coords are shifted

*units* value = *box* or *lattice* or *fraction*

### 17.174.2 Examples

```
fix 1 all recenter 0.0 0.5 0.0
fix 1 all recenter INIT INIT NULL
fix 1 all recenter INIT 0.0 0.0 units box
```

### 17.174.3 Description

Constrain the center-of-mass position of a group of atoms by adjusting the coordinates of the atoms every timestep. This is simply a small shift that does not alter the dynamics of the system or change the relative coordinates of any pair of atoms in the group. This can be used to insure the entire collection of atoms (or a portion of them) do not drift during the simulation due to random perturbations (e.g. *fix langevin* thermostating).

Distance units for the x,y,z values are determined by the setting of the *units* keyword, as discussed below. One or more x,y,z values can also be specified as NULL, which means exclude that dimension from this operation. Or it can be specified as INIT which means to constrain the center-of-mass to its initial value at the beginning of the run.

The center-of-mass (COM) is computed for the group specified by the *fix*. If the current COM is different than the specified x,y,z, then a group of atoms has their coordinates shifted by the difference. By default the shifted group is also the group specified by the *fix*. A different group can be shifted by using the *shift* keyword. For example, the COM could be computed on a protein to keep it in the center of the simulation box. But the entire system (protein + water) could be shifted.

If the *units* keyword is set to *box*, then the distance units of x,y,z are defined by the *units* command - e.g. Angstroms for *real* units. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing. A *fraction* value means a fractional distance between the lo/hi box boundaries, e.g. 0.5 = middle of the box. The default is to use lattice units.

Note that the *velocity* command can be used to create velocities with zero aggregate linear and/or angular momentum.

---

**Note:** This *fix* performs its operations at the same point in the timestep as other time integration fixes, such as *fix nve*, *fix nvt*, or *fix npt*. Thus *fix recenter* should normally be the last such *fix* specified in the input script, since the adjustments it makes to atom coordinates should come after the changes made by time integration. LAMMPS will warn you if your fixes are not ordered this way.

---

---

**Note:** If you use this *fix* on a small group of atoms (e.g. a molecule in solvent) without using the *shift* keyword to adjust the positions of all atoms in the system, then the results can be unpredictable. For example, if the molecule is pushed consistently in one direction by a flowing solvent, its velocity will increase. But its coordinates will be re-centered, meaning it is moved back towards the force. Thus over time, the velocity and effective temperature of the molecule could become very large, though it won't actually be moving due to the re-centering. If you are thermostating the entire system, then the solvent would be cooled to compensate. A better solution for this simulation scenario is to use the *fix spring* command to tether the molecule in place.

---

### 17.174.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this *fix* is written to *binary restart files*. None of the *fix\_modify* options are relevant to this *fix*.

This *fix* computes a global scalar which can be accessed by various *output commands*. The scalar is the distance the group is moved by *fix recenter*.

This *fix* also computes global 3-vector which can be accessed by various *output commands*. The 3 quantities in the vector are xyz components of displacement applied to the group of atoms by the *fix*.

The scalar and vector values calculated by this *fix* are “extensive”.

No parameter of this *fix* can be used with the *start/stop* keywords of the *run* command. This *fix* is not invoked during *energy minimization*.

## 17.174.5 Restrictions

This fix should not be used with an x,y,z setting that causes a large shift in the system on the first timestep, due to the requested COM being very different from the initial COM. This could cause atoms to be lost, especially in parallel. Instead, use the *displace\_atoms* command, which can be used to move atoms a large distance.

## 17.174.6 Related commands

*fix momentum, velocity*

## 17.174.7 Default

The option defaults are shift = fix group-ID, and units = lattice.

## 17.175 fix restrain command

### 17.175.1 Syntax

```
fix ID group-ID restrain keyword args ...
```

- ID, group-ID are documented in *fix* command
- restrain = style name of this fix command
- one or more keyword/arg pairs may be appended
- keyword = *bond* or *lbound* or *angle* or *dihedral*

```
bond args = atom1 atom2 Kstart Kstop r0start (r0stop)
  atom1,atom2 = IDs of 2 atoms in bond
  Kstart,Kstop = restraint coefficients at start/end of run (energy units)
  r0start = equilibrium bond distance at start of run (distance units)
  r0stop = equilibrium bond distance at end of run (optional) (distance units). If
↪not
  specified it is assumed to be equal to r0start
lbound args = atom1 atom2 Kstart Kstop r0start (r0stop)
  atom1,atom2 = IDs of 2 atoms in bond
  Kstart,Kstop = restraint coefficients at start/end of run (energy units)
  r0start = equilibrium bond distance at start of run (distance units)
  r0stop = equilibrium bond distance at end of run (optional) (distance units). If
↪not
  specified it is assumed to be equal to r0start
angle args = atom1 atom2 atom3 Kstart Kstop theta0
  atom1,atom2,atom3 = IDs of 3 atoms in angle, atom2 = middle atom
  Kstart,Kstop = restraint coefficients at start/end of run (energy units)
  theta0 = equilibrium angle theta (degrees)
dihedral args = atom1 atom2 atom3 atom4 Kstart Kstop phi0 keyword/value
  atom1,atom2,atom3,atom4 = IDs of 4 atoms in dihedral in linear order
  Kstart,Kstop = restraint coefficients at start/end of run (energy units)
  phi0 = equilibrium dihedral angle phi (degrees)
  keyword/value = optional keyword value pairs. supported keyword/value pairs:
  mult n = dihedral multiplicity n (integer >= 0, default = 1)
```

## 17.175.2 Examples

```

fix holdem all restrain bond 45 48 2000.0 2000.0 2.75
fix holdem all restrain lbound 45 48 2000.0 2000.0 2.75
fix holdem all restrain dihedral 1 2 3 4 2000.0 2000.0 120.0
fix holdem all restrain bond 45 48 2000.0 2000.0 2.75 dihedral 1 2 3 4 2000.0 2000.0 120.
↪0
fix texas_holdem all restrain dihedral 1 2 3 4 0.0 2000.0 120.0 dihedral 1 2 3 5 0.0 ↪
↪2000.0 -120.0 dihedral 1 2 3 6 0.0 2000.0 0.0

```

## 17.175.3 Description

Restrain the motion of the specified sets of atoms by making them part of a bond or angle or dihedral interaction whose strength can vary over time during a simulation. This is functionally similar to creating a bond or angle or dihedral for the same atoms in a data file, as specified by the *read\_data* command, albeit with a time-varying prefactor coefficient, and except for exclusion rules, as explained below.

For the purpose of force field parameter-fitting or mapping a molecular potential energy surface, this fix reduces the hassle and risk associated with modifying data files. In other words, use this fix to temporarily force a molecule to adopt a particular conformation. To create a permanent bond or angle or dihedral, you should modify the data file.

---

**Note:** Adding a bond/angle/dihedral with this command does not apply the exclusion rules and weighting factors specified by the *special\_bonds* command to atoms in the restraint that are now bonded (1-2,1-3,1-4 neighbors) as a result. If they are close enough to interact in a *pair\_style* sense (non-bonded interaction), then the bond/angle/dihedral restraint interaction will simply be superposed on top of that interaction.

---

The group-ID specified by this fix is ignored.

The second example above applies a restraint to hold the dihedral angle formed by atoms 1, 2, 3, and 4 near 120 degrees using a constant restraint coefficient. The fourth example applies similar restraints to multiple dihedral angles using a restraint coefficient that increases from 0.0 to 2000.0 over the course of the run.

---

**Note:** Adding a force to atoms implies a change in their potential energy as they move due to the applied force field. For dynamics via the *run* command, this energy can be added to the system's potential energy for thermodynamic output (see below). For energy minimization via the *minimize* command, this energy must be added to the system's potential energy to formulate a self-consistent minimization problem (see below).

---

In order for a restraint to be effective, the restraint force must typically be significantly larger than the forces associated with conventional force field terms. If the restraint is applied during a dynamics run (as opposed to during an energy minimization), a large restraint coefficient can significantly reduce the stable timestep size, especially if the atoms are initially far from the preferred conformation. You may need to experiment to determine what value of *K* works best for a given application.

For the case of finding a minimum energy structure for a single molecule with particular restraints (e.g. for fitting force field parameters or constructing a potential energy surface), commands such as the following may be useful:

```

# minimize molecule energy with restraints
velocity all create 600.0 8675309 mom yes rot yes dist gaussian
fix NVE all nve
fix TFIX all langevin 600.0 0.0 100 24601
fix REST all restrain dihedral 2 1 3 8 0.0 5000.0 ${angle1} dihedral 3 1 2 9 0.0 5000.0 $

```

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```

↪{angle2}
fix_modify REST energy yes
run 10000
fix TFIX all langevin 0.0 0.0 100 24601
fix REST all restrain dihedral 2 1 3 8 5000.0 5000.0 ${angle1} dihedral 3 1 2 9 5000.0.
↪5000.0 ${angle2}
fix_modify REST energy yes
run 10000
# sanity check for convergence
minimize 1e-6 1e-9 1000 1000000
# report unrestrained energies
unfix REST
run 0

```

The *bond* keyword applies a bond restraint to the specified atoms using the same functional form used by the *bond\_style harmonic* command. The potential associated with the restraint is

$$E = K(r - r_0)^2$$

with the following coefficients:

- $K$  (energy/distance<sup>2</sup>)
- $r_0$  (distance)

$K$  and  $r_0$  are specified with the *fix*. Note that the usual 1/2 factor is included in  $K$ .

The *lbond* keyword applies a lower bound bond restraint to the specified atoms using the same functional form used by the *bond\_style harmonic* command if the distance between the atoms is smaller than the equilibrium bond distance and 0 otherwise. The potential associated with the restraint is

$$E = K(r - r_0)^2, \text{ if } r < r_0$$

$$E = 0, \text{ if } r \geq r_0$$

with the following coefficients:

- $K$  (energy/distance<sup>2</sup>)
- $r_0$  (distance)

$K$  and  $r_0$  are specified with the *fix*. Note that the usual 1/2 factor is included in  $K$ .

The *angle* keyword applies an angle restraint to the specified atoms using the same functional form used by the *angle\_style harmonic* command. The potential associated with the restraint is

$$E = K(\theta - \theta_0)^2$$

with the following coefficients:

- $K$  (energy)
- $\theta_0$  (degrees)

$K$  and  $\theta_0$  are specified with the fix.  $\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence  $K$  is effectively energy per radian<sup>2</sup>. Note that the usual 1/2 factor is included in  $K$ .

---

The *dihedral* keyword applies a dihedral restraint to the specified atoms using a simplified form of the function used by the *dihedral\_style charmm* command. The potential associated with the restraint is

$$E = K[1 + \cos(n\phi - d)]$$

with the following coefficients:

- $K$  (energy)
- $n$  (multiplicity,  $\geq 0$ )
- $d$  (degrees) =  $\phi_0 + 180$

$K$  and  $\phi_0$  are specified with the fix. Note that the value of the dihedral multiplicity  $n$  is set by default to 1. You can use the optional *mult* keyword to set it to a different positive integer. Also note that the energy will be a minimum when the current dihedral angle  $\phi$  is equal to  $\phi_0$ .

---

#### 17.175.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy associated with this fix to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

---

**Note:** If you want the fictitious potential energy associated with the added forces to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

This fix computes a global scalar and a global vector of length 3, which can be accessed by various *output commands*. The scalar is the total potential energy for *all* the restraints as discussed above. The vector values are the sum of contributions to the following individual categories:

- 1 = bond energy
- 2 = angle energy
- 3 = dihedral energy

The scalar and vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

## 17.175.5 Restrictions

none

## 17.175.6 Related commands

none

## 17.175.7 Default

none

## 17.176 fix rhok command

### 17.176.1 Syntax

```
fix ID group-ID rhok nx ny nz K a
```

- ID, group-ID are documented in *fix* command
- nx, ny, nz = k-vector of collective density field
- K = spring constant of bias potential
- a = anchor point of bias potential

### 17.176.2 Examples

```
fix bias all rhok 16 0 0 4.0 16.0
fix 1 all npt temp 0.8 0.8 4.0 z 2.2 2.2 8.0
# output of 4 values from fix rhok: U_bias rho_k_RE rho_k_IM \|\rho_k\|
thermo_style custom step temp pzz lz f_bias f_bias[1] f_bias[2] f_bias[3]
```

### 17.176.3 Description

The fix applies a force to atoms given by the potential

$$U = \frac{1}{2}K(|\rho_{\vec{k}}| - a)^2$$

$$\rho_{\vec{k}} = \sum_j^N \exp(-i\vec{k} \cdot \vec{r}_j) / \sqrt{N}$$

$$\vec{k} = (2\pi n_x / L_x, 2\pi n_y / L_y, 2\pi n_z / L_z)$$

as described in (*Pedersen*).

This field, which biases configurations with long-range order, can be used to study crystal-liquid interfaces and determine melting temperatures (*Pedersen*).

An example of using the interface pinning method is located in the *examples/PACKAGES/rhok* directory.

### 17.176.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the potential energy calculated by the fix to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the potential energy discussed in the preceding paragraph. The scalar stored by this fix is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

This fix is not invoked during *energy minimization*.

### 17.176.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.176.6 Related commands

*thermo\_style*

### 17.176.7 Default

none

---

(Pedersen) Pedersen, J. Chem. Phys., 139, 104102 (2013).

## 17.177 fix rigid command

Accelerator Variants: *rigid/omp*

## 17.178 fix rigid/nve command

Accelerator Variants: *rigid/nve/omp*

## 17.179 fix rigid/nvt command

Accelerator Variants: *rigid/nvt/omp*

## 17.180 fix rigid/npt command

Accelerator Variants: *rigid/npt/omp*

## 17.181 fix rigid/nph command

Accelerator Variants: *rigid/nph/omp*

## 17.182 fix rigid/small command

Accelerator Variants: *rigid/small/omp*

## 17.183 fix rigid/nve/small command

## 17.184 fix rigid/nvt/small command

## 17.185 fix rigid/npt/small command

## 17.186 fix rigid/nph/small command

### 17.186.1 Syntax

```
fix ID group-ID style bodystyle args keyword values ...
```

- ID, group-ID are documented in *fix* command
- style = *rigid* or *rigid/nve* or *rigid/nvt* or *rigid/npt* or *rigid/nph* or *rigid/small* or *rigid/nve/small* or *rigid/nvt/small* or *rigid/npt/small* or *rigid/nph/small*
- bodystyle = *single* or *molecule* or *group*

*single* args = none

*molecule* args = none

*custom* args = *i\_propname* or *v\_varname*

*i\_propname* = a custom integer vector defined via *fix property/atom*

*v\_varname* = an atom-style or atomfile-style variable

*group* args = N groupID1 groupID2 ...

N = # of groups

groupID1, groupID2, ... = list of N group IDs

- zero or more keyword/value pairs may be appended
- keyword = *langevin* or *reinit* or *temp* or *iso* or *aniso* or *x* or *y* or *z* or *couple* or *tparam* or *pchain* or *dilate* or *force* or *torque* or *infile*

*langevin* values = Tstart Tstop Tperiod seed

Tstart, Tstop = desired temperature at start/stop of run (temperature units)

Tdamp = temperature damping parameter (time units)

```

    seed = random number seed to use for white noise (positive integer)
reinit value = yes or no
temp values = Tstart Tstop Tdamp
    Tstart,Tstop = desired temperature at start/stop of run (temperature units)
    Tdamp = temperature damping parameter (time units)
iso or aniso values = Pstart Pstop Pdamp
    Pstart,Pstop = scalar external pressure at start/end of run (pressure units)
    Pdamp = pressure damping parameter (time units)
x or y or z values = Pstart Pstop Pdamp
    Pstart,Pstop = external stress tensor component at start/end of run (pressure
↪units)
    Pdamp = stress damping parameter (time units)
couple value = none or xyz or xy or yz or xz
tparam values = Tchain Titer Torder
    Tchain = length of Nose/Hoover thermostat chain
    Titer = number of thermostat iterations performed
    Torder = 3 or 5 = Yoshida-Suzuki integration parameters
pchain values = Pchain
    Pchain = length of the Nose/Hoover thermostat chain coupled with the barostat
dilate value = dilate-group-ID
    dilate-group-ID = only dilate atoms in this group due to barostat volume changes
force values = M xflag yflag zflag
    M = which rigid body from 1-Nbody (see asterisk form below)
    xflag,yflag,zflag = off/on if component of center-of-mass force is active
torque values = M xflag yflag zflag
    M = which rigid body from 1-Nbody (see asterisk form below)
    xflag,yflag,zflag = off/on if component of center-of-mass torque is active
infile filename
    filename = file with per-body values of mass, center-of-mass, moments of inertia
mol value = template-ID
    template-ID = ID of molecule template specified in a separate molecule command

```

## 17.186.2 Examples

```

fix 1 clump rigid single reinit yes
fix 1 clump rigid/small molecule
fix 1 clump rigid single force 1 off off on langevin 1.0 1.0 1.0 428984
fix 1 polychains rigid/nvt molecule temp 1.0 1.0 5.0 reinit no
fix 1 polychains rigid molecule force 1*5 off off off force 6*10 off off on
fix 1 polychains rigid/small molecule langevin 1.0 1.0 1.0 428984
fix 2 fluid rigid group 3 clump1 clump2 clump3 torque * off off off
fix 1 rods rigid/npt molecule temp 300.0 300.0 100.0 iso 0.5 0.5 10.0
fix 1 particles rigid/npt molecule temp 1.0 1.0 5.0 x 0.5 0.5 1.0 z 0.5 0.5 1.0 couple xz
fix 1 water rigid/nph molecule iso 0.5 0.5 1.0
fix 1 particles rigid/npt/small molecule temp 1.0 1.0 1.0 iso 0.5 0.5 1.0

variable bodyid atom 1.0*gmask(clump1)+2.0*gmask(clump2)+3.0*gmask(clump3)
fix 1 clump rigid custom v_bodyid

variable bodyid atomfile bodies.txt
fix 1 clump rigid custom v_bodyid

```

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```
fix 0 all property/atom i_bodyid
read_restart data.rigid fix 0 NULL Bodies
fix 1 clump rigid/small custom i_bodyid
```

### 17.186.3 Description

Treat one or more sets of atoms as independent rigid bodies. This means that each timestep the total force and torque on each rigid body is computed as the sum of the forces and torques on its constituent particles. The coordinates, velocities, and orientations of the atoms in each body are then updated so that the body moves and rotates as a single entity. This is implemented by creating internal data structures for each rigid body and performing time integration on these data structures. Positions, velocities, and orientations of the constituent particles are regenerated from the rigid body data structures in every time step. This restricts which operations and fixes can be applied to rigid bodies. See below for a detailed discussion.

Examples of large rigid bodies are a colloidal particle, or portions of a biomolecule such as a protein.

Example of small rigid bodies are patchy nanoparticles, such as those modeled in [this paper](#) by Sharon Glotzer's group, clumps of granular particles, lipid molecules consisting of one or more point dipoles connected to other spheroids or ellipsoids, irregular particles built from line segments (2d) or triangles (3d), and coarse-grain models of nano or colloidal particles consisting of a small number of constituent particles. Note that the *fix shake* command can also be used to rigidify small molecules of 2, 3, or 4 atoms, e.g. water molecules. That fix treats the constituent atoms as point masses.

These fixes also update the positions and velocities of the atoms in each rigid body via time integration, in the NVE, NVT, NPT, or NPH ensemble, as described below.

There are two main variants of this fix, *fix rigid* and *fix rigid/small*. The NVE/NVT/NPT/NHT versions belong to one of the two variants, as their style names indicate.

---

**Note:** Not all of the *bodystyle* options and keyword/value options are available for both the *rigid* and *rigid/small* variants. See details below.

---

The *rigid* styles are typically the best choice for a system with a small number of large rigid bodies, each of which can extend across the domain of many processors. It operates by creating a single global list of rigid bodies, which all processors contribute to. MPI\_Allreduce operations are performed each timestep to sum the contributions from each processor to the force and torque on all the bodies. This operation will not scale well in parallel if large numbers of rigid bodies are simulated.

The *rigid/small* styles are typically best for a system with a large number of small rigid bodies. Each body is assigned to the atom closest to the geometrical center of the body. The fix operates using local lists of rigid bodies owned by each processor and information is exchanged and summed via local communication between neighboring processors when ghost atom info is accumulated.

---

**Note:** To use the *rigid/small* styles the ghost atom cutoff must be large enough to span the distance between the atom that owns the body and every other atom in the body. This distance value is printed out when the rigid bodies are defined. If the *pair\_style* cutoff plus neighbor skin does not span this distance, then you should use the *comm\_modify cutoff* command with a setting epsilon larger than the distance.

---

Which of the two variants is faster for a particular problem is hard to predict. The best way to decide is to perform a short test run. Both variants should give identical numerical answers for short runs. Long runs should give statistically similar results, but round-off differences may accumulate to produce divergent trajectories.

**Note:** You should not update the atoms in rigid bodies via other time-integration fixes (e.g. *fix nve*, *fix nvt*, *fix npt*, *fix move*), or you will have conflicting updates to positions and velocities resulting in unphysical behavior in most cases. When performing a hybrid simulation with some atoms in rigid bodies, and some not, a separate time integration fix like *fix nve* or *fix nvt* should be used for the non-rigid particles.

---

**Note:** These fixes are overkill if you simply want to hold a collection of atoms stationary or have them move with a constant velocity. A simpler way to hold atoms stationary is to not include those atoms in your time integration fix. E.g. use “fix 1 mobile nve” instead of “fix 1 all nve”, where “mobile” is the group of atoms that you want to move. You can move atoms with a constant velocity by assigning them an initial velocity (via the *velocity* command), setting the force on them to 0.0 (via the *fix setforce* command), and integrating them as usual (e.g. via the *fix nve* command).

---

**Warning:** The aggregate properties of each rigid body are calculated at the start of a simulation run and are maintained in internal data structures. The properties include the position and velocity of the center-of-mass of the body, its moments of inertia, and its angular momentum. This is done using the properties of the constituent atoms of the body at that point in time (or see the *infile* keyword option). Thereafter, changing these properties of individual atoms in the body will have no effect on a rigid body’s dynamics, unless they effect any computation of per-atom forces or torques. If the keyword *reinit* is set to *yes* (the default), the rigid body data structures will be recreated at the beginning of each *run* command; if the keyword *reinit* is set to *no*, the rigid body data structures will be built only at the very first *run* command and maintained for as long as the rigid fix is defined. For example, you might think you could displace the atoms in a body or add a large velocity to each atom in a body to make it move in a desired direction before a second run is performed, using the *set* or *displace\_atoms* or *velocity* commands. But these commands will not affect the internal attributes of the body unless *reinit* is set to *yes*. With *reinit* set to *no* (or using the *infile* option, which implies *reinit no*) the position and velocity of individual atoms in the body will be reset when time integration starts again.

---

Each rigid body must have two or more atoms. An atom can belong to at most one rigid body. Which atoms are in which bodies can be defined via several options.

---

**Note:** With the *rigid/small* styles, which require that *bodystyle* be specified as *molecule* or *custom*, you can define a system that has no rigid bodies initially. This is useful when you are using the *mol* keyword in conjunction with another fix that is adding rigid bodies on-the-fly as molecules, such as *fix deposit* or *fix pour*.

---

For *bodystyle single* the entire fix group of atoms is treated as one rigid body. This option is only allowed for the *rigid* styles.

For *bodystyle molecule*, atoms are grouped into rigid bodies by their respective molecule IDs: each set of atoms in the fix group with the same molecule ID is treated as a different rigid body. This option is allowed for both the *rigid* and *rigid/small* styles. Note that atoms with a molecule ID = 0 will be treated as a single rigid body. For a system with atomic solvent (typically this is atoms with molecule ID = 0) surrounding rigid bodies, this may not be what you want. Thus you should be careful to use a fix group that only includes atoms you want to be part of rigid bodies.

*Bodystyle custom* is similar to *bodystyle molecule* except that it is more flexible in using other per-atom properties to define the sets of atoms that form rigid bodies. A custom per-atom integer vector defined by the *fix property/atom* command can be used. Or an *atom-style* or *atomfile-style variable* can be used; the floating-point value produced by the variable is rounded to an integer. As with *bodystyle molecule*, each set of atoms in the fix groups with the same integer value is treated as a different rigid body. Since fix *property/atom* custom vectors and *atom-style* variables produce values for all atoms, you should be careful to use a fix group that only includes atoms you want to be part of rigid bodies.



**Note:** To compute the initial center-of-mass position and other properties of each rigid body, the image flags for each atom in the body are used to “unwrap” the atom coordinates. Thus you must insure that these image flags are consistent so that the unwrapping creates a valid rigid body (one where the atoms are close together), particularly if the atoms in a single rigid body straddle a periodic boundary. This means the input data file or restart file must define the image flags for each atom consistently or that you have used the *set* command to specify them correctly. If a dimension is non-periodic then the image flag of each atom must be 0 in that dimension, else an error is generated.

---

The *force* and *torque* keywords discussed next are only allowed for the *rigid* styles.

By default, each rigid body is acted on by other atoms which induce an external force and torque on its center of mass, causing it to translate and rotate. Components of the external center-of-mass force and torque can be turned off by the *force* and *torque* keywords. This may be useful if you wish a body to rotate but not translate, or vice versa, or if you wish it to rotate or translate continuously unaffected by interactions with other particles. Note that if you expect a rigid body not to move or rotate by using these keywords, you must insure its initial center-of-mass translational or angular velocity is 0.0. Otherwise the initial translational or angular momentum the body has will persist.

An *xflag*, *yflag*, or *zflag* set to *off* means turn off the component of force or torque in that dimension. A setting of *on* means turn on the component, which is the default. Which rigid body(s) the settings apply to is determined by the first argument of the *force* and *torque* keywords. It can be an integer *M* from 1 to *Nbody*, where *Nbody* is the number of rigid bodies defined. A wild-card asterisk can be used in place of, or in conjunction with, the *M* argument to set the flags for multiple rigid bodies. This takes the form “\*” or “\**n*” or “*n*\*” or “*m*\**n*”. If *N* = the number of rigid bodies, then an asterisk with no numeric values means all bodies from 1 to *N*. A leading asterisk means all bodies from 1 to *n* (inclusive). A trailing asterisk means all bodies from *n* to *N* (inclusive). A middle asterisk means all types from *m* to *n* (inclusive). Note that you can use the *force* or *torque* keywords as many times as you like. If a particular rigid body has its component flags set multiple times, the settings from the final keyword are used.

**Note:** For computational efficiency, you may wish to turn off pairwise and bond interactions within each rigid body, as they no longer contribute to the motion. The *neigh\_modify exclude* and *delete\_bonds* commands are used to do this. If the rigid bodies have strongly overlapping atoms, you may need to turn off these interactions to avoid numerical problems due to large equal/opposite intra-body forces swamping the contribution of small inter-body forces.

---

For computational efficiency, you should typically define one *fix rigid* or *fix rigid/small* command which includes all the desired rigid bodies. LAMMPS will allow multiple rigid fixes to be defined, but it is more expensive.

---

The constituent particles within a rigid body can be point particles (the default in LAMMPS) or finite-size particles, such as spheres or ellipsoids or line segments or triangles. See the *atom\_style sphere and ellipsoid and line and tri* commands for more details on these kinds of particles. Finite-size particles contribute differently to the moment of inertia of a rigid body than do point particles. Finite-size particles can also experience torque (e.g. due to *frictional granular interactions*) and have an orientation. These contributions are accounted for by these fixes.

Forces between particles within a body do not contribute to the external force or torque on the body. Thus for computational efficiency, you may wish to turn off pairwise and bond interactions between particles within each rigid body. The *neigh\_modify exclude* and *delete\_bonds* commands are used to do this. For finite-size particles this also means the particles can be highly overlapped when creating the rigid body.

---

The *rigid*, *rigid/nve*, *rigid/small*, and *rigid/small/nve* styles perform constant NVE time integration. They are referred to below as the 4 NVE rigid styles. The only difference is that the *rigid* and *rigid/small* styles use an integration technique based on Richardson iterations. The *rigid/nve* and *rigid/small/nve* styles uses the methods described in the paper by *Miller*, which are thought to provide better energy conservation than an iterative approach.

The *rigid/nvt* and *rigid/nvt/small* styles performs constant NVT integration using a Nose/Hoover thermostat with chains as described originally in (*Hoover*) and (*Martyna*), which thermostats both the translational and rotational degrees of

freedom of the rigid bodies. They are referred to below as the 2 NVT rigid styles. The rigid-body algorithm used by *rigid/nvt* is described in the paper by *Kamberaj*.

The *rigid/npt*, *rigid/nph*, *rigid/npt/small*, and *rigid/nph/small* styles perform constant NPT or NPH integration using a Nose/Hoover barostat with chains. They are referred to below as the 4 NPT and NPH rigid styles. For the NPT case, the same Nose/Hoover thermostat is also used as with *rigid/nvt* and *rigid/nvt/small*.

The barostat parameters are specified using one or more of the *iso*, *aniso*, *x*, *y*, *z* and *couple* keywords. These keywords give you the ability to specify 3 diagonal components of the external stress tensor, and to couple these components together so that the dimensions they represent are varied together during a constant-pressure simulation. The effects of these keywords are similar to those defined in *fix npt/nph*

---

**Note:** Currently the *rigid/npt*, *rigid/nph*, *rigid/npt/small*, and *rigid/nph/small* styles do not support triclinic (non-orthogonal) boxes.

---

The target pressures for each of the 6 components of the stress tensor can be specified independently via the *x*, *y*, *z* keywords, which correspond to the 3 simulation box dimensions. For each component, the external pressure or tensor component at each timestep is a ramped value during the run from *Pstart* to *Pstop*. If a target pressure is specified for a component, then the corresponding box dimension will change during a simulation. For example, if the *y* keyword is used, the *y*-box length will change. A box dimension will not change if that component is not specified, although you have the option to change that dimension via the *fix deform* command.

For all barostat keywords, the *Pdamp* parameter operates like the *Tdamp* parameter, determining the time scale on which pressure is relaxed. For example, a value of 10.0 means to relax the pressure in a timespan of (roughly) 10 time units (e.g.  $\tau$  or fs or ps - see the *units* command).

Regardless of what atoms are in the *fix* group (the only atoms which are time integrated), a global pressure or stress tensor is computed for all atoms. Similarly, when the size of the simulation box is changed, all atoms are re-scaled to new positions, unless the keyword *dilate* is specified with a *dilate-group-ID* for a group that represents a subset of the atoms. This can be useful, for example, to leave the coordinates of atoms in a solid substrate unchanged and controlling the pressure of a surrounding fluid. Another example is a system consisting of rigid bodies and point particles where the barostat is only coupled with the rigid bodies. This option should be used with care, since it can be unphysical to dilate some atoms and not others, because it can introduce large, instantaneous displacements between a pair of atoms (one dilated, one not) that are far from the dilation origin.

The *couple* keyword allows two or three of the diagonal components of the pressure tensor to be “coupled” together. The value specified with the keyword determines which are coupled. For example, *xz* means the *Pxx* and *Pzz* components of the stress tensor are coupled. *xyz* means all 3 diagonal components are coupled. Coupling means two things: the instantaneous stress will be computed as an average of the corresponding diagonal components, and the coupled box dimensions will be changed together in lockstep, meaning coupled dimensions will be dilated or contracted by the same percentage every timestep. The *Pstart*, *Pstop*, *Pdamp* parameters for any coupled dimensions must be identical. *Couple xyz* can be used for a 2d simulation; the *z* dimension is simply ignored.

The *iso* and *aniso* keywords are simply shortcuts that are equivalent to specifying several other keywords together.

The keyword *iso* means couple all 3 diagonal components together when pressure is computed (hydrostatic pressure), and dilate/contract the dimensions together. Using “iso Pstart Pstop Pdamp” is the same as specifying these 4 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
couple xyz
```

The keyword *aniso* means *x*, *y*, and *z* dimensions are controlled independently using the *Pxx*, *Pyy*, and *Pzz* components of the stress tensor as the driving forces, and the specified scalar external pressure. Using “aniso Pstart Pstop Pdamp” is the same as specifying these 4 keywords:

```
x Pstart Pstop Pdamp
y Pstart Pstop Pdamp
z Pstart Pstop Pdamp
couple none
```

The keyword/value option pairs are used in the following ways.

The *reinit* keyword determines, whether the rigid body properties are re-initialized between run commands. With the option *yes* (the default) this is done, with the option *no* this is not done. Turning off the re-initialization can be helpful to protect rigid bodies against unphysical manipulations between runs or when properties cannot be easily re-computed (e.g. when read from a file). When using the *infile* keyword, the *reinit* option is automatically set to *no*.

The *langevin* and *temp* and *tparam* keywords perform thermostating of the rigid bodies, altering both their translational and rotational degrees of freedom. What is meant by “temperature” of a collection of rigid bodies and how it can be monitored via the fix output is discussed below.

The *langevin* keyword applies a Langevin thermostat to the constant NVE time integration performed by any of the 4 NVE rigid styles: *rigid*, *rigid/nve*, *rigid/small*, *rigid/small/nve*. It cannot be used with the 2 NVT rigid styles: *rigid/nvt*, *rigid/small/nvt*. The desired temperature at each timestep is a ramped value during the run from *Tstart* to *Tstop*. The *Tdamp* parameter is specified in time units and determines how rapidly the temperature is relaxed. For example, a value of 100.0 means to relax the temperature in a timespan of (roughly) 100 time units ( $\tau$  or fs or ps - see the *units* command). The random # *seed* must be a positive integer.

The way that Langevin thermostating operates is explained on the [fix langevin](#) doc page. If you wish to simply viscously damp the rotational motion without thermostating, you can set *Tstart* and *Tstop* to 0.0, which means only the viscous drag term in the Langevin thermostat will be applied. See the discussion on the [fix viscous](#) page for details.

---

**Note:** When the *langevin* keyword is used with fix rigid versus fix rigid/small, different dynamics will result for parallel runs. This is because of the way random numbers are used in the two cases. The dynamics for the two cases should be statistically similar, but will not be identical, even for a single timestep.

---

The *temp* and *tparam* keywords apply a Nose/Hoover thermostat to the NVT time integration performed by the 2 NVT rigid styles. They cannot be used with the 4 NVE rigid styles. The desired temperature at each timestep is a ramped value during the run from *Tstart* to *Tstop*. The *Tdamp* parameter is specified in time units and determines how rapidly the temperature is relaxed. For example, a value of 100.0 means to relax the temperature in a timespan of (roughly) 100 time units (tau or fs or ps - see the *units* command).

Nose/Hoover chains are used in conjunction with this thermostat. The *tparam* keyword can optionally be used to change the chain settings used. *Tchain* is the number of thermostats in the Nose Hoover chain. This value, along with *Tdamp* can be varied to dampen undesirable oscillations in temperature that can occur in a simulation. As a rule of thumb, increasing the chain length should lead to smaller oscillations. The keyword *pchain* specifies the number of thermostats in the chain thermostating the barostat degrees of freedom.

---

**Note:** There are alternate ways to thermostat a system of rigid bodies. You can use [fix langevin](#) to treat the individual particles in the rigid bodies as effectively immersed in an implicit solvent, e.g. a Brownian dynamics model. For hybrid systems with both rigid bodies and solvent particles, you can thermostat only the solvent particles that surround one or more rigid bodies by appropriate choice of groups in the compute and fix commands for temperature and thermostating. The solvent interactions with the rigid bodies should then effectively thermostat the rigid body temperature as well without use of the Langevin or Nose/Hoover options associated with the fix rigid commands.

---

The *mol* keyword can only be used with the *rigid/small* styles. It must be used when other commands, such as [fix deposit](#) or [fix pour](#), add rigid bodies on-the-fly during a simulation. You specify a *template-ID* previously defined using the

*molecule* command, which reads a file that defines the molecule. You must use the same *template-ID* that the other fix which is adding rigid bodies uses. The coordinates, atom types, atom diameters, center-of-mass, and moments of inertia can be specified in the molecule file. See the *molecule* command for details. The only settings required to be in this file are the coordinates and types of atoms in the molecule, in which case the molecule command calculates the other quantities itself.

Note that these other fixes create new rigid bodies, in addition to those defined initially by this fix via the *bodystyle* setting.

Also note that when using the *mol* keyword, extra restart information about all rigid bodies is written out whenever a restart file is written out. See the NOTE in the next section for details.

---

The *infile* keyword allows a file of rigid body attributes to be read in from a file, rather than having LAMMPS compute them. There are 5 such attributes: the total mass of the rigid body, its center-of-mass position, its 6 moments of inertia, its center-of-mass velocity, and the 3 image flags of the center-of-mass position. For rigid bodies consisting of point particles or non-overlapping finite-size particles, LAMMPS can compute these values accurately. However, for rigid bodies consisting of finite-size particles which overlap each other, LAMMPS will ignore the overlaps when computing these 4 attributes. The amount of error this induces depends on the amount of overlap. To avoid this issue, the values can be pre-computed (e.g. using Monte Carlo integration).

The format of the file is as follows. Note that the file does not have to list attributes for every rigid body integrated by fix rigid. Only bodies which the file specifies will have their computed attributes overridden. The file can contain initial blank lines or comment lines starting with “#” which are ignored. The first non-blank, non-comment line should list N = the number of lines to follow. The N successive lines contain the following information:

```
ID1 masstotal xcm ycm zcm ix x iy izz ix y ixz iy z vxcm vy cm vzcm lx ly lz ixcm iy cm izcm
ID2 masstotal xcm ycm zcm ix x iy izz ix y ixz iy z vxcm vy cm vzcm lx ly lz ixcm iy cm izcm
...
IDN masstotal xcm ycm zcm ix x iy izz ix y ixz iy z vxcm vy cm vzcm lx ly lz ixcm iy cm izcm
```

The rigid body IDs are all positive integers. For the *single* bodystyle, only an ID of 1 can be used. For the *group* bodystyle, IDs from 1 to Ng can be used where Ng is the number of specified groups. For the *molecule* bodystyle, use the molecule ID for the atoms in a specific rigid body as the rigid body ID.

The masstotal and center-of-mass coordinates (xcm,ycm,zcm) are self-explanatory. The center-of-mass should be consistent with what is calculated for the position of the rigid body with all its atoms unwrapped by their respective image flags. If this produces a center-of-mass that is outside the simulation box, LAMMPS wraps it back into the box.

The 6 moments of inertia (ixx,iyy,izz,ixy,ixz,iyz) should be the values consistent with the current orientation of the rigid body around its center of mass. The values are with respect to the simulation box XYZ axes, not with respect to the principal axes of the rigid body itself. LAMMPS performs the latter calculation internally.

The (vxcm,vy cm,vzcm) values are the velocity of the center of mass. The (lx,ly,lz) values are the angular momentum of the body. The (vxcm,vy cm,vzcm) and (lx,ly,lz) values can simply be set to 0 if you wish the body to have no initial motion.

The (ixcm,iy cm,izcm) values are the image flags of the center of mass of the body. For periodic dimensions, they specify which image of the simulation box the body is considered to be in. An image of 0 means it is inside the box as defined. A value of 2 means add 2 box lengths to get the true value. A value of -1 means subtract 1 box length to get the true value. LAMMPS updates these flags as the rigid bodies cross periodic boundaries during the simulation.

---

**Note:** If you use the *infile* or *mol* keywords and write restart files during a simulation, then each time a restart file is written, the fix also write an auxiliary restart file with the name rfile.rigid, where “rfile” is the name of the restart file, e.g. tmp.restart.10000 and tmp.restart.10000.rigid. This auxiliary file is in the same format described above. Thus it can be used in a new input script that restarts the run and re-specifies a rigid fix using an *infile* keyword and the

appropriate filename. Note that the auxiliary file will contain one line for every rigid body, even if the original file only listed a subset of the rigid bodies.

---

If you use a *temperature compute* with a group that includes particles in rigid bodies, the degrees-of-freedom removed by each rigid body are accounted for in the temperature (and pressure) computation, but only if the temperature group includes all the particles in a particular rigid body.

A 3d rigid body has 6 degrees of freedom (3 translational, 3 rotational), except for a collection of point particles lying on a straight line, which has only 5, e.g a dimer. A 2d rigid body has 3 degrees of freedom (2 translational, 1 rotational).

---

**Note:** You may wish to explicitly subtract additional degrees-of-freedom if you use the *force* and *torque* keywords to eliminate certain motions of one or more rigid bodies. LAMMPS does not do this automatically.

---

The rigid body contribution to the pressure of the system (virial) is also accounted for by this fix.

---

If your simulation is a hybrid model with a mixture of rigid bodies and non-rigid particles (e.g. solvent) there are several ways these rigid fixes can be used in tandem with *fix nve*, *fix nvt*, *fix npt*, and *fix nph*.

If you wish to perform NVE dynamics (no thermostating or barostating), use one of 4 NVE rigid styles to integrate the rigid bodies, and *fix nve* to integrate the non-rigid particles.

If you wish to perform NVT dynamics (thermostating, but no barostating), you can use one of the 2 NVT rigid styles for the rigid bodies, and any thermostating fix for the non-rigid particles (*fix nvt*, *fix langevin*, *fix temp/berendsen*). You can also use one of the 4 NVE rigid styles for the rigid bodies and thermostat them using *fix langevin* on the group that contains all the particles in the rigid bodies. The net force added by *fix langevin* to each rigid body effectively thermostats its translational center-of-mass motion. Not sure how well it does at thermostating its rotational motion.

If you wish to perform NPT or NPH dynamics (barostating), you cannot use both *fix npt* and the NPT or NPH rigid styles. This is because there can only be one fix which monitors the global pressure and changes the simulation box dimensions. So you have 3 choices:

- Use one of the 4 NPT or NPH styles for the rigid bodies. Use the *dilate* all option so that it will dilate the positions of the non-rigid particles as well. Use *fix nvt* (or any other thermostat) for the non-rigid particles.
- Use *fix npt* for the group of non-rigid particles. Use the *dilate* all option so that it will dilate the center-of-mass positions of the rigid bodies as well. Use one of the 4 NVE or 2 NVT rigid styles for the rigid bodies.
- Use *fix press/berendsen* to compute the pressure and change the box dimensions. Use one of the 4 NVE or 2 NVT rigid styles for the rigid bodies. Use *fix nvt* (or any other thermostat) for the non-rigid particles.

In all case, the rigid bodies and non-rigid particles both contribute to the global pressure and the box is scaled the same by any of the barostating fixes.

You could even use the second and third options for a non-hybrid simulation consisting of only rigid bodies, assuming you give *fix npt* an empty group, though it's an odd thing to do. The barostating fixes (*fix npt* and *fix press/berendsen*) will monitor the pressure and change the box dimensions, but not time integrate any particles. The integration of the rigid bodies will be performed by *fix rigid/nvt*.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

---

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 17.186.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about the 4 NVE rigid styles is written to *binary restart files*. The exception is if the *infile* or *mol* keyword is used, in which case an auxiliary file is written out with rigid body information each time a restart file is written, as explained above for the *infile* keyword. For the 2 NVT rigid styles, the state of the Nose/Hoover thermostat is written to *binary restart files*. Ditto for the 4 NPT and NPH rigid styles, and the state of the Nose/Hoover barostat. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* and *press* options are supported by the 4 NPT and NPH rigid styles to change the computes used to calculate the instantaneous pressure tensor. Note that the 2 NVT rigid fixes do not use any external compute to compute instantaneous temperature.

The *fix\_modify bodyforces* option is supported by all rigid styles to set whether per-body forces and torques are computed early or late in a timestep, i.e. at the post-force stage or at the final-integrate stage or the timestep, respectively.

The cumulative energy change in the system imposed by the 6 NVT, NPT, NPH rigid fixes, via either thermostating and/or barostating, is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

The 2 NVE rigid fixes compute a global scalar which can be accessed by various *output commands*. The scalar value calculated by these fixes is “intensive”. The scalar is the current temperature of the collection of rigid bodies. This is averaged over all rigid bodies and their translational and rotational degrees of freedom. The translational energy of a rigid body is  $1/2 m v^2$ , where  $m$  = total mass of the body and  $v$  = the velocity of its center of mass. The rotational energy of a rigid body is  $1/2 I w^2$ , where  $I$  = the moment of inertia tensor of the body and  $w$  = its angular velocity. Degrees of freedom constrained by the *force* and *torque* keywords are removed from this calculation, but only for the *rigid* and *rigid/nve* fixes.

The 6 NVT, NPT, NPH rigid fixes compute a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to these fixes described above. The scalar value calculated by this fix is “extensive”.

The *fix\_modify virial* option is supported by these fixes to add the contribution due to the added forces on atoms to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial yes*.

All of the *rigid* styles (not the *rigid/small* styles) compute a global array of values which can be accessed by various *output commands*. Similar information about the bodies defined by the *rigid/small* styles can be accessed via the *compute rigid/local* command.

The number of rows in the array is equal to the number of rigid bodies. The number of columns is 15. Thus for each rigid body, 15 values are stored: the xyz coords of the center of mass (COM), the xyz components of the COM velocity, the xyz components of the force acting on the COM, the xyz components of the torque acting on the COM, and the xyz image flags of the COM.

The center of mass (COM) for each body is similar to unwrapped coordinates written to a dump file. It will always be inside (or slightly outside) the simulation box. The image flags have the same meaning as image flags for atom positions (see the “dump” command). This means you can calculate the unwrapped COM by applying the image flags to the COM, the same as when unwrapped coordinates are written to a dump file.

The force and torque values in the array are not affected by the *force* and *torque* keywords in the *fix rigid* command; they reflect values before any changes are made by those keywords.

The ordering of the rigid bodies (by row in the array) is as follows. For the *single* keyword there is just one rigid body. For the *molecule* keyword, the bodies are ordered by ascending molecule ID. For the *group* keyword, the list of group IDs determines the ordering of bodies.

The array values calculated by these fixes are “intensive”, meaning they are independent of the number of atoms in the simulation.

No parameter of these fixes can be used with the *start/stop* keywords of the *run* command. These fixes are not invoked during *energy minimization*.

## 17.186.5 Restrictions

These fixes are all part of the RIGID package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Assigning a temperature via the *velocity create* command to a system with *rigid bodies* may not have the desired outcome for two reasons. First, the velocity command can be invoked before the rigid-body fix is invoked or initialized and the number of adjusted degrees of freedom (DOFs) is known. Thus it is not possible to compute the target temperature correctly. Second, the assigned velocities may be partially canceled when constraints are first enforced, leading to a different temperature than desired. A workaround for this is to perform a *run 0* command, which insures all DOFs are accounted for properly, and then rescale the temperature to the desired value before performing a simulation. For example:

```
velocity all create 300.0 12345
run 0 # temperature may not be 300K
velocity all scale 300.0 # now it should be
```

## 17.186.6 Related commands

*delete\_bonds*, *neigh\_modify exclude*, *fix shake*

## 17.186.7 Default

The option defaults are force \* on on on and torque \* on on on, meaning all rigid bodies are acted on by center-of-mass force and torque. Also Tchain = Pchain = 10, Titer = 1, Torder = 3, reinit = yes.

**(Hoover)** Hoover, Phys Rev A, 31, 1695 (1985).

**(Kamberaj)** Kamberaj, Low, Neal, J Chem Phys, 122, 224114 (2005).

**(Martyna)** Martyna, Klein, Tuckerman, J Chem Phys, 97, 2635 (1992); Martyna, Tuckerman, Tobias, Klein, Mol Phys, 87, 1117.

**(Miller)** Miller, Eleftheriou, Pattnaik, Ndirango, and News, J Chem Phys, 116, 8649 (2002).

**(Zhang)** Zhang, Glotzer, Nanoletters, 4, 1407-1413 (2004).

## 17.187 fix rigid/meso command

### 17.187.1 Syntax

```
fix ID group-ID rigid/meso bodystyle args keyword values ...
```

- ID, group-ID are documented in *fix* command
- rigid/meso = style name of this fix command
- bodystyle = *single* or *molecule* or *group*

*single* args = none

*molecule* args = none

*custom* args = *i\_propname* or *v\_varname*

*i\_propname* = an integer property defined via *fix property/atom*

*v\_varname* = an atom-style or atomfile-style variable

*group* args = N groupID1 groupID2 ...

N = # of groups

groupID1, groupID2, ... = list of N group IDs

- zero or more keyword/value pairs may be appended
- keyword = *reinit* or *force* or *torque* or *infile*

*reinit* = *yes* or *no*

*force* values = M xflag yflag zflag

M = which rigid body from 1-Nbody (see asterisk form below)

xflag,yflag,zflag = off/on if component of center-of-mass force is active

*torque* values = M xflag yflag zflag

M = which rigid body from 1-Nbody (see asterisk form below)

xflag,yflag,zflag = off/on if component of center-of-mass torque is active

*infile* filename

filename = file with per-body values of mass, center-of-mass, moments of inertia

### 17.187.2 Examples

```
fix 1 ellipsoid rigid/meso single
fix 1 rods      rigid/meso molecule
fix 1 spheres  rigid/meso single force 1 off off on
fix 1 particles rigid/meso molecule force 1*5 off off off force 6*10 off off on
fix 2 spheres  rigid/meso group 3 sphere1 sphere2 sphere3 torque * off off off
```

### 17.187.3 Description

Treat one or more sets of mesoscopic SPH/SDPD particles as independent rigid bodies. This means that each timestep the total force and torque on each rigid body is computed as the sum of the forces and torques on its constituent particles. The coordinates and velocities of the particles in each body are then updated so that the body moves and rotates as a single entity using the methods described in the paper by (*Miller*). Density and internal energy of the particles will also be updated. This is implemented by creating internal data structures for each rigid body and performing time integration on these data structures. Positions and velocities of the constituent particles are regenerated from the rigid body data structures in every time step. This restricts which operations and fixes can be applied to rigid bodies. See below for a detailed discussion.



The operation of this fix is exactly like that described by the *fix rigid/mve* command, except that particles' density, internal energy and extrapolated velocity are also updated.

---

**Note:** You should not update the particles in rigid bodies via other time-integration fixes (e.g. *fix sph*, *fix sph/stationary*), or you will have conflicting updates to positions and velocities resulting in unphysical behavior in most cases. When performing a hybrid simulation with some atoms in rigid bodies, and some not, a separate time integration fix like *fix sph* should be used for the non-rigid particles.

---

**Note:** These fixes are overkill if you simply want to hold a collection of particles stationary or have them move with a constant velocity. To hold particles stationary use *fix sph/stationary* instead. If you would like to move particles with a constant velocity use *fix meso/move*.

---

**Warning:** The aggregate properties of each rigid body are calculated at the start of a simulation run and are maintained in internal data structures. The properties include the position and velocity of the center-of-mass of the body, its moments of inertia, and its angular momentum. This is done using the properties of the constituent particles of the body at that point in time (or see the *infile* keyword option). Thereafter, changing these properties of individual particles in the body will have no effect on a rigid body's dynamics, unless they effect any computation of per-particle forces or torques. If the keyword *reinit* is set to *yes* (the default), the rigid body data structures will be recreated at the beginning of each *run* command; if the keyword *reinit* is set to *no*, the rigid body data structures will be built only at the very first *run* command and maintained for as long as the rigid fix is defined. For example, you might think you could displace the particles in a body or add a large velocity to each particle in a body to make it move in a desired direction before a second run is performed, using the *set* or *displace\_atoms* or *velocity* commands. But these commands will not affect the internal attributes of the body unless *reinit* is set to *yes*. With *reinit* set to *no* (or using the *infile* option, which implies *reinit no*) the position and velocity of individual particles in the body will be reset when time integration starts again.

---

Each rigid body must have two or more particles. A particle can belong to at most one rigid body. Which particles are in which bodies can be defined via several options.

For bodystyle *single* the entire fix group of particles is treated as one rigid body.

For bodystyle *molecule*, particles are grouped into rigid bodies by their respective molecule IDs: each set of particles in the fix group with the same molecule ID is treated as a different rigid body. Note that particles with a molecule ID = 0 will be treated as a single rigid body. For a system with solvent (typically this is particles with molecule ID = 0) surrounding rigid bodies, this may not be what you want. Thus you should be careful to use a fix group that only includes particles you want to be part of rigid bodies.

Bodystyle *custom* is similar to bodystyle *molecule* except that it is more flexible in using other per-atom properties to define the sets of particles that form rigid bodies. An integer vector defined by the *fix property/atom* command can be used. Or an *atom-style* or *atomfile-style variable* can be used; the floating-point value produced by the variable is rounded to an integer. As with bodystyle *molecule*, each set of particles in the fix groups with the same integer value is treated as a different rigid body. Since fix property/atom vectors and atom-style variables produce values for all particles, you should be careful to use a fix group that only includes particles you want to be part of rigid bodies.

For bodystyle *group*, each of the listed groups is treated as a separate rigid body. Only particles that are also in the fix group are included in each rigid body.

---

**Note:** To compute the initial center-of-mass position and other properties of each rigid body, the image flags for each particle in the body are used to "unwrap" the particle coordinates. Thus you must insure that these image flags are consistent so that the unwrapping creates a valid rigid body (one where the particles are close together), particularly

if the particles in a single rigid body straddle a periodic boundary. This means the input data file or restart file must define the image flags for each particle consistently or that you have used the *set* command to specify them correctly. If a dimension is non-periodic then the image flag of each particle must be 0 in that dimension, else an error is generated.

---

By default, each rigid body is acted on by other particles which induce an external force and torque on its center of mass, causing it to translate and rotate. Components of the external center-of-mass force and torque can be turned off by the *force* and *torque* keywords. This may be useful if you wish a body to rotate but not translate, or vice versa, or if you wish it to rotate or translate continuously unaffected by interactions with other particles. Note that if you expect a rigid body not to move or rotate by using these keywords, you must insure its initial center-of-mass translational or angular velocity is 0.0. Otherwise the initial translational or angular momentum, the body has, will persist.

An *xflag*, *yflag*, or *zflag* set to *off* means turn off the component of force or torque in that dimension. A setting of *on* means turn on the component, which is the default. Which rigid body(s) the settings apply to is determined by the first argument of the *force* and *torque* keywords. It can be an integer *M* from 1 to *Nbody*, where *Nbody* is the number of rigid bodies defined. A wild-card asterisk can be used in place of, or in conjunction with, the *M* argument to set the flags for multiple rigid bodies. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If *N* = the number of rigid bodies, then an asterisk with no numeric values means all bodies from 1 to *N*. A leading asterisk means all bodies from 1 to *n* (inclusive). A trailing asterisk means all bodies from *n* to *N* (inclusive). A middle asterisk means all bodies from *m* to *n* (inclusive). Note that you can use the *force* or *torque* keywords as many times as you like. If a particular rigid body has its component flags set multiple times, the settings from the final keyword are used.

For computational efficiency, you should typically define one *fix rigid/meso* command which includes all the desired rigid bodies. LAMMPS will allow multiple *fix rigid/meso* fixes to be defined, but it is more expensive.

---

The keyword/value option pairs are used in the following ways.

The *reinit* keyword determines, whether the rigid body properties are re-initialized between run commands. With the option *yes* (the default) this is done, with the option *no* this is not done. Turning off the re-initialization can be helpful to protect rigid bodies against unphysical manipulations between runs or when properties cannot be easily re-computed (e.g. when read from a file). When using the *infile* keyword, the *reinit* option is automatically set to *no*.

---

The *infile* keyword allows a file of rigid body attributes to be read in from a file, rather than having LAMMPS compute them. There are 5 such attributes: the total mass of the rigid body, its center-of-mass position, its 6 moments of inertia, its center-of-mass velocity, and the 3 image flags of the center-of-mass position. For rigid bodies consisting of point particles or non-overlapping finite-size particles, LAMMPS can compute these values accurately. However, for rigid bodies consisting of finite-size particles which overlap each other, LAMMPS will ignore the overlaps when computing these 4 attributes. The amount of error this induces depends on the amount of overlap. To avoid this issue, the values can be pre-computed (e.g. using Monte Carlo integration).

The format of the file is as follows. Note that the file does not have to list attributes for every rigid body integrated by *fix rigid*. Only bodies which the file specifies will have their computed attributes overridden. The file can contain initial blank lines or comment lines starting with “#” which are ignored. The first non-blank, non-comment line should list *N* = the number of lines to follow. The *N* successive lines contain the following information:

```
ID1 masstotal xcm ycm zcm ixc iyc izz ixc izc iyc vxcm vycm vzcm lx ly lz ixcm iycm izcm
ID2 masstotal xcm ycm zcm ixc iyc izz ixc izc iyc vxcm vycm vzcm lx ly lz ixcm iycm izcm
...
IDN masstotal xcm ycm zcm ixc iyc izz ixc izc iyc vxcm vycm vzcm lx ly lz ixcm iycm izcm
```

The rigid body IDs are all positive integers. For the *single* bodystyle, only an ID of 1 can be used. For the *group* bodystyle, IDs from 1 to *Ng* can be used where *Ng* is the number of specified groups. For the *molecule* bodystyle, use the molecule ID for the atoms in a specific rigid body as the rigid body ID.

---

The `masstotal` and center-of-mass coordinates (`xcm,ycm,zcm`) are self-explanatory. The center-of-mass should be consistent with what is calculated for the position of the rigid body with all its atoms unwrapped by their respective image flags. If this produces a center-of-mass that is outside the simulation box, LAMMPS wraps it back into the box.

The 6 moments of inertia (`ixx,iyy,izz,ixy,ixz,iyz`) should be the values consistent with the current orientation of the rigid body around its center of mass. The values are with respect to the simulation box XYZ axes, not with respect to the principal axes of the rigid body itself. LAMMPS performs the latter calculation internally.

The (`vxcm,vycm,vzcm`) values are the velocity of the center of mass. The (`lx,ly,lz`) values are the angular momentum of the body. The (`vxcm,vycm,vzcm`) and (`lx,ly,lz`) values can simply be set to 0 if you wish the body to have no initial motion.

The (`ixcm,iycm,izcm`) values are the image flags of the center of mass of the body. For periodic dimensions, they specify which image of the simulation box the body is considered to be in. An image of 0 means it is inside the box as defined. A value of 2 means add 2 box lengths to get the true value. A value of -1 means subtract 1 box length to get the true value. LAMMPS updates these flags as the rigid bodies cross periodic boundaries during the simulation.

---

**Note:** If you use the `infile` keyword and write restart files during a simulation, then each time a restart file is written, the fix also write an auxiliary restart file with the name `rfile.rigid`, where “`rfile`” is the name of the restart file, e.g. `tmp.restart.10000` and `tmp.restart.10000.rigid`. This auxiliary file is in the same format described above. Thus it can be used in a new input script that restarts the run and re-specifies a rigid fix using an `infile` keyword and the appropriate filename. Note that the auxiliary file will contain one line for every rigid body, even if the original file only listed a subset of the rigid bodies.

---

#### 17.187.4 Restart, fix\_modify, output, run start/stop, minimize info

No information is written to *binary restart files*. If the `infile` keyword is used, an auxiliary file is written out with rigid body information each time a restart file is written, as explained above for the `infile` keyword.

None of the `fix_modify` options are relevant to this fix.

This fix computes a global array of values which can be accessed by various *output commands*.

The number of rows in the array is equal to the number of rigid bodies. The number of columns is 28. Thus for each rigid body, 28 values are stored: the xyz coords of the center of mass (COM), the xyz components of the COM velocity, the xyz components of the force acting on the COM, the components of the 4-vector quaternion representing the orientation of the rigid body, the xyz components of the angular velocity of the body around its COM, the xyz components of the torque acting on the COM, the 3 principal components of the moment of inertia, the xyz components of the angular momentum of the body around its COM, and the xyz image flags of the COM.

The center of mass (COM) for each body is similar to unwrapped coordinates written to a dump file. It will always be inside (or slightly outside) the simulation box. The image flags have the same meaning as image flags for particle positions (see the “`dump`” command). This means you can calculate the unwrapped COM by applying the image flags to the COM, the same as when unwrapped coordinates are written to a dump file.

The force and torque values in the array are not affected by the `force` and `torque` keywords in the fix rigid command; they reflect values before any changes are made by those keywords.

The ordering of the rigid bodies (by row in the array) is as follows. For the `single` keyword there is just one rigid body. For the `molecule` keyword, the bodies are ordered by ascending molecule ID. For the `group` keyword, the list of group IDs determines the ordering of bodies.

The array values calculated by this fix are “intensive”, meaning they are independent of the number of particles in the simulation.

No parameter of this fix can be used with the `start/stop` keywords of the `run` command.

This fix is not invoked during *energy minimization*.

---

### 17.187.5 Restrictions

This fix is part of the DPD-SMOOTH package and also depends on the RIGID package. It is only enabled if LAMMPS was built with both packages. See the *Build package* page for more info.

This fix requires that atoms store density and internal energy as defined by the *atom\_style sph* command.

All particles in the group must be mesoscopic SPH/SDPD particles.

### 17.187.6 Related commands

*fix meso/move*, *fix rigid*, *neigh\_modify exclude*

### 17.187.7 Default

The option defaults are force \* on on on and torque \* on on on, meaning all rigid bodies are acted on by center-of-mass force and torque. Also reinit = yes.

---

(Miller) Miller, Eleftheriou, Pattnaik, Ndirango, and Newns, J Chem Phys, 116, 8649 (2002).

## 17.188 fix rx command

Accelerator Variants: *rx/kk*

### 17.188.1 Syntax

```
fix ID group-ID rx file localTemp matrix solver minSteps ...
```

- ID, group-ID are documented in *fix* command
- rx = style name of this fix command
- file = filename containing the reaction kinetic equations and Arrhenius parameters
- localTemp = *none*, *lucy* = no local temperature averaging or local temperature defined through Lucy weighting function
- matrix = *sparse*, *dense* format for the stoichiometric matrix
- solver = *lammps\_rk4*, *rkf45* = rk4 is an explicit fourth order Runge-Kutta method; rkf45 is an adaptive fourth-order Runge-Kutta-Fehlberg method
- minSteps = # of steps for rk4 solver or minimum # of steps for rkf45 (rk4 or rkf45)
- maxSteps = maximum number of steps for the rkf45 solver (rkf45 only)
- relTol = relative tolerance for the rkf45 solver (rkf45 only)
- absTol = absolute tolerance for the rkf45 solver (rkf45 only)
- diag = Diagnostics frequency for the rkf45 solver (optional, rkf45 only)

## 17.188.2 Examples

```
fix 1 all rx kinetics.rx none dense lammmps_rk4
fix 1 all rx kinetics.rx none sparse lammmps_rk4 1
fix 1 all rx kinetics.rx lucy sparse lammmps_rk4 10
fix 1 all rx kinetics.rx none dense rkf45 1 100 1e-6 1e-8
fix 1 all rx kinetics.rx none dense rkf45 1 100 1e-6 1e-8 -1
```

## 17.188.3 Description

Fix *rx* solves the reaction kinetic ODEs for a given reaction set that is defined within the file associated with this command.

For a general reaction such that



the reaction rate equation is defined to be of the form

$$r = k(T)[A]^{\nu_A}[B]^{\nu_B}$$

In the current implementation, the exponents are defined to be equal to the stoichiometric coefficients. A given reaction set consisting of  $n$  reaction equations will contain a total of  $m$  species. A set of  $m$  ordinary differential equations (ODEs) that describe the change in concentration of a given species as a function of time are then constructed based on the  $n$  reaction rate equations.

The ODE systems are solved over the full DPD timestep  $dt$  using either a fourth order Runge-Kutta *rk4* method with a fixed step-size  $h$ , specified by the *lammmps\_rk4* keyword, or a fourth order Runge-Kutta-Fehlberg (rkf45) method with an adaptive step-size for  $h$ . The number of ODE steps per DPD timestep for the rk4 method is optionally specified immediately after the rk4 keyword. The ODE step-size is set as  $dt/num\_steps$ . Smaller step-sizes tend to yield more accurate results but there is not control on the error. For error control, use the rkf45 ODE solver.

The rkf45 method adjusts the step-size so that the local truncation error is held within the specified absolute and relative tolerances. The initial step-size  $h0$  can be specified by the user or estimated internally. It is recommended that the user specify  $h0$  since this will generally reduced the number of ODE integration steps required.  $h0$  is defined as  $dt/min\_steps$  if  $min\_steps \geq 1$ . If  $min\_steps == 0$ ,  $h0$  is estimated such that an explicit Euler method would likely produce an acceptable solution. This is generally overly conservative for the fourth-order method and users are advised to specify  $h0$  as some fraction of the DPD timestep. For small DPD timesteps, only one step may be necessary depending upon the tolerances. Note that more than  $min\_steps$  ODE steps may be taken depending upon the ODE stiffness but no more than  $max\_steps$  will be taken. If  $max\_steps$  is reached, an error warning is printed and the simulation is stopped.

After each ODE step, the solution error  $e$  is tested and weighted using the *absTol* and *relTol* values. The error vector is weighted as  $e / (relTol * |u| + absTol)$  where  $u$  is the solution vector. If the norm of the error is  $\leq 1$ , the solution is accepted,  $h$  is increased by a proportional amount, and the next ODE step is begun. Otherwise,  $h$  is shrunk and the ODE step is repeated.

Run-time diagnostics are available for the rkf45 ODE solver. The frequency (in timesteps) that diagnostics are reported is controlled by the last (optional) 12th argument. A negative frequency means that diagnostics are reported once at the end of each run. A positive value  $N$  means that the diagnostics are reported once per  $N$  timesteps.

The diagnostics report the average # of integrator steps and RHS function evaluations and run-time per ODE as well as the average/RMS/min/max per process. If the reporting frequency is 1, the RMS/min/max per ODE are also reported. The per ODE statistics can be used to adjust the tolerance and min/max step parameters. The statistics per MPI process can be useful to examine any load imbalance caused by the adaptive ODE solver. (Some DPD particles can take longer to solve than others. This can lead to an imbalance across the MPI processes.)

The filename specifies a file that contains the entire set of reaction kinetic equations and corresponding Arrhenius parameters. The format of this file is described below.

There is no restriction on the total number or reaction equations that are specified. The species names are arbitrary string names that are associated with the species concentrations. Each species in a given reaction must be preceded by its stoichiometric coefficient. The only delimiters that are recognized between the species are either a + or = character. The = character corresponds to an irreversible reaction. After specifying the reaction, the reaction rate constant is determined through the temperature dependent Arrhenius equation:

$$k = AT^n e^{\frac{-E_a}{k_B T}}$$

where  $A$  is the Arrhenius factor in time units or concentration/time units,  $n$  is the unitless exponent of the temperature dependence, and  $E_a$  is the activation energy in energy units. The temperature dependence can be removed by specifying the exponent as zero.

The internal temperature of the coarse-grained particles can be used in constructing the reaction rate constants at every DPD timestep by specifying the keyword *none*. Alternatively, the keyword *lucy* can be specified to compute a local-average particle internal temperature for use in the reaction rate constant expressions. The local-average particle internal temperature is defined as:

$$\theta_i^{-1} = \frac{\sum_{j=1} \omega_{Lucy}(r_{ij}) \theta_j^{-1}}{\sum_{j=1} \omega_{Lucy}(r_{ij})}$$

where the Lucy function is expressed as:

$$\omega_{Lucy}(r_{ij}) = \left(1 + \frac{3r_{ij}}{r_c}\right) \left(1 - \frac{r_{ij}}{r_c}\right)^3$$

The self-particle interaction is included in the above equation.

The stoichiometric coefficients for the reaction mechanism are stored in either a sparse or dense matrix format. The dense matrix should only be used for small reaction mechanisms. The sparse matrix should be used when there are many reactions (e.g., more than 5). This allows the number of reactions and species to grow while keeping the computational cost tractable. The matrix format can be specified as using either the *sparse* or *dense* keywords. If all stoichiometric coefficients for a reaction are small integers (whole numbers  $\leq 3$ ), a fast exponential function is used. This can save significant computational time so users are encouraged to use integer coefficients where possible.

The format of a tabulated file is as follows (without the parenthesized comments):

```
# Rxn equations and parameters (one or more comment or blank lines)
1.0 hcn + 1.0 no2 = 1.0 no + 0.5 n2 + 0.5 h2 + 1.0 co 2.49E+01 0.0 1.34 (rxn equation, A, n, Ea)
1.0 hcn + 1.0 no = 1.0 co + 1.0 n2 + 0.5 h2 2.16E+00 0.0 1.52
...
1.0 no + 1.0 co = 0.5 n2 + 1.0 co2 1.66E+06 0.0 0.69
```

A section begins with a non-blank line whose first character is not a "#"; blank lines or lines starting with "#" can be used as comments between sections.

Following a blank line, the next  $N$  lines list the  $N$  reaction equations. Each species within the reaction equation is specified through its stoichiometric coefficient and a species tag. Reactant species are specified on the left-hand side of the equation and product species are specified on the right-hand side of the equation. After specifying the reactant and product species, the final three arguments of each line represent the Arrhenius parameter  $A$ , the temperature exponent  $n$ , and the activation energy  $E_a$ .

Note that the species tags that are defined in the reaction equations are used by the *fix eos/table/rx* command to define the thermodynamic properties of each species. Furthermore, the number of species molecules (i.e., concentration) can be specified either with the *set* command using the “d\_” prefix or by reading directly the concentrations from a data file. For the latter case, the *read\_data* command with the *fix* keyword should be specified, where the *fix*-ID will be the “fix rx`ID with a <SPECIES>`\_ suffix, e.g.

```
fix foo all rx reaction.file ... read_data data.dpd fix foo_SPECIES NULL Species
```

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

## 17.188.4 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This command also requires use of the *atom\_style dpd* command.

This command can only be used with a constant energy or constant enthalpy DPD simulation.

## 17.188.5 Related commands

*fix eos/table/rx*, *fix shardlow*, *pair dpd/fdt/energy*

## 17.188.6 Default

none

## 17.189 fix saed/vtk command

### 17.189.1 Syntax

```
fix ID group-ID saed/vtk Nevery Nrepeat Nfreak c_ID attribute args ... keyword args ...
```

- ID, group-ID are documented in *fix* command
- saed/vtk = style name of this *fix* command
- Nevery = use input values every this many timesteps
- Nrepeat = # of times to use input values for calculating averages

- Nfreq = calculate averages every this many timesteps
- c\_ID = saed compute ID

```
keyword = file or ave or start or file or overwrite:l
ave args = one or running or window M
  one = output a new average value every Nfreq steps
  running = output cumulative average of all previous Nfreq steps
  window M = output average of M most recent Nfreq steps
start args = Nstart
  Nstart = start averaging on this timestep
file arg = filename
  filename = name of file to output time averages to
```

## 17.189.2 Examples

```
compute 1 all saed 0.0251 Al 0 Kmax 1.70 Zone 0 0 1 dR_Ewald 0.01 c 0.5 0.5 0.5
compute 2 all saed 0.0251 Ni Kmax 1.70 Zone 0 0 0 c 0.05 0.05 0.05 manual echo

fix 1 all saed/vtk 1 1 1 c_1 file Al203_001.saed
fix 2 all saed/vtk 1 1 1 c_2 file Ni_000.saed
```

## 17.189.3 Description

Time average computed intensities from *compute saed* and write output to a file in the third generation vtk image data format for visualization directly in parallelized visualization software packages like ParaView and VisIt. Note that if no time averaging is done, this command can be used as a convenient way to simply output diffraction intensities at a single snapshot.

To produce output in the image data vtk format ghost data is added outside the *Kmax* range assigned in the *compute saed*. The ghost data is assigned a value of -1 and can be removed setting a minimum isovolume of 0 within the visualization software. SAED images can be created by visualizing a spherical slice of the data that is centered at  $R\_Ewald * [h k l] / \text{norm}([h k l])$ , where  $R\_Ewald = 1/\lambda$ .

The group specified within this command is ignored. However, note that specified values may represent calculations performed by saed computes which store their own “group” definitions.

Fix saed/vtk is designed to work only with *compute saed* values, e.g.

```
compute 3 top saed 0.0251 Al 0
fix saed/vtk 1 1 1 c_3 file Al203_001.saed
```

---

The *Nevery*, *Nrepeat*, and *Nfreq* arguments specify on what timesteps the input values will be used in order to contribute to the average. The final averaged quantities are generated on timesteps that are a multiple of *Nfreq*. The average is over *Nrepeat* quantities, computed in the preceding portion of the simulation every *Nevery* timesteps. *Nfreq* must be a multiple of *Nevery* and *Nevery* must be non-zero even if *Nrepeat* is 1. Also, the timesteps contributing to the average value cannot overlap, i.e.  $Nrepeat * Nevery$  can not exceed *Nfreq*.

For example, if *Nevery*=2, *Nrepeat*=6, and *Nfreq*=100, then values on timesteps 90,92,94,96,98,100 will be used to compute the final average on timestep 100. Similarly for timesteps 190,192,194,196,198,200 on timestep 200, etc. If *Nrepeat*=1 and *Nfreq* = 100, then no time averaging is done; values are simply generated on timesteps 100,200,etc.

---



The output for `fix ave/time/saed` is a file written with the third generation vtk image data formatting. The filename assigned by the `file` keyword is appended with `_N.vtk` where `N` is an index (0,1,2...) to account for multiple diffraction intensity outputs.

By default the header contains the following information (with example data):

```
# vtk DataFile Version 3.0 c_SAED
Image data set
ASCII
DATASET STRUCTURED_POINTS
DIMENSIONS 337 219 209
ASPECT_RATIO 0.00507953 0.00785161 0.00821458
ORIGIN -0.853361 -0.855826 -0.854316
POINT_DATA 15424827
SCALARS intensity float
LOOKUP_TABLE default
...data
```

In this example, `kspace` is sampled across a  $337 \times 219 \times 209$  point mesh where the mesh spacing is approximately 0.005, 0.007, and 0.008  $\text{inv}(\text{length})$  units in the `k1`, `k2`, and `k3` directions, respectively. The data is shifted by -0.85, -0.85, -0.85  $\text{inv}(\text{length})$  units so that the origin will lie at 0, 0, 0. Here, 15,424,827 `kspace` points are sampled in total.

Additional optional keywords also affect the operation of this `fix`.

The `ave` keyword determines how the values produced every `Nfreq` steps are averaged with values produced on previous steps that were multiples of `Nfreq`, before they are accessed by another output command or written to a file.

If the `ave` setting is `one`, then the values produced on timesteps that are multiples of `Nfreq` are independent of each other; they are output as-is without further averaging.

If the `ave` setting is `running`, then the values produced on timesteps that are multiples of `Nfreq` are summed and averaged in a cumulative sense before being output. Each output value is thus the average of the value produced on that timestep with all preceding values. This running average begins when the `fix` is defined; it can only be restarted by deleting the `fix` via the `unfix` command, or by re-defining the `fix` by re-specifying it.

If the `ave` setting is `window`, then the values produced on timesteps that are multiples of `Nfreq` are summed and averaged within a moving “window” of time, so that the last `M` values are used to produce the output. E.g. if `M = 3` and `Nfreq = 1000`, then the output on step 10000 will be the average of the individual values on steps 8000,9000,10000. Outputs on early steps will average over less than `M` values if they are not available.

The `start` keyword specifies what timestep averaging will begin on. The default is step 0. Often input values can be 0.0 at time 0, so setting `start` to a larger value can avoid including a 0.0 in a running or windowed average.

The `file` keyword allows a filename to be specified. Every `Nfreq` steps, the vector of `saed` intensity data is written to a new file using the third generation vtk format. The base of each file is assigned by the `file` keyword and this string is appended with `_N.vtk` where `N` is an index (0,1,2...) to account for situations with multiple diffraction intensity outputs.

### 17.189.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.189.5 Restrictions

The attributes for *fix\_saed\_vtk* must match the values assigned in the associated *compute\_saed* command.

### 17.189.6 Related commands

*compute\_saed*

### 17.189.7 Default

The option defaults are *ave = one*, *start = 0*, no file output.

---

(Coleman) Coleman, Spearot, Capolungo, MSMSE, 21, 055020 (2013).

## 17.190 fix setforce command

Accelerator Variants: *setforce/kk*

## 17.191 fix setforce/spin command

### 17.191.1 Syntax

```
fix ID group-ID setforce fx fy fz keyword value ...
```

- ID, group-ID are documented in *fix* command
- setforce = style name of this fix command
- fx,fy,fz = force component values
- any of fx,fy,fz can be a variable (see below)
- zero or more keyword/value pairs may be appended to args
- keyword = *region*

*region* value = region-ID

region-ID = ID of region atoms must be in to have added force

## 17.191.2 Examples

```
fix freeze indenter setforce 0.0 0.0 0.0
fix 2 edge setforce NULL 0.0 0.0
fix 1 edge setforce/spin 0.0 0.0 0.0
fix 2 edge setforce NULL 0.0 v_oscillate
```

## 17.191.3 Description

Set each component of force on each atom in the group to the specified values  $f_x, f_y, f_z$ . This erases all previously computed forces on the atom, though additional fixes could add new forces. This command can be used to freeze certain atoms in the simulation by zeroing their force, either for running dynamics or performing an energy minimization. For dynamics, this assumes their initial velocity is also zero.

Any of the  $f_x, f_y, f_z$  values can be specified as NULL which means do not alter the force component in that dimension.

Any of the 3 quantities defining the force components can be specified as an equal-style or atom-style *variable*, namely  $f_x, f_y, f_z$ . If the value is a variable, it should be specified as  $v\_name$ , where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the force component.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent force field.

Atom-style variables can specify the same formulas as equal-style variables but can also include per-atom values, such as atom coordinates. Thus it is easy to specify a spatially-dependent force field with optional time-dependence as well.

If the *region* keyword is used, the atom must also be in the specified geometric *region* in order to have force added to it.

---

Style *spin* suffix sets the components of the magnetic precession vectors instead of the mechanical forces. This also erases all previously computed magnetic precession vectors on the atom, though additional magnetic fixes could add new forces.

This command can be used to freeze the magnetic moment of certain atoms in the simulation by zeroing their precession vector.

All options defined above remain valid, they just apply to the magnetic precession vectors instead of the forces.

---

Styles with a *gpu, intel, kk, omp, or opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

**Note:** The region keyword is supported by Kokkos, but a Kokkos-enabled region must be used. See the region *region* command for more information.

---

### 17.191.4 Restart, `fix_modify`, `output`, `run start/stop`, minimize info

No information about this fix is written to *binary restart files*.

The `fix_modify respa` option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is setting the forces to the desired values; on all other levels, the force is set to 0.0 for the atoms in the fix group, so that setforce values are not counted multiple times. Default is to to override forces at the outermost level.

This fix computes a global 3-vector of forces, which can be accessed by various *output commands*. This is the total force on the group of atoms before the forces on individual atoms are changed by the fix. The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the `run` command.

The forces due to this fix are imposed during an energy minimization, invoked by the `minimize` command, but you cannot set forces to any value besides zero when performing a minimization. Use the `fix addforce` command if you want to apply a non-zero force to atoms during a minimization.

### 17.191.5 Restrictions

The fix `setforce/spin` only makes sense when LAMMPS was built with the SPIN package.

### 17.191.6 Related commands

`fix addforce`, `fix aveforce`

### 17.191.7 Default

none

## 17.192 `fix shake` command

Accelerator Variants: `shake/kk`

## 17.193 `fix rattle` command

### 17.193.1 Syntax

```
fix ID group-ID style tol iter N constraint values ... keyword value ...
```

- ID, group-ID are documented in `fix` command
- style = shake or rattle = style name of this fix command
- tol = accuracy tolerance of SHAKE solution
- iter = max # of iterations in each SHAKE solution

- N = print SHAKE statistics every this many timesteps (0 = never)
- one or more constraint/value pairs are appended
- constraint = *b* or *a* or *t* or *m*
  - b* values = one or more bond types
  - a* values = one or more angle types
  - t* values = one or more atom types
  - m* value = one or more mass values
- zero or more keyword/value pairs may be appended
- keyword = *mol* or *kbond*
  - mol* value = template-ID
    - template-ID = ID of molecule template specified in a separate *molecule* command
  - kbond* value = force constant
    - force constant = force constant used to apply a restraint force when used during ↵  
↵minimization

## 17.193.2 Examples

```
fix 1 sub shake 0.0001 20 10 b 4 19 a 3 5 2
fix 1 sub shake 0.0001 20 10 t 5 6 m 1.0 a 31
fix 1 sub shake 0.0001 20 10 t 5 6 m 1.0 a 31 mol myMol
fix 1 sub rattle 0.0001 20 10 t 5 6 m 1.0 a 31
fix 1 sub rattle 0.0001 20 10 t 5 6 m 1.0 a 31 mol myMol
```

## 17.193.3 Description

Apply bond and angle constraints to specified bonds and angles in the simulation by either the SHAKE or RATTLE algorithms. This typically enables a longer timestep. The SHAKE or RATTLE algorithms, however, can *only* be applied during molecular dynamics runs. When this fix is used during a minimization, the constraints are *approximated* by strong harmonic restraints.

### SHAKE vs RATTLE:

The SHAKE algorithm was invented for schemes such as standard Verlet timestepping, where only the coordinates are integrated and the velocities are approximated as finite differences to the trajectories (*Ryckaert et al. (1977)*). If the velocities are integrated explicitly, as with velocity Verlet which is what LAMMPS uses as an integration method, a second set of constraining forces is required in order to eliminate velocity components along the bonds (*Andersen (1983)*).

In order to formulate individual constraints for SHAKE and RATTLE, focus on a single molecule whose bonds are constrained. Let  $\mathbf{r}_i$  and  $\mathbf{v}_i$  be the position and velocity of atom  $i$  at time  $n$ , for  $i=1,\dots,N$ , where  $N$  is the number of sites of our reference molecule. The distance vector between sites  $i$  and  $j$  is given by

$$\mathbf{r}_{ij}^{n+1} = \mathbf{r}_j^n - \mathbf{r}_i^n$$

The constraints can then be formulated as

$$\begin{aligned} \mathbf{r}_{ij}^{n+1} \cdot \mathbf{r}_{ij}^{n+1} &= d_{ij}^2 \quad \text{and} \\ \mathbf{v}_{ij}^{n+1} \cdot \mathbf{r}_{ij}^{n+1} &= 0 \end{aligned}$$

The SHAKE algorithm satisfies the first condition, i.e. the sites at time  $n+1$  will have the desired separations  $D_{ij}$  immediately after the coordinates are integrated. If we also enforce the second condition, the velocity components

along the bonds will vanish. RATTLE satisfies both conditions. As implemented in LAMMPS, *fix rattle* uses *fix shake* for satisfying the coordinate constraints. Therefore the settings and optional keywords are the same for both fixes, and all the information below about SHAKE is also relevant for RATTLE.

### SHAKE:

Each timestep the specified bonds and angles are reset to their equilibrium lengths and angular values via the SHAKE algorithm (*Ryckaert et al. (1977)*). This is done by applying an additional constraint force so that the new positions preserve the desired atom separations. The equations for the additional force are solved via an iterative method that typically converges to an accurate solution in a few iterations. The desired tolerance (e.g.  $1.0e-4 = 1$  part in 10000) and maximum # of iterations are specified as arguments. Setting the *N* argument will print statistics to the screen and log file about regarding the lengths of bonds and angles that are being constrained. Small delta values mean SHAKE is doing a good job.

In LAMMPS, only small clusters of atoms can be constrained. This is so the constraint calculation for a cluster can be performed by a single processor, to enable good parallel performance. A cluster is defined as a central atom connected to others in the cluster by constrained bonds. LAMMPS allows for the following kinds of clusters to be constrained: one central atom bonded to 1 or 2 or 3 atoms, or one central atom bonded to 2 others and the angle between the 3 atoms also constrained. This means water molecules or CH<sub>2</sub> or CH<sub>3</sub> groups may be constrained, but not all the C-C backbone bonds of a long polymer chain.

The *b* constraint lists bond types that will be constrained. The *t* constraint lists atom types. All bonds connected to an atom of the specified type will be constrained. The *m* constraint lists atom masses. All bonds connected to atoms of the specified masses will be constrained (within a fudge factor of MASSDELTA specified in *fix\_shake.cpp*). The *a* constraint lists angle types. If both bonds in the angle are constrained then the angle will also be constrained if its type is in the list.

For all constraints, a particular bond is only constrained if both atoms in the bond are in the group specified with the SHAKE fix.

The degrees-of-freedom removed by SHAKE bonds and angles are accounted for in temperature and pressure computations. Similarly, the SHAKE contribution to the pressure of the system (virial) is also accounted for.

---

**Note:** This command works by using the current forces on atoms to calculate an additional constraint force which when added will leave the atoms in positions that satisfy the SHAKE constraints (e.g. bond length) after the next time integration step. If you define fixes (e.g. *fix efield*) that add additional force to the atoms after *fix shake* operates, then this fix will not take them into account and the time integration will typically not satisfy the SHAKE constraints. The solution for this is to make sure that *fix shake* is defined in your input script after any other fixes which add or change forces (to atoms that *fix shake* operates on).

---

The *mol* keyword should be used when other commands, such as *fix deposit* or *fix pour*, add molecules on-the-fly during a simulation, and you wish to constrain the new molecules via SHAKE. You specify a *template-ID* previously defined using the *molecule* command, which reads a file that defines the molecule. You must use the same *template-ID* that the command adding molecules uses. The coordinates, atom types, special bond restrictions, and SHAKE info can be specified in the molecule file. See the *molecule* command for details. The only settings required to be in this file (by this command) are the SHAKE info of atoms in the molecule.

The *kbond* keyword sets the restraint force constant when *fix shake* or *fix rattle* are used during minimization. In that case the constraint algorithms are *not* applied and restraint forces are used instead to maintain the geometries similar to the constraints. How well the geometries are maintained and how quickly a minimization converges, depends largely on the force constant *kbond*: larger values will reduce the deviation from the desired geometry, but can also lead to slower convergence of the minimization or lead to instabilities depending on the minimization algorithm requiring to reduce the value of *timestep*. Even though the restraints will not preserve the bond lengths and angles as closely as the constraints during the MD, they are generally close enough so that the constraints will be fulfilled to the desired

accuracy within a few MD steps following the minimization. The default value for *kbond* depends on the *units* setting and is  $1.0e6*k_B$ .

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### RATTLE:

The velocity constraints lead to a linear system of equations which can be solved analytically. The implementation of the algorithm in LAMMPS closely follows (*Andersen (1983)*).

---

**Note:** The *fix rattle* command modifies forces and velocities and thus should be defined after all other integration fixes in your input script. If you define other fixes that modify velocities or forces after *fix rattle* operates, then *fix rattle* will not take them into account and the overall time integration will typically not satisfy the RATTLE constraints. You can check whether the constraints work correctly by setting the value of RATTLE\_DEBUG in *src/fix\_rattle.cpp* to 1 and recompiling LAMMPS.

---

## 17.193.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about these fixes is written to *binary restart files*.

Both *fix shake* and *fix rattle* behave differently during a minimization in comparison to a molecular dynamics run:

- When used during a minimization, the SHAKE or RATTLE constraint algorithms themselves are **not** applied. Instead the constraints are replaced by harmonic restraint forces. The energy and virial contributions due to the restraint forces are tallied into global and per-atom accumulators. The total restraint energy is also accessible as a global scalar property of the fix.
- During molecular dynamics runs, however, the fixes do apply the requested SHAKE or RATTLE constraint algorithms.

The *fix\_modify virial* option is supported by these fixes to add the contribution due to the added constraint forces on atoms to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*.

The default setting for this fix is *fix\_modify virial yes*. No global or per-atom quantities are stored by these fixes for access by various *output commands* during an MD run. No parameter of these fixes can be used with the *start/stop* keywords of the *run* command.

### 17.193.5 Restrictions

These fixes are part of the RIGID package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

For computational efficiency, there can only be one shake or rattle fix defined in a simulation.

If you use a tolerance that is too large or a max-iteration count that is too small, the constraints will not be enforced very strongly, which can lead to poor energy conservation. You can test for this in your system by running a constant NVE simulation with a particular set of SHAKE parameters and monitoring the energy versus time.

SHAKE or RATTLE should not be used to constrain an angle at 180 degrees (e.g. a linear CO<sub>2</sub> molecule). This causes a divergence when solving the constraint equations numerically. You can use *fix rigid or fix rigid/small* instead to make a linear molecule rigid.

When used during minimization choosing a too large value of the *kbond* can make minimization very inefficient and also cause stability problems with some minimization algorithms. Sometimes those can be avoided by reducing the *timestep*.

### 17.193.6 Related commands

*fix rigid* <fix\_rigid>, *fix ehex* <fix\_ehex>, *fix nve/manifold/rattle* <fix\_nve\_manifold\_rattle>

### 17.193.7 Default

*kbond* = 1.0e9\*k<sub>B</sub>

---

**(Ryckaert)** J.-P. Ryckaert, G. Ciccotti and H. J. C. Berendsen, J of Comp Phys, 23, 327-341 (1977).

**(Andersen)** H. Andersen, J of Comp Phys, 52, 24-34 (1983).

## 17.194 fix shardlow command

Accelerator Variants: *shardlow/kk*

### 17.194.1 Syntax

```
fix ID group-ID shardlow
```

- ID, group-ID are documented in *fix* command
- shardlow = style name of this fix command



## 17.194.2 Examples

```
fix 1 all shardlow
```

## 17.194.3 Description

Specifies that the Shardlow splitting algorithm (SSA) is to be used to integrate the DPD equations of motion. The SSA splits the integration into a stochastic and deterministic integration step. The fix *shardlow* performs the stochastic integration step and must be used in conjunction with a deterministic integrator (e.g. *fix nve* or *fix nph*). The stochastic integration of the dissipative and random forces is performed prior to the deterministic integration of the conservative force. Further details regarding the method are provided in (*Lisal*) and (*Larentzos1*).

The fix *shardlow* must be used with the *pair\_style dpd/fdt* or *pair\_style dpd/fdt/energy* command to properly initialize the fluctuation-dissipation theorem parameter(s) sigma (and kappa, if necessary).

Note that numerous variants of DPD can be specified by choosing an appropriate combination of the integrator and *pair\_style dpd/fdt* command. DPD under isothermal conditions can be specified by using *fix shardlow*, *fix nve* and *pair\_style dpd/fdt*. DPD under isoenergetic conditions can be specified by using *fix shardlow*, *fix nve* and *pair\_style dpd/fdt/energy*. DPD under isobaric conditions can be specified by using *fix shardlow*, *fix nph* and *pair\_style dpd/fdt*. DPD under isoenthalpic conditions can be specified by using *fix shardlow*, *fix nph* and *pair\_style dpd/fdt/energy*. Examples of each DPD variant are provided in the examples/PACKAGES/dpd-react directory.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

## 17.194.4 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix is currently limited to orthogonal simulation cell geometries.

This fix must be used with an additional fix that specifies time integration, e.g. *fix nve* or *fix nph*.

The Shardlow splitting algorithm requires the sizes of the sub-domain lengths to be larger than twice the cutoff+skin. Generally, the domain decomposition is dependent on the number of processors requested.

## 17.194.5 Related commands

*pair\_style dpd/fdt, fix eos/cv*

## 17.194.6 Default

none

---

**(Lisal)** M. Lisal, J.K. Brennan, J. Bonet Avalos, “Dissipative particle dynamics as isothermal, isobaric, isoenergetic, and isoenthalpic conditions using Shardlow-like splitting algorithms.”, J. Chem. Phys., 135, 204105 (2011).

**(Larentzos1)** J.P. Larentzos, J.K. Brennan, J.D. Moore, M. Lisal and W.D. Mattson, “Parallel Implementation of Isothermal and Isoenergetic Dissipative Particle Dynamics Using Shardlow-Like Splitting Algorithms”, Comput. Phys. Commun., 185, 1987-1998 (2014).

**(Larentzos2)** J.P. Larentzos, J.K. Brennan, J.D. Moore, and W.D. Mattson, “LAMMPS Implementation of Constant Energy Dissipative Particle Dynamics (DPD-E)”, ARL-TR-6863, U.S. Army Research Laboratory, Aberdeen Proving Ground, MD (2014).

## 17.195 fix smd command

### 17.195.1 Syntax

<code>fix ID group-ID smd type values keyword values</code>
---

- ID, group-ID are documented in *fix* command
- smd = style name of this fix command
- mode = *cvel* or *cfor* to select constant velocity or constant force SMD
  - cvel* values = K vel
    - K = spring constant (force/distance units)
    - vel = velocity of pulling (distance/time units)
  - cfor* values = force
    - force = pulling force (force units)
- keyword = *tether* or *couple*
  - tether* values = x y z R0
    - x,y,z = point to which spring is tethered
    - R0 = distance of end of spring from tether point (distance units)
  - couple* values = group-ID2 x y z R0
    - group-ID2 = 2nd group to couple to fix group with a spring
    - x,y,z = direction of spring, automatically computed with 'auto'
    - R0 = distance of end of spring (distance units)

## 17.195.2 Examples

```

fix pull      cterm smd cvel 20.0 -0.00005 tether NULL NULL 100.0 0.0
fix pull      cterm smd cvel 20.0 -0.0001 tether 25.0 25 25.0 0.0
fix stretch cterm smd cvel 20.0 0.0001 couple nterm auto auto 0.0
fix pull      cterm smd cfor 5.0 tether 25.0 25.0 25.0 0.0

```

## 17.195.3 Description

This fix implements several options of steered MD (SMD) as reviewed in (*Izrailev*), which allows to induce conformational changes in systems and to compute the potential of mean force (PMF) along the assumed reaction coordinate (*Park*) based on Jarzynski's equality (*Jarzynski*). This fix borrows a lot from *fix spring* and *fix setforce*.

You can apply a moving spring force to a group of atoms (*tether* style) or between two groups of atoms (*couple* style). The spring can then be used in either constant velocity (*cvel*) mode or in constant force (*cfor*) mode to induce transitions in your systems. When running in *tether* style, you may need some way to fix some other part of the system (e.g. via *fix spring/self*)

The *tether* style attaches a spring between a point at a distance of  $R_0$  away from a fixed point  $x,y,z$  and the center of mass of the fix group of atoms. A restoring force of magnitude  $K (R - R_0) M_i / M$  is applied to each atom in the group where  $K$  is the spring constant,  $M_i$  is the mass of the atom, and  $M$  is the total mass of all atoms in the group. Note that  $K$  thus represents the total force on the group of atoms, not a per-atom force.

In *cvel* mode the distance  $R$  is incremented or decremented monotonously according to the pulling (or pushing) velocity. In *cfor* mode a constant force is added and the actual distance in direction of the spring is recorded.

The *couple* style links two groups of atoms together. The first group is the fix group; the second is specified by group-ID2. The groups are coupled together by a spring that is at equilibrium when the two groups are displaced by a vector in direction  $x,y,z$  with respect to each other and at a distance  $R_0$  from that displacement. Note that  $x,y,z$  only provides a direction and will be internally normalized. But since it represents the *absolute* displacement of group-ID2 relative to the fix group, (1,1,0) is a different spring than (-1,-1,0). For each vector component, the displacement can be described with the *auto* parameter. In this case the direction is re-computed in every step, which can be useful for steering a local process where the whole object undergoes some other change. When the relative positions and distance between the two groups are not in equilibrium, the same spring force described above is applied to atoms in each of the two groups.

For both the *tether* and *couple* styles, any of the  $x,y,z$  values can be specified as NULL which means do not include that dimension in the distance calculation or force application.

For constant velocity pulling (*cvel* mode), the running integral over the pulling force in direction of the spring is recorded and can then later be used to compute the potential of mean force (PMF) by averaging over multiple independent trajectories along the same pulling path.

## 17.195.4 Restart, fix\_modify, output, run start/stop, minimize info

The fix stores the direction of the spring, current pulling target distance and the running PMF to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify virial* option is supported by this fix to add the contribution due to the added forces on atoms to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a vector list of 7 quantities, which can be accessed by various *output commands*. The quantities in the vector are in this order: the x-, y-, and z-component of the pulling force, the total force in direction of the pull, the equilibrium distance of the spring, the distance between the two reference points, and finally the accumulated PMF (the sum of pulling forces times displacement).

The force is the total force on the group of atoms by the spring. In the case of the *couple* style, it is the force on the fix group (group-ID) or the negative of the force on the second group (group-ID2). The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.195.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.195.6 Related commands

*fix drag, fix spring, fix spring/self, fix spring/rg, fix colvars, fix plumed*

### 17.195.7 Default

none

---

**(Izrailev)** Izrailev, Stepaniants, Isralewitz, Kosztin, Lu, Molnar, Wriggers, Schulten. Computational Molecular Dynamics: Challenges, Methods, Ideas, volume 4 of Lecture Notes in Computational Science and Engineering, pp. 39-65. Springer-Verlag, Berlin, 1998.

**(Park)** Park, Schulten, J. Chem. Phys. 120 (13), 5946 (2004)

**(Jarzynski)** Jarzynski, Phys. Rev. Lett. 78, 2690 (1997)

## 17.196 fix smd/adjust\_dt command

### 17.196.1 Syntax

```
fix ID group-ID smd/adjust_dt arg
```

- ID, group-ID are documented in *fix* command
- smd/adjust\_dt = style name of this fix command
- arg = *s\_fact*  
*s\_fact* = safety factor

## 17.196.2 Examples

```
fix 1 all smd/adjust_dt 0.1
```

## 17.196.3 Description

The fix calculates a new stable time increment for use with the SMD time integrators.

The stable time increment is based on multiple conditions. For the SPH pair styles, a CFL criterion (Courant, Friedrichs & Lewy, 1928) is evaluated, which determines the speed of sound cannot propagate further than a typical spacing between particles within a single time step to ensure no information is lost. For the contact pair styles, a linear analysis of the pair potential determines a stable maximum time step.

This fix inquires the minimum stable time increment across all particles contained in the group for which this fix is defined. An additional safety factor *s\_fact* is applied to the time increment.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

## 17.196.4 Restart, fix\_modify, output, run start/stop, minimize info

Currently, no part of MACHDYN supports restarting nor minimization.

## 17.196.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 17.196.6 Related commands

*smd/tlsph\_dt*

## 17.196.7 Default

none

## 17.197 fix smd/integrate\_tlsph command

### 17.197.1 Syntax

```
fix ID group-ID smd/integrate_tlsph keyword values
```

- ID, group-ID are documented in *fix* command
- smd/integrate\_tlsph = style name of this fix command
- zero or more keyword/value pairs may be appended
- keyword = *limit\_velocity*

*limit\_velocity* value = *max\_vel*  
*max\_vel* = maximum allowed velocity

## 17.197.2 Examples

```
fix 1 all smd/integrate_tlsph  
fix 1 all smd/integrate_tlsph limit_velocity 1000
```

## 17.197.3 Description

The fix performs explicit time integration for particles which interact according with the Total-Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

The *limit\_velocity* keyword will control the velocity, scaling the norm of the velocity vector to *max\_vel* in case it exceeds this velocity limit.

## 17.197.4 Restart, fix\_modify, output, run start/stop, minimize info

Currently, no part of MACHDYN supports restarting nor minimization. This fix has no outputs.

## 17.197.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 17.197.6 Related commands

*smd/integrate\_ulsph*

## 17.197.7 Default

none

# 17.198 fix smd/integrate\_ulsph command

## 17.198.1 Syntax

```
fix ID group-ID smd/integrate_ulsph keyword
```

- ID, group-ID are documented in *fix* command
- smd/integrate\_ulsph = style name of this fix command
- zero or more keyword/value pairs may be appended
- keyword = *adjust\_radius* or *limit\_velocity*

**adjust\_radius values = adjust\_radius\_factor min\_nn max\_nn**

adjust\_radius\_factor = factor which scale the smooth/kernel radius  
 min\_nn = minimum number of neighbors  
 max\_nn = maximum number of neighbors

**limit\_velocity values = max\_velocity**

max\_velocity = maximum allowed velocity.

## 17.198.2 Examples

```
fix 1 all smd/integrate_ulsph adjust_radius 1.02 25 50
fix 1 all smd/integrate_ulsph limit_velocity 1000
```

## 17.198.3 Description

The fix performs explicit time integration for particles which interact with the updated Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

The *adjust\_radius* keyword activates dynamic adjustment of the per-particle SPH smoothing kernel radius such that the number of neighbors per particles remains within the interval *min\_nn* to *max\_nn*. The parameter *adjust\_radius\_factor* determines the amount of adjustment per timestep. Typical values are *adjust\_radius\_factor* =1.02, *min\_nn* =15, and *max\_nn* =20.

The *limit\_velocity* keyword will control the velocity, scaling the norm of the velocity vector to *max\_vel* in case it exceeds this velocity limit.

## 17.198.4 Restart, fix\_modify, output, run start/stop, minimize info

Currently, no part of MACHDYN supports restarting nor minimization. This fix has no outputs.

## 17.198.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 17.198.6 Related commands

none

## 17.198.7 Default

none

## 17.199 fix smd/move\_tri\_surf command

### 17.199.1 Syntax

```
fix ID group-ID smd/move_tri_surf keyword
```

- ID, group-ID are documented in *fix* command
- smd/move\_tri\_surf keyword = style name of this fix command
- keyword = *\*LINEAR* or *\*WIGGLE* or *\*ROTATE*
  - \*LINEAR* args = Vx Vy Vz  
Vx,Vy,Vz = components of velocity vector (velocity units), any component can be specified as NULL
  - \*WIGGLE* args = Vx Vy Vz max\_travel  
vx,vy,vz = components of velocity vector (velocity units), any component can be specified as NULL  
max\_travel = wiggle amplitude
  - \*ROTATE* args = Px Py Pz Rx Ry Rz period  
Px,Py,Pz = origin point of axis of rotation (distance units)  
Rx,Ry,Rz = axis of rotation vector  
period = period of rotation (time units)

### 17.199.2 Examples

```
fix 1 tool smd/move_tri_surf *LINEAR 20 20 10  
fix 2 tool smd/move_tri_surf *WIGGLE 20 20 10  
fix 2 tool smd/move_tri_surf *ROTATE 0 0 0 5 2 1
```

### 17.199.3 Description

This fix applies only to rigid surfaces read from .STL files via fix *smd/wall\_surface* . It updates position and velocity for the particles in the group each timestep without regard to forces on the particles. The rigid surfaces can thus be moved along simple trajectories during the simulation.

The *\*LINEAR* style moves particles with the specified constant velocity vector  $V = (V_x, V_y, V_z)$ . This style also sets the velocity of each particle to  $V = (V_x, V_y, V_z)$ .

The *\*WIGGLE* style moves particles in an oscillatory fashion. Particles are moved along  $(v_x, v_y, v_z)$  with constant velocity until a displacement of *max\_travel* is reached. Then, the velocity vector is reversed. This process is repeated.

The *\*ROTATE* style rotates particles around a rotation axis  $R = (R_x, R_y, R_z)$  that goes through a point  $P = (P_x, P_y, P_z)$ . The period of the rotation is also specified. This style also sets the velocity of each particle to  $(\omega \text{ cross } R_{\text{perp}})$  where  $\omega$  is its angular velocity around the rotation axis and  $R_{\text{perp}}$  is a perpendicular vector from the rotation axis to the particle.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.



### 17.199.4 Restart, fix\_modify, output, run start/stop, minimize info

Currently, no part of MACHDYN supports restarting nor minimization. This fix has no outputs.

### 17.199.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 17.199.6 Related commands

*smd/triangle\_mesh\_vertices, smd/wall\_surface*

### 17.199.7 Default

none

## 17.200 fix smd/setvel command

### 17.200.1 Syntax

```
fix ID group-ID smd/setvel vx vy vz keyword value ...
```

- ID, group-ID are documented in *fix* command
- smd/setvel = style name of this fix command
- vx,vy,vz = velocity component values
- any of vx,vy,vz can be a variable (see below)
- zero or more keyword/value pairs may be appended to args
- keyword = *region*

*region* value = region-ID

region-ID = ID of region particles must be in to have their velocities set

### 17.200.2 Examples

```
fix top_velocity top_group smd/setvel 1.0 0.0 0.0
```

### 17.200.3 Description

Set each component of velocity on each particle in the group to the specified values  $v_x, v_y, v_z$ , regardless of the forces acting on the particle. This command can be used to impose velocity boundary conditions.

Any of the  $v_x, v_y, v_z$  values can be specified as NULL which means do not alter the velocity component in that dimension.

This fix is indented to be used together with a time integration fix.

Any of the 3 quantities defining the velocity components can be specified as an equal-style or atom-style *variable*, namely  $v_x, v_y, v_z$ . If the value is a variable, it should be specified as  $v\_name$ , where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the force component.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent velocity field.

Atom-style variables can specify the same formulas as equal-style variables but can also include per-atom values, such as atom coordinates. Thus it is easy to specify a spatially-dependent velocity field with optional time-dependence as well.

If the *region* keyword is used, the particle must also be in the specified geometric *region* in order to have its velocity set by this command.

---

### 17.200.4 Restart, fix\_modify, output, run start/stop, minimize info

Currently, no part of MACHDYN supports restarting nor minimization. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global 3-vector of forces, which can be accessed by various *output commands*. This is the total force on the group of atoms. The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

### 17.200.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.200.6 Related commands

none

### 17.200.7 Default

none

## 17.201 fix smd/wall\_surface command

### 17.201.1 Syntax

```
fix ID group-ID smd/wall_surface arg type mol-ID
```

- ID, group-ID are documented in *fix* command
- smd/wall\_surface = style name of this fix command
- arg = *file*  
*file* = file name of a triangular mesh in stl format
- type = particle type to be given to the new particles created by this fix
- mol-ID = molecule-ID to be given to the new particles created by this fix (must be  $\geq 65535$ )

### 17.201.2 Examples

```
fix stl_surf all smd/wall_surface tool.stl 2 65535
```

### 17.201.3 Description

This fix creates reads a triangulated surface from a file in .STL format. For each triangle, a new particle is created which stores the barycenter of the triangle and the vertex positions. The radius of the new particle is that of the minimum circle which encompasses the triangle vertices.

The triangulated surface can be used as a complex rigid wall via the *smd/tri\_surface* pair style. It is possible to move the triangulated surface via the *smd/move\_tri\_surf* fix style.

Immediately after a .STL file has been read, the simulation needs to be run for 0 timesteps in order to properly register the new particles in the system. See the “funnel\_flow” example in the MACHDYN examples directory.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

### 17.201.4 Restart, fix\_modify, output, run start/stop, minimize info

Currently, no part of MACHDYN supports restarting nor minimization. This fix has no outputs.

### 17.201.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

The molecule ID given to the particles created by this fix have to be equal to or larger than 65535.

Within each .STL file, only a single triangulated object must be present, even though the STL format allows for the possibility of multiple objects in one file.

## 17.201.6 Related commands

*smd/triangle\_mesh\_vertices, smd/move\_tri\_surf, smd/tri\_surface*

## 17.201.7 Default

none

## 17.202 fix sph command

### 17.202.1 Syntax

```
fix ID group-ID sph
```

- ID, group-ID are documented in *fix* command
- sph = style name of this fix command

### 17.202.2 Examples

```
fix 1 all sph
```

### 17.202.3 Description

Perform time integration to update position, velocity, internal energy and local density for atoms in the group each timestep. This fix is needed to time-integrate SPH systems where particles carry internal variables such as internal energy. SPH stands for Smoothed Particle Hydrodynamics.

See [this PDF guide](#) to using SPH in LAMMPS.

### 17.202.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.202.5 Restrictions

This fix is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 17.202.6 Related commands

*fix sph/stationary*

## 17.202.7 Default

none

# 17.203 fix sph/stationary command

## 17.203.1 Syntax

```
fix ID group-ID sph/stationary
```

- ID, group-ID are documented in *fix* command
- sph = style name of this fix command

## 17.203.2 Examples

```
fix 1 boundary sph/stationary
```

## 17.203.3 Description

Perform time integration to update internal energy and local density, but not position or velocity for atoms in the group each timestep. This fix is needed for SPH simulations to correctly time-integrate fixed boundary particles which constrain a fluid to a given region in space. SPH stands for Smoothed Particle Hydrodynamics.

See [this PDF guide](#) to using SPH in LAMMPS.

## 17.203.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.203.5 Restrictions

This fix is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 17.203.6 Related commands

*fix sph*

## 17.203.7 Default

none

# 17.204 fix spring command

## 17.204.1 Syntax

```
fix ID group-ID spring keyword values
```

- ID, group-ID are documented in *fix* command
- spring = style name of this fix command
- keyword = *tether* or *couple*

*tether* values = K x y z R0

K = spring constant (force/distance units)

x,y,z = point to which spring is tethered

R0 = equilibrium distance from tether point (distance units)

*couple* values = group-ID2 K x y z R0

group-ID2 = 2nd group to couple to fix group with a spring

K = spring constant (force/distance units)

x,y,z = direction of spring

R0 = equilibrium distance of spring (distance units)

## 17.204.2 Examples

```
fix pull ligand spring tether 50.0 0.0 0.0 0.0 0.0
fix pull ligand spring tether 50.0 0.0 0.0 0.0 5.0
fix pull ligand spring tether 50.0 NULL NULL 2.0 3.0
fix 5 bilayer1 spring couple bilayer2 100.0 NULL NULL 10.0 0.0
fix longitudinal pore spring couple ion 100.0 NULL NULL -20.0 0.0
fix radial pore spring couple ion 100.0 0.0 0.0 NULL 5.0
```

## 17.204.3 Description

Apply a spring force to a group of atoms or between two groups of atoms. This is useful for applying an umbrella force to a small molecule or lightly tethering a large group of atoms (e.g. all the solvent or a large molecule) to the center of the simulation box so that it does not wander away over the course of a long simulation. It can also be used to hold the centers of mass of two groups of atoms at a given distance or orientation with respect to each other.

The *tether* style attaches a spring between a fixed point  $x,y,z$  and the center of mass of the fix group of atoms. The equilibrium position of the spring is  $R_0$ . At each timestep the distance  $R$  from the center of mass of the group of atoms to the tethering point is computed, taking account of wrap-around in a periodic simulation box. A restoring force of magnitude  $K (R - R_0) M_i / M$  is applied to each atom in the group where  $K$  is the spring constant,  $M_i$  is the mass of

the atom, and  $M$  is the total mass of all atoms in the group. Note that  $K$  thus represents the spring constant for the total force on the group of atoms, not for a spring applied to each atom.

The *couple* style links two groups of atoms together. The first group is the fix group; the second is specified by group-ID2. The groups are coupled together by a spring that is at equilibrium when the two groups are displaced by a vector  $x,y,z$  with respect to each other and at a distance  $R0$  from that displacement. Note that  $x,y,z$  is the equilibrium displacement of group-ID2 relative to the fix group. Thus (1,1,0) is a different spring than (-1,-1,0). When the relative positions and distance between the two groups are not in equilibrium, the same spring force described above is applied to atoms in each of the two groups.

For both the *tether* and *couple* styles, any of the  $x,y,z$  values can be specified as NULL which means do not include that dimension in the distance calculation or force application.

The first example above pulls the ligand towards the point (0,0,0). The second example holds the ligand near the surface of a sphere of radius 5 around the point (0,0,0). The third example holds the ligand a distance 3 away from the  $z=2$  plane (on either side).

The fourth example holds 2 bilayers a distance 10 apart in  $z$ . For the last two examples, imagine a pore (a slab of atoms with a cylindrical hole cut out) oriented with the pore axis along  $z$ , and an ion moving within the pore. The fifth example holds the ion a distance of -20 below the  $z = 0$  center plane of the pore (umbrella sampling). The last example holds the ion a distance 5 away from the pore axis (assuming the center-of-mass of the pore in  $x,y$  is the pore axis).

---

**Note:** The center of mass of a group of atoms is calculated in “unwrapped” coordinates using atom image flags, which means that the group can straddle a periodic boundary. See the *dump* doc page for a discussion of unwrapped coordinates. It also means that a spring connecting two groups or a group and the tether point can cross a periodic boundary and its length be calculated correctly.

---

#### 17.204.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the energy stored in the spring to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the spring energy  $= 0.5 * K * r^2$ .

This fix also computes global 4-vector which can be accessed by various *output commands*. The first 3 quantities in the vector are xyz components of the total force added to the group of atoms by the spring. In the case of the *couple* style, it is the force on the fix group (group-ID) or the negative of the force on the second group (group-ID2). The fourth quantity in the vector is the magnitude of the force added by the spring, as a positive value if  $(r-R0) > 0$  and a negative value if  $(r-R0) < 0$ . This sign convention can be useful when using the spring force to compute a potential of mean force (PMF).

The scalar and vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the spring energy to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

## 17.204.5 Restrictions

none

## 17.204.6 Related commands

*fix drag*, *fix spring/self*, *fix spring/rg*, *fix smd*

## 17.204.7 Default

none

# 17.205 fix spring/chunk command

## 17.205.1 Syntax

```
fix ID group-ID spring/chunk K chunkID comID
```

- ID, group-ID are documented in *fix* command
- spring/chunk = style name of this fix command
- K = spring constant for each chunk (force/distance units)
- chunkID = ID of *compute chunk/atom* command
- comID = ID of *compute com/chunk* command

## 17.205.2 Examples

```
fix restrain all spring/chunk 100 chunkID comID
```

## 17.205.3 Description

Apply a spring force to the center-of-mass (COM) of chunks of atoms as defined by the *compute chunk/atom* command. Chunks can be molecules or spatial bins or other groupings of atoms. This is a way of tethering each chunk to its initial COM coordinates.

The *chunkID* is the ID of a compute chunk/atom command defined in the input script. It is used to define the chunks. The *comID* is the ID of a compute com/chunk command defined in the input script. It is used to compute the COMs of each chunk.

At the beginning of the first *run* or *minimize* command after this fix is defined, the initial COM of each chunk is calculated and stored as  $R_{0m}$ , where M is the chunk number. Thereafter, at every timestep (or minimization iteration), the current COM of each chunk is calculated as  $R_m$ . A restoring force of magnitude  $K (R_m - R_{0m}) M_i / M_m$  is applied to each atom in each chunk where  $K$  is the specified spring constant,  $M_i$  is the mass of the atom, and  $M_m$  is the total mass of all atoms in the chunk. Note that  $K$  thus represents the spring constant for the total force on each chunk of atoms, not for a spring applied to each atom.



## 17.205.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the locations of the initial per-chunk center of mass coordinates to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the fix continues in an uninterrupted fashion. Since this fix depends on an instance of *compute chunk/atom* it will check when reading the restart if the chunk still exists and will define the same number of chunks. The restart data is only applied when the number of chunks matches. Otherwise the center of mass coordinates are recomputed.

The *fix\_modify energy* option is supported by this fix to add the energy stored in all the springs to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the energy of all the springs, i.e.  $0.5 * K * r^2$  per-spring.

The scalar value calculated by this fix is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the spring energies to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

## 17.205.5 Restrictions

none

## 17.205.6 Related commands

*fix spring*, *fix spring/self*, *fix spring/rg*

## 17.205.7 Default

none

## 17.206 fix spring/rg command

### 17.206.1 Syntax

```
fix ID group-ID spring/rg K RG0
```

- ID, group-ID are documented in *fix* command
- spring/rg = style name of this fix command
- K = harmonic force constant (force/distance units)
- RG0 = target radius of gyration to constrain to (distance units)

```
if RG0 = NULL, use the current RG as the target value
```

## 17.206.2 Examples

```
fix 1 protein spring/rg 5.0 10.0
fix 2 micelle spring/rg 5.0 NULL
```

## 17.206.3 Description

Apply a harmonic restraining force to atoms in the group to affect their central moment about the center of mass (radius of gyration). This fix is useful to encourage a protein or polymer to fold/unfold and also when sampling along the radius of gyration as a reaction coordinate (i.e. for protein folding).

The radius of gyration is defined as RG in the first formula. The energy of the constraint and associated force on each atom is given by the second and third formulas, when the group is at a different RG than the target value RG0.

$$R_G^2 = \frac{1}{M} \sum_i^N m_i \left( x_i - \frac{1}{M} \sum_j^N m_j x_j \right)^2$$

$$E = K (R_G - R_{G0})^2$$

$$F_i = 2K \frac{m_i}{M} \left( 1 - \frac{R_{G0}}{R_G} \right) \left( x_i - \frac{1}{M} \sum_j^N m_j x_j \right)$$

The ( $x_i$  - center-of-mass) term is computed taking into account periodic boundary conditions,  $m_i$  is the mass of the atom, and  $M$  is the mass of the entire group. Note that  $K$  is thus a force constant for the aggregate force on the group of atoms, not a per-atom force.

If  $R_{G0}$  is specified as NULL, then the RG of the group is computed at the time the fix is specified, and that value is used as the target.

## 17.206.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the currently used reference RG ( $R_{G0}$ ) to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the fix continues in an uninterrupted fashion.

None of the *fix\_modify* options are relevant to this fix.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the reference radius of gyration  $R_{G0}$  used by the fix. energy change due to this fix. The scalar value calculated by this fix is “intensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

## 17.206.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 17.206.6 Related commands

*fix spring, fix spring/self fix drag, fix smd*

## 17.206.7 Default

none

## 17.207 fix spring/self command

### 17.207.1 Syntax

```
fix ID group-ID spring/self K dir
```

- ID, group-ID are documented in *fix* command
- spring/self = style name of this fix command
- K = spring constant (force/distance units)
- dir = xyz, xy, xz, yz, x, y, or z (optional, default: xyz)

### 17.207.2 Examples

```
fix tether boundary-atoms spring/self 10.0
fix zrest move spring/self 10.0 z
```

### 17.207.3 Description

Apply a spring force independently to each atom in the group to tether it to its initial position. The initial position for each atom is its location at the time the fix command was issued. At each timestep, the magnitude of the force on each atom is  $-Kr$ , where  $r$  is the displacement of the atom from its current position to its initial position. The distance  $r$  correctly takes into account any crossings of periodic boundary by the atom since it was in its initial position.

With the (optional) dir flag, one can select in which direction the spring force is applied. By default, the restraint is applied in all directions, but it can be limited to the xy-, xz-, yz-plane and the x-, y-, or z-direction, thus restraining the atoms to a line or a plane, respectively.

## 17.207.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the original coordinates of tethered atoms to *binary restart files*, so that the spring effect will be the same in a restarted simulation. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify energy* option is supported by this fix to add the energy stored in the per-atom springs to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is an energy which is the sum of the spring energy for each atom, where the per-atom energy is  $0.5 * K * r^2$ . The scalar value calculated by this fix is “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the per-atom spring energy to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

## 17.207.5 Restrictions

none

## 17.207.6 Related commands

*fix drag*, *fix spring*, *fix smd*, *fix spring/rg*

## 17.207.7 Default

none

## 17.208 fix srd command

### 17.208.1 Syntax

```
fix ID group-ID srd N groupbig-ID Tsrđ hgrid seed keyword value ...
```

- ID, group-ID are documented in *fix* command
- srd = style name of this fix command
- N = reset SRD particle velocities every this many timesteps
- groupbig-ID = ID of group of large particles that SRDs interact with
- Tsrđ = temperature of SRD particles (temperature units)
- hgrid = grid spacing for SRD grouping (distance units)

- seed = random # seed (positive integer)
- zero or more keyword/value pairs may be appended
- keyword = *lamda* or *collision* or *overlap* or *inside* or *exact* or *radius* or *bounce* or *search* or *cubic* or *shift* or *tstat* or *rescale*

*lamda* value = mean free path of SRD particles (distance units)  
*collision* value = *noslip* or *slip* = collision model  
*overlap* value = *yes* or *no* = whether big particles may overlap  
*inside* value = *error* or *warn* or *ignore* = how SRD particles which end up inside a  
↳big particle are treated  
*exact* value = *yes* or *no*  
*radius* value = *rfactor* = scale collision radius by this factor  
*bounce* value = *Nbounce* = max # of collisions an SRD particle can undergo in one  
↳timestep  
*search* value = *sgrid* = grid spacing for collision partner searching (distance units)  
*cubic* values = style tolerance  
style = *error* or *warn*  
tolerance = fractional difference allowed (0 <= tol <= 1)  
*shift* values = flag shiftseed  
flag = *yes* or *no* or *possible* = SRD bin shifting for better statistics  
yes = perform bin shifting each time SRD velocities are rescaled  
no = no shifting  
*possible* = shift depending on mean free path and bin size  
shiftseed = random # seed (positive integer)  
*tstat* value = *yes* or *no* = thermostat SRD particles or not  
*rescale* value = *yes* or *no* or *rotate* or *collide* = rescaling of SRD velocities  
yes = rescale during velocity rotation and collisions  
no = no rescaling  
*rotate* = rescale during velocity rotation, but not collisions  
*collide* = rescale during collisions, but not velocity rotation

## 17.208.2 Examples

```
fix 1 srd srd 10 big 1.0 0.25 482984
fix 1 srd srd 10 big 0.5 0.25 482984 collision slip search 0.5
```

## 17.208.3 Description

Treat a group of particles as stochastic rotation dynamics (SRD) particles that serve as a background solvent when interacting with big (colloidal) particles in groupbig-ID. The SRD formalism is described in ([Hecht](#)). The key idea behind using SRD particles as a cheap coarse-grained solvent is that SRD particles do not interact with each other, but only with the solute particles, which in LAMMPS can be spheroids, ellipsoids, or line segments, or triangles, or rigid bodies containing multiple spheroids or ellipsoids or line segments or triangles. The collision and rotation properties of the model imbue the SRD particles with fluid-like properties, including an effective viscosity. Thus simulations with large solute particles can be run more quickly, to measure solute properties like diffusivity and viscosity in a background fluid. The usual LAMMPS fixes for such simulations, such as *fix deform*, *fix viscosity*, and *fix nvt/sllod*, can be used in conjunction with the SRD model.

For more details on how the SRD model is implemented in LAMMPS, [this paper](#) describes the implementation and usage of pure SRD fluids. [This paper](#), which is nearly complete, describes the implementation and usage of mixture systems (solute particles in an SRD fluid). See the examples/srd directory for sample input scripts using SRD particles in both settings.

This fix does two things:

1. It advects the SRD particles, performing collisions between SRD and big particles or walls every timestep, imparting force and torque to the big particles. Collisions also change the position and velocity of SRD particles.
2. It resets the velocity distribution of SRD particles via random rotations every N timesteps.

SRD particles have a mass, temperature, characteristic timestep  $dt_{SRD}$ , and mean free path between collisions ( $\lambda$ ). The fundamental equation relating these 4 quantities is

$$\lambda = dt_{SRD} \sqrt{\frac{k_B T_{SRD}}{m}}$$

The mass  $m$  of SRD particles is set by the *mass* command elsewhere in the input script. The SRD timestep  $dt_{SRD}$  is N times the step  $dt$  defined by the *timestep* command. Big particles move in the normal way via a time integration *fix* with a short timestep  $dt$ . SRD particles advect with a large timestep  $dt_{SRD} \geq dt$ .

If the *lamda* keyword is not specified, the SRD temperature  $T_{SRD}$  is used in the above formula to compute  $\lambda$ . If the *lamda* keyword is specified, then the *Tsrd* setting is ignored and the above equation is used to compute the SRD temperature.

The characteristic length scale for the SRD fluid is set by *hgrid* which is used to bin SRD particles for purposes of resetting their velocities. Normally *hgrid* is set to be 1/4 of the big particle diameter or smaller, to adequately resolve fluid properties around the big particles.

$\lambda$  cannot be smaller than  $0.6 * hgrid$ , else an error is generated (unless the *shift* keyword is used, see below). The velocities of SRD particles are bounded by  $V_{max}$ , which is set so that an SRD particle will not advect further than  $D_{max} = 4\lambda$  in  $dt_{SRD}$ . This means that roughly speaking,  $D_{max}$  should not be larger than a big particle diameter, else SRDs may pass through big particles without colliding. A warning is generated if this is the case.

Collisions between SRD particles and big particles or walls are modeled as a lightweight SRD point particle hitting a heavy big particle of given diameter or a wall at a point on its surface and bouncing off with a new velocity. The collision changes the momentum of the SRD particle. It imparts a force and torque to the big particle. It imparts a force to a wall. Static or moving SRD walls are setup via the *fix wall/srd* command. For the remainder of this doc page, a collision of an SRD particle with a wall can be viewed as a collision with a big particle of infinite radius and mass.

The *collision* keyword sets the style of collisions. The *slip* style means that the tangential component of the SRD particle momentum is preserved. Thus a force is imparted to a big particle, but no torque. The normal component of the new SRD velocity is sampled from a Gaussian distribution at temperature *Tsrd*.

For the *noslip* style, both the normal and tangential components of the new SRD velocity are sampled from a Gaussian distribution at temperature *Tsrd*. Additionally, a new tangential direction for the SRD velocity is chosen randomly. This collision style imparts torque to a big particle. Thus a time integrator *fix* that rotates the big particles appropriately should be used.

---

The *overlap* keyword should be set to *yes* if two (or more) big particles can ever overlap. This depends on the pair potential interaction used for big-big interactions, or could be the case if multiple big particles are held together as rigid bodies via the *fix rigid* command. If the *overlap* keyword is *no* and big particles do in fact overlap, then SRD/big collisions can generate an error if an SRD ends up inside two (or more) big particles at once. How this error is treated is determined by the *inside* keyword. Running with *overlap* set to *no* allows for faster collision checking, so it should only be set to *yes* if needed.

The *inside* keyword determines how a collision is treated if the computation determines that the timestep started with the SRD particle already inside a big particle. If the setting is *error* then this generates an error message and LAMMPS stops. If the setting is *warn* then this generates a warning message and the code continues. If the setting is *ignore* then no message is generated. One of the output quantities logged by the *fix* (see below) tallies the number of such events, so it can be monitored. Note that once an SRD particle is inside a big particle, it may remain there for several steps until it drifts outside the big particle.

The *exact* keyword determines how accurately collisions are computed. A setting of *yes* computes the time and position of each collision as SRD and big particles move together. A setting of *no* estimates the position of each collision based on the end-of-timestep positions of the SRD and big particle. If *overlap* is set to *yes*, the setting of the *exact* keyword is ignored since time-accurate collisions are needed.

The *radius* keyword scales the effective size of big particles. If big particles will overlap as they undergo dynamics, then this keyword can be used to scale down their effective collision radius by an amount *rfactor*, so that SRD particle will only collide with one big particle at a time. For example, in a Lennard-Jones system at a temperature of 1.0 (in reduced LJ units), the minimum separation between two big particles is as small as about 0.88 sigma. Thus an *rfactor* value of 0.85 should prevent dual collisions.

The *bounce* keyword can be used to limit the maximum number of collisions an SRD particle undergoes in a single timestep as it bounces between nearby big particles. Note that if the limit is reached, the SRD can be left inside a big particle. A setting of 0 is the same as no limit.

---

There are 2 kinds of bins created and maintained when running an SRD simulation. The first are “SRD bins” which are used to bin SRD particles and reset their velocities, as discussed above. The second are “search bins” which are used to identify SRD/big particle collisions.

The *search* keyword can be used to choose a search bin size for identifying SRD/big particle collisions. The default is to use the *hgrid* parameter for SRD bins as the search bin size. Choosing a smaller or large value may be more efficient, depending on the problem. But, in a statistical sense, it should not change the simulation results.

The *cubic* keyword can be used to generate an error or warning when the bin size chosen by LAMMPS creates SRD bins that are non-cubic or different than the requested value of *hgrid* by a specified *tolerance*. Note that using non-cubic SRD bins can lead to undetermined behavior when rotating the velocities of SRD particles, hence LAMMPS tries to protect you from this problem.

LAMMPS attempts to set the SRD bin size to exactly *hgrid*. However, there must be an integer number of bins in each dimension of the simulation box. Thus the actual bin size will depend on the size and shape of the overall simulation box. The actual bin size is printed as part of the SRD output when a simulation begins.

If the actual bin size is non-cubic by an amount exceeding the tolerance, an error or warning is printed, depending on the style of the *cubic* keyword. Likewise, if the actual bin size differs from the requested *hgrid* value by an amount exceeding the tolerance, then an error or warning is printed. The *tolerance* is a fractional difference. E.g. a tolerance setting of 0.01 on the shape means that if the ratio of any 2 bin dimensions exceeds (1 +/- tolerance) then an error or warning is generated. Similarly, if the ratio of any bin dimension with *hgrid* exceeds (1 +/- tolerance), then an error or warning is generated.

---

**Note:** The `fix srd` command can be used with simulations where the size and/or shape of the simulation box changes. This can be due to non-periodic boundary conditions or the use of fixes such as the `fix deform` or `fix wall/srd` commands to impose a shear on an SRD fluid or an interaction with an external wall. If the box size changes then the size of SRD bins must be recalculated every reneighboring. This is not necessary if only the box shape changes. This re-binning is always done so as to fit an integer number of bins in the current box dimension, whether it be a fixed, shrink-wrapped, or periodic boundary, as set by the `boundary` command. If the box size or shape changes, then the size of the search bins must be recalculated every reneighboring. Note that changing the SRD bin size may alter the properties of the SRD fluid, such as its viscosity.

---

The *shift* keyword determines whether the coordinates of SRD particles are randomly shifted when binned for purposes of rotating their velocities. When no shifting is performed, SRD particles are binned and the velocity distribution of the set of SRD particles in each bin is adjusted via a rotation operator. This is a statistically valid operation if SRD particles move sufficiently far between successive rotations. This is determined by their mean-free path  $\lambda$ . If  $\lambda$  is less than 0.6 of the SRD bin size, then shifting is required. A shift means that all of the SRD particles are shifted by a vector whose coordinates are chosen randomly in the range [-1/2 bin size, 1/2 bin size]. Note that all particles are shifted by

the same vector. The specified random number *shiftseed* is used to generate these vectors. This operation sufficiently randomizes which SRD particles are in the same bin, even if *lambda* is small.

If the *shift* flag is set to *no*, then no shifting is performed, but bin data will be communicated if bins overlap processor boundaries. An error will be generated if  $\lambda < 0.6$  of the SRD bin size. If the *shift* flag is set to *possible*, then shifting is performed only if  $\lambda < 0.6$  of the SRD bin size. A warning is generated to let you know this is occurring. If the *shift* flag is set to *yes* then shifting is performed regardless of the magnitude of  $\lambda$ . Note that the *shiftseed* is not used if the *shift* flag is set to *no*, but must still be specified.

Note that shifting of SRD coordinates requires extra communication, hence it should not normally be enabled unless required.

The *tstat* keyword will thermostat the SRD particles to the specified *Tsrd*. This is done every N timesteps, during the velocity rotation operation, by rescaling the thermal velocity of particles in each SRD bin to the desired temperature. If there is a streaming velocity associated with the system, e.g. due to use of the *fix deform* command to perform a simulation undergoing shear, then that is also accounted for. The mean velocity of each bin of SRD particles is set to the position-dependent streaming velocity, based on the coordinates of the center of the SRD bin. Note that collisions of SRD particles with big particles or walls has a thermostating effect on the colliding particles, so it may not be necessary to thermostat the SRD particles on a bin by bin basis in that case. Also note that for streaming simulations, if no thermostating is performed (the default), then it may take a long time for the SRD fluid to come to equilibrium with a velocity profile that matches the simulation box deformation.

The *rescale* keyword enables rescaling of an SRD particle's velocity if it would travel more than 4 mean-free paths in an SRD timestep. If an SRD particle exceeds this velocity it is possible it will be lost when migrating to other processors or that collisions with big particles will be missed, either of which will generate errors. Thus the safest mode is to run with rescaling enabled. However rescaling removes kinetic energy from the system (the particle's velocity is reduced). The latter will not typically be a problem if thermostating is enabled via the *tstat* keyword or if SRD collisions with big particles or walls effectively thermostat the system. If you wish to turn off rescaling (on is the default), e.g. for a pure SRD system with no thermostating so that the temperature does not decline over time, the *rescale* keyword can be used. The *no* value turns rescaling off during collisions and the per-bin velocity rotation operation. The *collide* and *rotate* values turn it on for one of the operations and off for the other.

---

**Note:** This fix is normally used for simulations with a huge number of SRD particles relative to the number of big particles, e.g. 100 to 1. In this scenario, computations that involve only big particles (neighbor list creation, communication, time integration) can slow down dramatically due to the large number of background SRD particles.

---

Three other input script commands will largely overcome this effect, speeding up an SRD simulation by a significant amount. These are the *atom\_modify first*, *neigh\_modify include*, and *comm\_modify group* commands. Each takes a group-ID as an argument, which in this case should be the group-ID of the big solute particles.

Additionally, when a *pair\_style* for big/big particle interactions is specified, the *pair\_coeff* command should be used to turn off big/SRD interactions, e.g. by setting their epsilon or cutoff length to 0.0.

The “delete\_atoms overlap” command may be useful in setting up an SRD simulation to insure there are no initial overlaps between big and SRD particles.

---



### 17.208.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix tabulates several SRD statistics which are stored in a vector of length 12, which can be accessed by various *output commands*. The vector values calculated by this fix are “intensive”, meaning they do not scale with the size of the simulation. Technically, the first 8 do scale with the size of the simulation, but treating them as intensive means they are not scaled when printed as part of thermodynamic output.

These are the 12 quantities. All are values for the current timestep, except for quantity 5 and the last three, each of which are cumulative quantities since the beginning of the run.

- (1) # of SRD/big collision checks performed
- (2) # of SRDs which had a collision
- (3) # of SRD/big collisions (including multiple bounces)
- (4) # of SRD particles inside a big particle
- (5) # of SRD particles whose velocity was rescaled to be < Vmax
- (6) # of bins for collision searching
- (7) # of bins for SRD velocity rotation
- (8) # of bins in which SRD temperature was computed
- (9) SRD temperature
- (10) # of SRD particles which have undergone max # of bounces
- (11) max # of bounces any SRD particle has had in a single step
- (12) # of reneighboring due to SRD particles moving too far

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.208.5 Restrictions

This command can only be used if LAMMPS was built with the SRD package. See the *Build package* doc page for more info.

### 17.208.6 Related commands

*fix wall/srd*

### 17.208.7 Default

The option defaults are: *lamda* ( $\lambda$ ) is inferred from *Tsrd*, collision = noslip, overlap = no, inside = error, exact = yes, radius = 1.0, bounce = 0, search = hgrid, cubic = error 0.01, shift = no, tstat = no, and rescale = yes.

---

**(Hecht)** Hecht, Harting, Ihle, Herrmann, Phys Rev E, 72, 011408 (2005).

**(Petersen)** Petersen, Lechman, Plimpton, Grest, in’ t Veld, Schunk, J Chem Phys, 132, 174106 (2010).

**(Lechman)** Lechman, et al, in preparation (2010).

## 17.209 fix store/force command

### 17.209.1 Syntax

```
fix ID group-ID store/force
```

- ID, group-ID are documented in *fix* command
- store/force = style name of this fix command

### 17.209.2 Examples

```
fix 1 all store/force
```

### 17.209.3 Description

Store the forces on atoms in the group at the point during each timestep when the fix is invoked, as described below. This is useful for storing forces before constraints or other boundary conditions are computed which modify the forces, so that unmodified forces can be *written to a dump file* or accessed by other *output commands* that use per-atom quantities.

This fix is invoked at the point in the velocity-Verlet timestepping immediately after *pair*, *bond*, *angle*, *dihedral*, *improper*, and *long-range* forces have been calculated. It is the point in the timestep when various fixes that compute constraint forces are calculated and potentially modify the force on each atom. Examples of such fixes are *fix shake*, *fix wall*, and *fix indent*.

---

**Note:** The order in which various fixes are applied which operate at the same point during the timestep, is the same as the order they are specified in the input script. Thus normally, if you want to store per-atom forces due to force field interactions, before constraints are applied, you should list this fix first within that set of fixes, i.e. before other fixes that apply constraints. However, if you wish to include certain constraints (e.g. *fix shake*) in the stored force, then it could be specified after some fixes and before others.

---

### 17.209.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix produces a per-atom array which can be accessed by various *output commands*. The number of columns for each atom is 3, and the columns store the x,y,z forces on each atom. The per-atom values be accessed on any timestep.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.209.5 Restrictions

none

## 17.209.6 Related commands

*fix store\_state*

## 17.209.7 Default

none

## 17.210 fix store/state command

### 17.210.1 Syntax

```
fix ID group-ID store/state N input1 input2 ... keyword value ...
```

- ID, group-ID are documented in *fix* command
- store/state = style name of this fix command
- N = store atom attributes every N steps, N = 0 for initial store only
- input = one or more atom attributes

```
possible attributes = id, mol, type, mass,
                    x, y, z, xs, ys, zs, xu, yu, zu, xsu, ysu, zsu, ix, iy, iz,
                    vx, vy, vz, fx, fy, fz,
                    q, mux, muy, muz, mu,
                    radius, diameter, omegax, omegay, omegaz,
                    angmomx, angmomy, angmomz, tqx, tqy, tqz,
                    c_ID, c_ID[I], f_ID, f_ID[I], v_name,
                    d_name, i_name, i2_name[I], d2_name[I],
```

```
id = atom ID
mol = molecule ID
type = atom type
mass = atom mass
x,y,z = unscaled atom coordinates
xs,ys,zs = scaled atom coordinates
xu,yu,zu = unwrapped atom coordinates
xsu,ysu,zsu = scaled unwrapped atom coordinates
ix,iy,iz = box image that the atom is in
vx,vy,vz = atom velocities
fx,fy,fz = forces on atoms
q = atom charge
mux,muy,muz = orientation of dipolar atom
mu = magnituded of dipole moment of atom
radius,diameter = radius.diameter of spherical particle
omegax,omegay,omegaz = angular velocity of spherical particle
angmomx,angmomy,angmomz = angular momentum of aspherical particle
```

`tqx,tqy,tqz` = torque on finite-size particles  
`c_ID` = per-atom vector calculated by a compute with ID  
`c_ID[I]` = Ith column of per-atom array calculated by a compute with ID  
`f_ID` = per-atom vector calculated by a fix with ID  
`f_ID[I]` = Ith column of per-atom array calculated by a fix with ID  
`v_name` = per-atom vector calculated by an atom-style variable with name  
`i_name` = custom integer vector with name  
`d_name` = custom floating point vector with name  
`i2_name[I]` = Ith column of custom integer array with name  
`d2_name[I]` = Ith column of custom floating-point array with name

- zero or more keyword/value pairs may be appended
- keyword = `com`  
`com` value = `yes` or `no`

## 17.210.2 Examples

```
fix 1 all store/state 0 x y z
fix 1 all store/state 0 xu yu zu com yes
fix 2 all store/state 1000 vx vy vz
```

## 17.210.3 Description

Define a fix that stores attributes for each atom in the group at the time the fix is defined. If  $N$  is 0, then the values are never updated, so this is a way of archiving an atom attribute at a given time for future use in a calculation or output. See the discussion of *output commands* that take fixes as inputs.

If  $N$  is not zero, then the attributes will be updated every  $N$  steps.

---

**Note:** Actually, only atom attributes specified by keywords like `xu` or `vy` or `radius` are initially stored immediately at the point in your input script when the fix is defined. Attributes specified by a compute, fix, or variable are not initially stored until the first run following the fix definition begins. This is because calculating those attributes may require quantities that are not defined in between runs.

---

The list of possible attributes is the same as that used by the *dump custom* command, which describes their meaning.

If the `com` keyword is set to `yes` then the `xu`, `yu`, and `zu` inputs store the position of each atom relative to the center-of-mass of the group of atoms, instead of storing the absolute position.

The requested values are stored in a per-atom vector or array as discussed below. Zeroes are stored for atoms not in the specified group.

### 17.210.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the per-atom values it stores to *binary restart files*, so that the values can be restored when a simulation is restarted. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

**Warning:** When reading data from a restart file, this fix command has to be specified **exactly** the same way as before. LAMMPS will only check whether a fix is of the same style and has the same fix ID and in case of a match will then try to initialize the fix with the data stored in the binary restart file. If the fix store/state command does not match exactly, data can be corrupted or LAMMPS may crash.

None of the *fix\_modify* options are relevant to this fix.

If a single input is specified, this fix produces a per-atom vector. If multiple inputs are specified, a per-atom array is produced where the number of columns for each atom is the number of inputs. These can be accessed by various *output commands*. The per-atom values be accessed on any timestep.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.210.5 Restrictions

none

### 17.210.6 Related commands

*dump custom, compute property/atom, fix property/atom, variable*

### 17.210.7 Default

The option default is com = no.

## 17.211 fix temp/berendsen command

### 17.211.1 Syntax

```
fix ID group-ID temp/berendsen Tstart Tstop Tdamp
```

- ID, group-ID are documented in *fix* command
- temp/berendsen = style name of this fix command
- Tstart,Tstop = desired temperature at start/end of run

Tstart can be a variable (see below)

- Tdamp = temperature damping parameter (time units)

## 17.211.2 Examples

```
fix 1 all temp/berendsen 300.0 300.0 100.0
```

---

## 17.211.3 Description

Reset the temperature of a group of atoms by using a Berendsen thermostat (*Berendsen*), which rescales their velocities every timestep.

The thermostat is applied to only the translational degrees of freedom for the particles, which is an important consideration for finite-size particles which have rotational degrees of freedom are being thermostatted with this fix. The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

The desired temperature at each timestep is a ramped value during the run from  $T_{start}$  to  $T_{stop}$ . The  $T_{damp}$  parameter is specified in time units and determines how rapidly the temperature is relaxed. For example, a value of 100.0 means to relax the temperature in a timespan of (roughly) 100 time units (tau or fs or ps - see the *units* command).

$T_{start}$  can be specified as an equal-style *variable*. In this case, the  $T_{stop}$  setting is ignored. If the value is a variable, it should be specified as  $v\_name$ , where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the target temperature.

---

**Note:** This thermostat will generate an error if the current temperature is zero at the end of a timestep. It cannot rescale a zero temperature.

---

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent temperature.

---

**Note:** Unlike the *fix nvt* command which performs Nose/Hoover thermostating AND time integration, this fix does NOT perform time integration. It only modifies velocities to effect thermostating. Thus you must use a separate time integration fix, like *fix nve* to actually update the positions of atoms using the modified velocities. Likewise, this fix should not normally be used on atoms that also have their temperature controlled by another fix - e.g. by *fix nvt* or *fix langevin* commands.

---

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostating.

This fix computes a temperature each timestep. To do this, the fix creates its own compute of style “temp”, as if this command had been issued:

```
compute fix-ID_temp group-ID temp
```

---

See the *compute temp* command for details. Note that the ID of the new compute is the fix-ID + underscore + “temp”, and the group for the new compute is the same as the fix group.

Note that this is NOT the compute used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp*. This means you can change the attributes of this fix’s temperature (e.g. its degrees-of-freedom) via the *compute\_modify* command or print this temperature during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

#### 17.211.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the cumulative global energy change to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the fix continues in an uninterrupted fashion.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a temperature *compute* you have defined to this fix which will be used in its thermostating procedure, as described above. For consistency, the group used by this fix and by the compute should be the same.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

#### 17.211.5 Restrictions

This fix can be used with dynamic groups as defined by the *group* command. Likewise it can be used with groups to which atoms are added or deleted over time, e.g. a deposition simulation. However, the conservation properties of the thermostat and barostat are defined for systems with a static set of atoms. You may observe odd behavior if the atoms in a group vary dramatically over time or the atom count becomes very small.

#### 17.211.6 Related commands

*fix nve*, *fix nvt*, *fix temp/rescale*, *fix langevin*, *fix\_modify*, *compute temp*, *fix press/berendsen*

#### 17.211.7 Default

none

---

**(Berendsen)** Berendsen, Postma, van Gunsteren, DiNola, Haak, J Chem Phys, 81, 3684 (1984).

## 17.212 fix temp/csvr command

## 17.213 fix temp/csld command

### 17.213.1 Syntax

```
fix ID group-ID temp/csvr Tstart Tstop Tdamp seed
fix ID group-ID temp/csld Tstart Tstop Tdamp seed
```

- ID, group-ID are documented in *fix* command
- temp/csvr or temp/csld = style name of this fix command
- Tstart, Tstop = desired temperature at start/end of run

```
Tstart can be a variable (see below)
```

- Tdamp = temperature damping parameter (time units)
- seed = random number seed to use for white noise (positive integer)

### 17.213.2 Examples

```
fix 1 all temp/csvr 300.0 300.0 100.0 54324
fix 1 all temp/csld 100.0 300.0 10.0 123321
```

### 17.213.3 Description

Adjust the temperature with a canonical sampling thermostat that uses global velocity rescaling with Hamiltonian dynamics (*temp/csvr*) (*Bussi1*), or Langevin dynamics (*temp/csld*) (*Bussi2*). In the case of *temp/csvr* the thermostat is similar to the empirical Berendsen thermostat in *temp/berendsen*, but chooses the actual scaling factor from a suitably chosen (gaussian) distribution rather than having it determined from the time constant directly. In the case of *temp/csld* the velocities are updated to a linear combination of the current velocities with a gaussian distribution of velocities at the desired temperature. Both thermostats are applied every timestep.

The thermostat is applied to only the translational degrees of freedom for the particles, which is an important consideration for finite-size particles which have rotational degrees of freedom are being thermostatted with these fixes. The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

The desired temperature at each timestep is a ramped value during the run from *Tstart* to *Tstop*. The *Tdamp* parameter is specified in time units and determines how rapidly the temperature is relaxed. For example, a value of 100.0 means to relax the temperature in a timespan of (roughly) 100 time units (tau or fs or ps - see the *units* command).

*Tstart* can be specified as an equal-style *variable*. In this case, the *Tstop* setting is ignored. If the value is a variable, it should be specified as *v\_name*, where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the target temperature.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent temperature.



---

**Note:** Unlike the *fix nvt* command which performs Nose/Hoover thermostating AND time integration, these fixes do NOT perform time integration. They only modify velocities to effect thermostating. Thus you must use a separate time integration fix, like *fix nve* to actually update the positions of atoms using the modified velocities. Likewise, these fixes should not normally be used on atoms that also have their temperature controlled by another fix - e.g. by *fix nvt* or *fix langevin* commands.

---

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostating. These fixes compute a temperature each timestep. To do this, the fix creates its own compute of style “temp”, as if this command had been issued:

```
compute fix-ID_temp group-ID temp
```

See the *compute temp* command for details. Note that the ID of the new compute is the fix-ID + underscore + “temp”, and the group for the new compute is the same as the fix group.

Note that this is NOT the compute used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp*. This means you can change the attributes of this fix’s temperature (e.g. its degrees-of-freedom) via the *compute\_modify* command or print this temperature during thermodynamic output via the *thermo\_style custom* command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

An important feature of these thermostats is that they have an associated effective energy that is a constant of motion. The effective energy is the total energy (kinetic + potential) plus the accumulated kinetic energy changes due to the thermostat. The latter quantity is the global scalar computed by these fixes. This feature is useful to check the integration of the equations of motion against discretization errors. In other words, the conservation of the effective energy can be used to choose an appropriate integration *timestep*. This is similar to the usual paradigm of checking the conservation of the total energy in the microcanonical ensemble.

---

#### 17.213.4 Restart, fix\_modify, output, run start/stop, minimize info

These fixes write the cumulative global energy change and the random number generator states to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the selected fix continues in an uninterrupted fashion. The random number generator state can only be restored when the number of processors remains unchanged from what is recorded in the restart file.

The *fix\_modify temp* option is supported by these fixes. You can use it to assign a temperature *compute* you have defined to these fixes which will be used in its thermostating procedure, as described above. For consistency, the group used by these fixes and by the compute should be the same.

The cumulative energy change in the system imposed by these fixes is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

These fixes compute a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

These fixes can ramp their target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

These fixes are not invoked during *energy minimization*.

### 17.213.5 Restrictions

Fix *temp/csld* is not compatible with *fix shake*.

These fixes are part of the EXTRA-FIX package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These fixes can be used with dynamic groups as defined by the *group* command. Likewise it can be used with groups to which atoms are added or deleted over time, e.g. a deposition simulation. However, the conservation properties of the thermostat and barostat are defined for systems with a static set of atoms. You may observe odd behavior if the atoms in a group vary dramatically over time or the atom count becomes very small.

### 17.213.6 Related commands

*fix nve*, *fix nvt*, *fix temp/rescale*, *fix langevin*, *fix\_modify*, *compute temp*, *fix temp/berendsen*

### 17.213.7 Default

none

---

**(Bussi1)** Bussi, Donadio and Parrinello, J. Chem. Phys. 126, 014101(2007)

**(Bussi2)** Bussi and Parrinello, Phys. Rev. E 75, 056707 (2007)

## 17.214 fix temp/rescale command

### 17.214.1 Syntax

```
fix ID group-ID temp/rescale N Tstart Tstop window fraction
```

- ID, group-ID are documented in *fix* command
- temp/rescale = style name of this fix command
- N = perform rescaling every N steps
- Tstart, Tstop = desired temperature at start/end of run (temperature units)

```
Tstart can be a variable (see below)
```

- window = only rescale if temperature is outside this window (temperature units)
- fraction = rescale to target temperature by this fraction

## 17.214.2 Examples

```
fix 3 flow temp/rescale 100 1.0 1.1 0.02 0.5
fix 3 boundary temp/rescale 1 1.0 1.5 0.05 1.0
fix 3 boundary temp/rescale 1 1.0 1.5 0.05 1.0
```

## 17.214.3 Description

Reset the temperature of a group of atoms by explicitly rescaling their velocities.

The rescaling is applied to only the translational degrees of freedom for the particles, which is an important consideration if finite-size particles which have rotational degrees of freedom are being thermostatted with this fix. The translational degrees of freedom can also have a bias velocity removed from them before thermostating takes place; see the description below.

Rescaling is performed every N timesteps. The target temperature is a ramped value between the *Tstart* and *Tstop* temperatures at the beginning and end of the run.

---

**Note:** This thermostat will generate an error if the current temperature is zero at the end of a timestep it is invoked on. It cannot rescale a zero temperature.

---

*Tstart* can be specified as an equal-style *variable*. In this case, the *Tstop* setting is ignored. If the value is a variable, it should be specified as *v\_name*, where name is the variable name. In this case, the variable will be evaluated each timestep, and its value used to determine the target temperature.

Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent temperature.

Rescaling is only performed if the difference between the current and desired temperatures is greater than the *window* value. The amount of rescaling that is applied is a *fraction* (from 0.0 to 1.0) of the difference between the actual and desired temperature. E.g. if *fraction* = 1.0, the temperature is reset to exactly the desired value.

---

**Note:** Unlike the *fix nvt* command which performs Nose/Hoover thermostating AND time integration, this fix does NOT perform time integration. It only modifies velocities to effect thermostating. Thus you must use a separate time integration fix, like *fix nve* to actually update the positions of atoms using the modified velocities. Likewise, this fix should not normally be used on atoms that also have their temperature controlled by another fix - e.g. by *fix nvt* or *fix langevin* commands.

---

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostating.

This fix computes a temperature each timestep. To do this, the fix creates its own compute of style “temp”, as if one of this command had been issued:

```
compute fix-ID_temp group-ID temp
```

See the *compute temp* for details. Note that the ID of the new compute is the fix-ID + underscore + “temp”, and the group for the new compute is the same as the fix group.

Note that this is NOT the compute used by thermodynamic output (see the *thermo\_style* command) with ID = *thermo\_temp*. This means you can change the attributes of this fix’s temperature (e.g. its degrees-of-freedom) via the *compute\_modify* command or print this temperature during thermodynamic output via the *thermo\_style custom*

command using the appropriate compute-ID. It also means that changing attributes of *thermo\_temp* will have no effect on this fix.

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

#### 17.214.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the cumulative global energy change to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the fix continues in an uninterrupted fashion.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a temperature *compute* you have defined to this fix which will be used in its thermostating procedure, as described above. For consistency, the group used by this fix and by the compute should be the same.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

#### 17.214.5 Restrictions

none

#### 17.214.6 Related commands

*fix langevin*, *fix nvt*, *fix\_modify*

#### 17.214.7 Default

none

## 17.215 fix temp/rescale/eff command

### 17.215.1 Syntax

```
fix ID group-ID temp/rescale/eff N Tstart Tstop window fraction
```

- ID, group-ID are documented in *fix* command
- temp/rescale/eff = style name of this fix command
- N = perform rescaling every N steps
- Tstart,Tstop = desired temperature at start/end of run (temperature units)
- window = only rescale if temperature is outside this window (temperature units)
- fraction = rescale to target temperature by this fraction

### 17.215.2 Examples

```
fix 3 flow temp/rescale/eff 10 1.0 100.0 0.02 1.0
```

### 17.215.3 Description

Reset the temperature of a group of nuclei and electrons in the *electron force field* model by explicitly rescaling their velocities.

The operation of this fix is exactly like that described by the *fix temp/rescale* command, except that the rescaling is also applied to the radial electron velocity for electron particles.

### 17.215.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify temp* option is supported by this fix. You can use it to assign a temperature *compute* you have defined to this fix which will be used in its thermostating procedure, as described above. For consistency, the group used by this fix and by the compute should be the same.

The cumulative energy change in the system imposed by this fix is included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* doc page for details.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

This fix can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

### 17.215.5 Restrictions

This fix is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.215.6 Related commands

*fix langevin/eff*, *fix nvt/eff*, *fix\_modify*, *fix temp rescale*,

### 17.215.7 Default

none

## 17.216 fix tfmc command

### 17.216.1 Syntax

```
fix ID group-ID tfmc Delta Temp seed keyword value
```

- ID, group-ID are documented in *fix* command
- tfmc = style name of this fix command
- Delta = maximal displacement length (distance units)
- Temp = imposed temperature of the system
- seed = random number seed (positive integer)
- zero or more keyword/arg pairs may be appended
- keyword = *com* or *rot*
  - com* args = xflag yflag zflag  
xflag,yflag,zflag = 0/1 to exclude/include each dimension
  - rot* args = none

### 17.216.2 Examples

```
fix 1 all tfmc 0.1 1000.0 159345  
fix 1 all tfmc 0.05 600.0 658943 com 1 1 0  
fix 1 all tfmc 0.1 750.0 387068 com 1 1 1 rot
```

### 17.216.3 Description

Perform uniform-acceptance force-bias Monte Carlo (fbMC) simulations, using the time-stamped force-bias Monte Carlo (tfMC) algorithm described in (*Mees*) and (*Bal*).

One successful use case of force-bias Monte Carlo methods is that they can be used to extend the time scale of atomistic simulations, in particular when long time scale relaxation effects must be considered; some interesting examples are given in the review by (*Neyts*). An example of a typical use case would be the modelling of chemical vapor deposition (CVD) processes on a surface, in which impacts by gas-phase species can be performed using MD, but subsequent relaxation of the surface is too slow to be done using MD only. Using tfMC can allow for a much faster relaxation of the surface, so that higher fluxes can be used, effectively extending the time scale of the simulation. (Such an alternating simulation approach could be set up using a *loop*.)

The initial version of tfMC algorithm in (*Mees*) contained an estimation of the effective time scale of such a simulation, but it was later shown that the speed-up one can gain from a tfMC simulation is system- and process-dependent, ranging from none to several orders of magnitude. In general, solid-state processes such as (re)crystallization or growth can be accelerated by up to two or three orders of magnitude, whereas diffusion in the liquid phase is not accelerated at all. The observed pseudodynamics when using the tfMC method is not the actual dynamics one would obtain using MD, but the relative importance of processes can match the actual relative dynamics of the system quite well, provided *Delta* is chosen with care. Thus, the system's equilibrium is reached faster than in MD, along a path that is generally roughly similar to a typical MD simulation (but not necessarily so). See (*Bal*) for details.

Each step, all atoms in the selected group are displaced using the stochastic tfMC algorithm, which is designed to sample the canonical (NVT) ensemble at the temperature *Temp*. Although tfMC is a Monte Carlo algorithm and thus strictly speaking does not perform time integration, it is similar in the sense that it uses the forces on all atoms in order to update their positions. Therefore, it is implemented as a time integration fix, and no other fixes of this type (such as *fix nve*) should be used at the same time. Because velocities do not play a role in this kind of Monte Carlo simulations, instantaneous temperatures as calculated by *temperature computes* or *thermodynamic output* have no meaning: the only relevant temperature is the sampling temperature *Temp*. Similarly, performing tfMC simulations does not require setting a *timestep* and the *simulated time* as calculated by LAMMPS is meaningless.

The critical parameter determining the success of a tfMC simulation is *Delta*, the maximal displacement length of the lightest element in the system: the larger it is, the longer the effective time scale of the simulation will be (there is an approximately quadratic dependence). However, *Delta* must also be chosen sufficiently small in order to comply with detailed balance; in general values between 5 and 10 % of the nearest neighbor distance are found to be a good choice. For a more extensive discussion with specific examples, please refer to (*Bal*), which also describes how the code calculates element-specific maximal displacements from *Delta*, based on the fourth root of their mass.

Because of the uncorrelated movements of the atoms, the center-of-mass of the fix group will not necessarily be stationary, just like its orientation. When the *com* keyword is used, all atom positions will be shifted (after every tfMC iteration) in order to fix the position of the center-of-mass along the included directions, by setting the corresponding flag to 1. The *rot* keyword does the same for the rotational component of the tfMC displacements after every iteration.

---

**Note:** the *com* and *rot* keywords should not be used if an external force is acting on the specified fix group, along the included directions. This can be either a true external force (e.g. through *fix wall*) or forces due to the interaction with atoms not included in the fix group. This is because in such cases, translations or rotations of the fix group could be induced by these external forces, and removing them will lead to a violation of detailed balance.

---

## 17.216.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

None of the *fix\_modify* options are relevant to this fix.

This fix is not invoked during *energy minimization*.

## 17.216.5 Restrictions

This fix is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

This fix is not compatible with *fix shake*.

## 17.216.6 Related commands

*fix gcmc*, *fix nvt*

## 17.216.7 Default

The option default is `com = 0 0 0`

---

**(Bal)** K. M Bal and E. C. Neyts, J. Chem. Phys. 141, 204104 (2014).

**(Mees)** M. J. Mees, G. Pourtois, E. C. Neyts, B. J. Thijsse, and A. Stesmans, Phys. Rev. B 85, 134301 (2012).

**(Neyts)** E. C. Neyts and A. Bogaerts, Theor. Chem. Acc. 132, 1320 (2013).

## 17.217 fix tgnvt/drude command

## 17.218 fix tgnpt/drude command

### 17.218.1 Syntax

```
fix ID group-ID style_name keyword values ...
```

- ID, group-ID are documented in *fix* command
- style\_name = *tgnvt/drude* or *tgnpt/drude*
- one or more keyword/values pairs may be appended

keyword = *temp iso* or *aniso* or *tri* or *x* or *y* or *z* or *xy* or *yz* or *xz* or *couple* or   
↪ *tchain* or *pchain* or *mtk* or *tloop* or *ploop* or *nreset* or *scalexy* or *scaleyz* or   
↪ *scalexz* or *flip* or *fixedpoint*

*temp* values = Tstart Tstop Tdamp Tdrude Tdamp\_drude

Tstart, Tstop = external temperature at start/end of run (temperature units)

Tdamp = temperature damping parameter (time units)

Tdrude = desired temperature of Drude oscillators (temperature units)

Tdamp\_drude = temperature damping parameter for Drude oscillators (time units)



*iso* or *aniso* or *tri* values = Pstart Pstop Pdamp  
 Pstart,Pstop = scalar external pressure at start/end of run (pressure units)  
 Pdamp = pressure damping parameter (time units)  
*x* or *y* or *z* or *xy* or *yz* or *xz* values = Pstart Pstop Pdamp  
 Pstart,Pstop = external stress tensor component at start/end of run (pressure\_↵  
 ↵units)  
 Pdamp = stress damping parameter (time units)  
*couple* = *none* or *xyz* or *xy* or *yz* or *xz*  
*tchain* value = N  
 N = length of thermostat chain (1 = single thermostat)  
*pchain* value = N  
 N length of thermostat chain on barostat (0 = no thermostat)  
*mtk* value = *yes* or *no* = add in MTK adjustment term or not  
*tloop* value = M  
 M = number of sub-cycles to perform on thermostat  
*ploop* value = M  
 M = number of sub-cycles to perform on barostat thermostat  
*nreset* value = reset reference cell every this many timesteps  
*scalexy* value = *yes* or *no* = scale xy with ly  
*scaleyz* value = *yes* or *no* = scale yz with lz  
*scalexz* value = *yes* or *no* = scale xz with lz  
*flip* value = *yes* or *no* = allow or disallow box flips when it becomes highly skewed  
*fixedpoint* values = x y z  
 x,y,z = perform barostat dilation/contraction around this point (distance units)

## 17.218.2 Examples

```

comm_modify vel yes
fix 1 all tgnvt/drude temp 300.0 300.0 100.0 1.0 20.0
fix 1 water tgnpt/drude temp 300.0 300.0 100.0 1.0 20.0 iso 0.0 0.0 1000.0
fix 2 jello tgnpt/drude temp 300.0 300.0 100.0 1.0 20.0 tri 5.0 5.0 1000.0
fix 2 ice tgnpt/drude temp 250.0 250.0 100.0 1.0 20.0 x 1.0 1.0 0.5 y 2.0 2.0 0.5 z 3.0_↵
↵3.0 0.5 yz 0.1 0.1 0.5 xz 0.2 0.2 0.5 xy 0.3 0.3 0.5 nreset 1000
  
```

Example input scripts available: examples/PACKAGES/drude

## 17.218.3 Description

These commands are variants of the Nose-Hoover fix styles *fix nvt* and *fix npt* for thermalized Drude polarizable models. They apply temperature-grouped Nose-Hoover thermostat (TGNH) proposed by (Son). When there are fast vibrational modes with frequencies close to Drude oscillators (e.g. double bonds or out-of-plane torsions), this thermostat can provide better kinetic energy equipartitioning.

The difference between TGNH and the original Nose-Hoover thermostat is that, TGNH separates the kinetic energy of the group into three contributions: molecular center of mass (COM) motion, motion of COM of atom-Drude pairs or non-polarizable atoms relative to molecular COM, and relative motion of atom-Drude pairs. An independent Nose-Hoover chain is applied to each type of motion. The temperatures for these three types of motion are denoted as molecular translational temperature ( $T_M$ ), real atomic temperature ( $T_R$ ) and Drude temperature ( $T_D$ ), which are defined in terms of their associated degrees of freedom (DOF):

$$T_M = \frac{\sum_i^{N_{\text{mol}}} M_i V_i^2}{3 \left( N_{\text{mol}} - \frac{N_{\text{mol}}}{N_{\text{mol,sys}}} \right) k_B}$$

$$T_R = \frac{\sum_i^{N_{\text{real}}} m_i (v_i - v_{M,i})^2}{(N_{\text{DOF}} - 3N_{\text{mol}} + 3\frac{N_{\text{mol}}}{N_{\text{mol,sys}}} - 3N_{\text{drude}})k_B}$$

$$T_D = \frac{\sum_i^{N_{\text{drude}}} m'_i v'_i{}^2}{3N_{\text{drude}}k_B}$$

Here  $N_{\text{mol}}$  and  $N_{\text{mol,sys}}$  are the numbers of molecules in the group and in the whole system, respectively.  $N_{\text{real}}$  is the number of atom-Drude pairs and non-polarizable atoms in the group.  $N_{\text{drude}}$  is the number of Drude particles in the group.  $N_{\text{DOF}}$  is the DOF of the group.  $M_i$  and  $V_i$  are the mass and the COM velocity of the  $i$ -th molecule.  $m_i$  is the mass of the  $i$ -th atom-Drude pair or non-polarizable atom.  $v_i$  is the velocity of COM of  $i$ -th atom-Drude pair or non-polarizable atom.  $v_{M,i}$  is the COM velocity of the molecule the  $i$ -th atom-Drude pair or non-polarizable atom belongs to.  $m'_i$  and  $v'_i$  are the reduced mass and the relative velocity of the  $i$ -th atom-Drude pair.

---

**Note:** These fixes require that each atom knows whether it is a Drude particle or not. You must therefore use the *fix drude* command to specify the Drude status of each atom type.

Because the TGNH thermostat thermostats the molecular COM motion, all atoms belonging to the same molecule must be in the same group. That is, these fixes can not be applied to a subset of a molecule.

For this fix to act correctly, ghost atoms need to know their velocity. You must use the *comm\_modify* command to enable this.

These fixes assume that the translational DOF of the whole system is removed. It is therefore recommended to invoke *fix momentum* command so that the  $T_M$  is calculated correctly.

---

The thermostat parameters are specified using the *temp* keyword. The thermostat is applied to only the translational DOF for the particles. The translational DOF can also have a bias velocity removed before thermostating takes place; see the description below. The desired temperature for molecular and real atomic motion is a ramped value during the run from  $T_{\text{start}}$  to  $T_{\text{stop}}$ . The *Tdamp* parameter is specified in time units and determines how rapidly the temperature is relaxed. For example, a value of 10.0 means to relax the temperature in a timespan of (roughly) 10 time units (e.g.  $\tau$  or fs or ps - see the *units* command). The parameter *Tdrude* is the desired temperature for Drude motion at each timestep. Similar to *Tdamp*, the *Tdamp\_drude* parameter determines the relaxation speed for Drude motion. Fix group are the only ones whose velocities and positions are updated by the velocity/position update portion of the integration. Other thermostat-related keywords are *tchain* and *tloop*, which are detailed in *fix nvt*.

---

**Note:** A Nose-Hoover thermostat will not work well for arbitrary values of *Tdamp*. If *Tdamp* is too small, the temperature can fluctuate wildly; if it is too large, the temperature will take a very long time to equilibrate. A good choice for many models is a *Tdamp* of around 100 timesteps. A smaller *Tdamp\_drude* value would be required to maintain Drude motion at low temperature.

---

```
fix 1 all nvt temp 300.0 300.0 $(100.0*dt) 1.0 $(20.0*dt)
```

The barostat parameters for fix style *ignpt/drude* is specified using one or more of the *iso*, *aniso*, *tri*, *x*, *y*, *z*, *xy*, *xz*, *yz*, and *couple* keywords. These keywords give you the ability to specify all 6 components of an external stress tensor, and to couple various of these components together so that the dimensions they represent are varied together during a constant-pressure simulation. Other barostat-related keywords are *pchain*, *mtk*, *ploop*, *nreset*, *scalexy*, *scaleyz*, *scalexz*, *flipand* *fixedpoint*. The meaning of barostat parameters are detailed in *fix npt*.

Regardless of what atoms are in the fix group (the only atoms which are time integrated), a global pressure or stress tensor is computed for all atoms. Similarly, when the size of the simulation box is changed, all atoms are re-scaled to new positions.

**Note:** Unlike the *fix temp/berendsen* command which performs thermostating but NO time integration, these fixes perform thermostating/barostatting AND time integration. Thus you should not use any other time integration fix, such as *fix nve* on atoms to which this fix is applied. Likewise, these fixes should not be used on atoms that also have their temperature controlled by another fix - e.g. by *fix langevin/drude* command.

---

See the *Howto thermostat* and *Howto barostat* doc pages for a discussion of different ways to compute temperature and perform thermostating and barostatting.

---

Like other fixes that perform thermostating, this fix can be used with *compute commands* that remove a “bias” from the atom velocities. E.g. to apply the thermostat only to atoms within a spatial *region*, or to remove the center-of-mass velocity from a group of atoms, or to remove the x-component of velocity from the calculation.

This is not done by default, but only if the *fix\_modify* command is used to assign a temperature compute to this fix that includes such a bias term. See the doc pages for individual *compute temp commands* to determine which ones include a bias. In this case, the thermostat works in the following manner: bias is removed from each atom, thermostating is performed on the remaining thermal degrees of freedom, and the bias is added back in.

---

**Note:** However, not all temperature compute commands are valid to be used with these fixes. Precisely, only temperature compute that does not modify the DOF of the group can be used. E.g. *compute temp/ramp* and *compute viscosity/cos* compute the kinetic energy after remove a velocity gradient without affecting the DOF of the group, then they can be invoked in this way. In contrast, *compute temp/partial* may remove the DOF at one or more dimensions, therefore it cannot be used with these fixes.

---

#### 17.218.4 Restart, fix\_modify, output, run start/stop, minimize info

These fixes writes the state of all the thermostat and barostat variables to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify temp* and *press* options are supported by these fixes. You can use them to assign a *compute* you have defined to this fix which will be used in its thermostating or barostatting procedure, as described above. If you do this, note that the kinetic energy derived from the compute temperature should be consistent with the virial term computed using all atoms for the pressure. LAMMPS will warn you if you choose to compute temperature on a subset of atoms.

---

**Note:** If both the *temp* and *press* keywords are used in a single *thermo\_modify* command (or in two separate commands), then the order in which the keywords are specified is important. Note that a *pressure compute* defines its own temperature compute as an argument when it is specified. The *temp* keyword will override this (for the pressure compute being used by *fix npt*), but only if the *temp* keyword comes after the *press* keyword. If the *temp* keyword comes before the *press* keyword, then the new pressure compute specified by the *press* keyword will be unaffected by the *temp* setting.

---

The cumulative energy change in the system imposed by these fixes, due to thermostating and/or barostatting, are included in the *thermodynamic output* keywords *ecouple* and *econserve*. See the *thermo\_style* page for details.

These fixes compute a global scalar which can be accessed by various *output commands*. The scalar is the same cumulative energy change due to this fix described in the previous paragraph. The scalar value calculated by this fix is “extensive”.

---

These fixes also compute a global vector of quantities, which can be accessed by various *output commands*. The vector values are “intensive”. The vector stores the three temperatures  $T_M$ ,  $T_R$  and  $T_D$ .

These fixes can ramp their external temperature and pressure over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

These fixes are not invoked during *energy minimization*.

---

## 17.218.5 Restrictions

These fixes are only available when LAMMPS was built with the DRUDE package. These fixes cannot be used with dynamic groups as defined by the *group* command. These fixes cannot be used in 2D simulations.

$X$ ,  $y$ ,  $z$  cannot be barostatted if the associated dimension is not periodic.  $Xy$ ,  $xz$ , and  $yz$  can only be barostatted if the simulation domain is triclinic and the second dimension in the keyword ( $y$  dimension in  $xy$ ) is periodic. The *create\_box*, *read\_data*, and *read\_restart* commands specify whether the simulation box is orthogonal or non-orthogonal (triclinic) and explain the meaning of the  $xy,xz,yz$  tilt factors.

For the *temp* keyword, the final  $T_{stop}$  cannot be 0.0 since it would make the external  $T = 0.0$  at some timestep during the simulation which is not allowed in the Nose/Hoover formulation.

The *scaleyz yes*, *scalexz yes*, and *scalexy yes* options can only be used if the second dimension in the keyword is periodic, and if the tilt factor is not coupled to the barostat via keywords *tri*, *yz*, *xz*, and *xy*.

## 17.218.6 Related commands

*fix drude*, *fix nvt*, *fix\_npt*, *fix\_modify*

## 17.218.7 Default

The keyword defaults are  $tchain = 3$ ,  $pchain = 3$ ,  $mtk = yes$ ,  $tloop = 1$ ,  $ploop = 1$ ,  $nreset = 0$ ,  $couple = none$ ,  $flip = yes$ ,  $scaleyz = scalexz = scalexy = yes$  if periodic in second dimension and not coupled to barostat, otherwise no.

---

(Son) Son, McDaniel, Cui and Yethiraj, J Phys Chem Lett, 10, 7523 (2019).

## 17.219 fix thermal/conductivity command

### 17.219.1 Syntax

```
fix ID group-ID thermal/conductivity N edim Nbin keyword value ...
```

- ID, group-ID are documented in *fix* command
- thermal/conductivity = style name of this fix command
- N = perform kinetic energy exchange every N steps
- edim =  $x$  or  $y$  or  $z$  = direction of kinetic energy transfer
- Nbin = # of layers in edim direction (must be even number)

- zero or more keyword/value pairs may be appended
- keyword = *swap*  
*swap* value = *Nswap* = number of swaps to perform every *N* steps

## 17.219.2 Examples

```
fix 1 all thermal/conductivity 100 z 20
fix 1 all thermal/conductivity 50 z 20 swap 2
```

## 17.219.3 Description

Use the Muller-Plathe algorithm described in [this paper](#) to exchange kinetic energy between two particles in different regions of the simulation box every *N* steps. This induces a temperature gradient in the system. As described below this enables the thermal conductivity of a material to be calculated. This algorithm is sometimes called a reverse non-equilibrium MD (reverse NEMD) approach to computing thermal conductivity. This is because the usual NEMD approach is to impose a temperature gradient on the system and measure the response as the resulting heat flux. In the Muller-Plathe method, the heat flux is imposed, and the temperature gradient is the system's response.

See the [compute heat/flux](#) command for details on how to compute thermal conductivity in an alternate way, via the Green-Kubo formalism.

The simulation box is divided into *Nbin* layers in the *edim* direction, where the layer 1 is at the low end of that dimension and the layer *Nbin* is at the high end. Every *N* steps, *Nswap* pairs of atoms are chosen in the following manner. Only atoms in the *fix* group are considered. The hottest *Nswap* atoms in layer 1 are selected. Similarly, the coldest *Nswap* atoms in the “middle” layer (see below) are selected. The two sets of *Nswap* atoms are paired up and their velocities are exchanged. This effectively swaps their kinetic energies, assuming their masses are the same. If the masses are different, an exchange of velocities relative to center of mass motion of the 2 atoms is performed, to conserve kinetic energy. Over time, this induces a temperature gradient in the system which can be measured using commands such as the following, which writes the temperature profile (assuming *z* = *edim*) to the file *tmp.profile*:

```
compute ke all ke/atom
variable temp atom c_ke/1.5
compute layers all chunk/atom bin/1d z lower 0.05 units reduced
fix 3 all ave/chunk 10 100 1000 layers v_temp file tmp.profile
```

Note that by default, *Nswap* = 1, though this can be changed by the optional *swap* keyword. Setting this parameter appropriately, in conjunction with the swap rate *N*, allows the heat flux to be adjusted across a wide range of values, and the kinetic energy to be exchanged in large chunks or more smoothly.

The “middle” layer for velocity swapping is defined as the  $Nbin/2 + 1$  layer. Thus if *Nbin* = 20, the two swapping layers are 1 and 11. This should lead to a symmetric temperature profile since the two layers are separated by the same distance in both directions in a periodic sense. This is why *Nbin* is restricted to being an even number.

As described below, the total kinetic energy transferred by these swaps is computed by the *fix* and can be output. Dividing this quantity by time and the cross-sectional area of the simulation box yields a heat flux. The ratio of heat flux to the slope of the temperature profile is proportional to the thermal conductivity of the fluid, in appropriate units. See the [Muller-Plathe paper](#) for details.

---

**Note:** If your system is periodic in the direction of the heat flux, then the flux is going in 2 directions. This means the effective heat flux in one direction is reduced by a factor of 2. You will see this in the equations for thermal conductivity

(kappa) in the Muller-Plathe paper. LAMMPS is simply tallying kinetic energy which does not account for whether or not your system is periodic; you must use the value appropriately to yield a kappa for your system.

---

**Note:** After equilibration, if the temperature gradient you observe is not linear, then you are likely swapping energy too frequently and are not in a regime of linear response. In this case you cannot accurately infer a thermal conductivity and should try increasing the `Nevery` parameter.

---

#### 17.219.4 Restart, `fix_modify`, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the cumulative kinetic energy transferred between the bottom and middle of the simulation box (in the *edim* direction) is stored as a scalar quantity by this fix. This quantity is zeroed when the fix is defined and accumulates thereafter, once every `N` steps. The units of the quantity are energy; see the *units* command for details. The scalar value calculated by this fix is “intensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.219.5 Restrictions

Swaps conserve both momentum and kinetic energy, even if the masses of the swapped atoms are not equal. Thus you should not need to thermostat the system. If you do use a thermostat, you may want to apply it only to the non-swapped dimensions (other than *vdim*).

LAMMPS does not check, but you should not use this fix to swap the kinetic energy of atoms that are in constrained molecules, e.g. via *fix shake* or *fix rigid*. This is because application of the constraints will alter the amount of transferred momentum. You should, however, be able to use flexible molecules. See the *Zhang paper* for a discussion and results of this idea.

When running a simulation with large, massive particles or molecules in a background solvent, you may want to only exchange kinetic energy between solvent particles.

#### 17.219.6 Related commands

*fix ehex*, *fix heat*, *fix ave/chunk*, *fix viscosity*, *compute heat/flux*

#### 17.219.7 Default

The option defaults are `swap = 1`.

---

**(Muller-Plathe)** Muller-Plathe, J Chem Phys, 106, 6082 (1997).

**(Zhang)** Zhang, Lussetti, de Souza, Muller-Plathe, J Phys Chem B, 109, 15060-15067 (2005).

## 17.220 fix ti/spring command

### 17.220.1 Syntax

```
fix ID group-ID ti/spring k t_s t_eq keyword value ...
```

- ID, group-ID are documented in *fix* command
- ti/spring = style name of this fix command
- k = spring constant (force/distance units)
- t\_eq = number of steps for the equilibration procedure
- t\_s = number of steps for the switching procedure
- zero or more keyword/value pairs may be appended to args

- keyword = *function*

*function* value = function-ID

function-ID = ID of the switching function (1 or 2)

### 17.220.2 Example

```
fix 1 all ti/spring 50.0 2000 1000 function 2
```

### 17.220.3 Description

This fix allows you to compute the free energy of crystalline solids by performing a nonequilibrium thermodynamic integration between the solid of interest and an Einstein crystal. A detailed explanation of how to use this command and choose its parameters for optimal performance and accuracy is given in the paper by *Freitas*. The paper also presents a short summary of the theory of nonequilibrium thermodynamic integration.

The thermodynamic integration procedure is performed by rescaling the force on each atom. Given an atomic configuration the force ( $F$ ) on each atom is given by

$$F = (1 - \lambda) F_{\text{solid}} + \lambda F_{\text{harm}}$$

where  $F_{\text{solid}}$  is the force that acts on an atom due to an interatomic potential (*e.g.* EAM potential),  $F_{\text{harm}}$  is the force due to the Einstein crystal harmonic spring, and  $\lambda$  is the coupling parameter of the thermodynamic integration. An Einstein crystal is a solid where each atom is attached to its equilibrium position by a harmonic spring with spring constant  $k$ . With this fix a spring force is applied independently to each atom in the group defined by the fix to tether it to its initial position. The initial position of each atom is its position at the time the fix command was issued.

The fix acts as follows: during the first  $t_{\text{eq}}$  steps after the fix is defined the value of  $\lambda$  is zero. This is the period to equilibrate the system in the  $\lambda = 0$  state. After this the value of  $\lambda$  changes dynamically during the simulation from 0 to 1 according to the function defined using the keyword *function* (described below), this switching from  $\lambda$  from 0 to 1 is done in  $t_{\text{s}}$  steps. Then comes the second equilibration period of  $t_{\text{eq}}$  to equilibrate the system in the  $\lambda = 1$  state. After that, the switching back to the  $\lambda = 0$  state is made using  $t_{\text{s}}$  timesteps and following the same switching function. After this period the value of  $\lambda$  is kept equal to zero and the fix has no other effect on the dynamics of the system.

The processes described above is known as nonequilibrium thermodynamic integration and is has been shown (*Freitas*) to present a much superior efficiency when compared to standard equilibrium methods. The reason why the switching

it is made in both directions (potential to Einstein crystal and back) is to eliminate the dissipated heat due to the nonequilibrium process. Further details about nonequilibrium thermodynamic integration and its implementation in LAMMPS is available in *Freitas*.

The *function* keyword allows the use of two different lambda paths. Option 1 results in a constant rate of change of lambda with time:

$$\lambda(\tau) = \tau$$

where  $\tau$  is the scaled time variable  $t/t_s$ . The option 2 performs the lambda switching at a rate defined by the following switching function

$$\lambda(\tau) = \tau^5 (70\tau^4 - 315\tau^3 + 540\tau^2 - 420\tau + 126)$$

This function has zero slope as lambda approaches its extreme values (0 and 1), according to *de Koning* this results in smaller fluctuations on the integral to be computed on the thermodynamic integration. The use of option 2 is recommended since it results in better accuracy and less dissipation without any increase in computational resources cost.

---

**Note:** As described in *Freitas*, it is important to keep the center-of-mass fixed during the thermodynamic integration. A nonzero total velocity will result in divergences during the integration due to the fact that the atoms are ‘attached’ to their equilibrium positions by the Einstein crystal. Check the option *zero* of *fix langevin* and *velocity*. The use of the Nose-Hoover thermostat (*fix nvt*) is *NOT* recommended due to its well documented issues with the canonical sampling of harmonic degrees of freedom (notice that the *chain* option will *NOT* solve this problem). The Langevin thermostat (*fix langevin*) correctly thermostats the system and we advise its usage with *ti/spring* command.

---

#### 17.220.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the original coordinates of tethered atoms to *binary restart files*, so that the spring effect will be the same in a restarted simulation. See the *read restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

The *fix\_modify energy* option is supported by this fix to add the energy stored in the per-atom springs to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

This fix computes a global scalar and a global vector quantities which can be accessed by various *output commands*. The scalar is an energy which is the sum of the spring energy for each atom, where the per-atom energy is  $0.5 \cdot k \cdot r^2$ . The vector stores 2 values. The first value is the coupling parameter lambda. The second value is the derivative of lambda with respect to the integer timestep  $s$ , i.e.  $\frac{d\lambda}{ds}$ . In order to obtain  $\frac{d\lambda}{dt}$ , where  $t$  is simulation time, this 2nd value needs to be divided by the timestep size (e.g. 0.5 fs). The scalar and vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the per-atom spring energy to be included in the total potential energy of the system (the quantity being minimized), you **MUST** enable the *fix modify energy* option for this fix.

---



## 17.220.5 Related commands

*fix spring*, *fix adapt*

## 17.220.6 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 17.220.7 Default

The keyword default is function = 1.

---

**(Freitas)** Freitas, Asta, and de Koning, Computational Materials Science, 112, 333 (2016).

**(de Koning)** de Koning and Antonelli, Phys Rev E, 53, 465 (1996).

## 17.221 fix tmd command

### 17.221.1 Syntax

```
fix ID group-ID tmd rho_final file1 N file2
```

- ID, group-ID are documented in *fix* command
- tmd = style name of this fix command
- rho\_final = desired value of rho at the end of the run (distance units)
- file1 = filename to read target structure from
- N = dump TMD statistics every this many timesteps, 0 = no dump
- file2 = filename to write TMD statistics to (only needed if N > 0)

### 17.221.2 Examples

```
fix 1 all nve
fix 2 tmdatoms tmd 1.0 target_file 100 tmd_dump_file
```

### 17.221.3 Description

Perform targeted molecular dynamics (TMD) on a group of atoms. A holonomic constraint is used to force the atoms to move towards (or away from) the target configuration. The parameter “rho” is monotonically decreased (or increased) from its initial value to rho\_final at the end of the run.

Rho has distance units and is a measure of the root-mean-squared distance (RMSD) between the current configuration of the atoms in the group and the target coordinates listed in file1. Thus a value of rho\_final = 0.0 means move the atoms all the way to the final structure during the course of the run.

The target file1 can be ASCII text or a gzipped text file (detected by a .gz suffix). The format of the target file1 is as follows:

```
0.0 25.0 xlo xhi
0.0 25.0 ylo yhi
0.0 25.0 zlo zhi
125      24.97311    1.69005      23.46956 0 0 -1
126      1.94691    2.79640      1.92799 1 0 0
127      0.15906    3.46099      0.79121 1 0 0
...
```

The first 3 lines may or may not be needed, depending on the format of the atoms to follow. If image flags are included with the atoms, the first 3 lo/hi lines **must** appear in the file. If image flags are not included, the first 3 lines **must not** appear. The 3 lines contain the simulation box dimensions for the atom coordinates, in the same format as in a LAMMPS data file (see the *read\_data* command).

The remaining lines each contain an atom ID and its target x,y,z coordinates. The atom lines (all or none of them) can optionally be followed by 3 integer values: nx,ny,nz. For periodic dimensions, they specify which image of the box the atom is considered to be in, i.e. a value of N (positive or negative) means add N times the box length to the coordinate to get the true value. Those 3 integers either must be given for all atoms or none.

The atom lines can be listed in any order, but every atom in the group must be listed in the file. Atoms not in the fix group may also be listed; they will be ignored.

Comments starting with '#' and empty lines may be included as well.

TMD statistics are written to file2 every N timesteps, unless N is specified as 0, which means no statistics.

The atoms in the fix tmd group should be integrated (via a fix nve, nvt, npt) along with other atoms in the system.

Restarts can be used with a fix tmd command. For example, imagine a 10000 timestep run with a rho\_initial = 11 and a rho\_final = 1. If a restart file was written after 2000 time steps, then the configuration in the file would have a rho value of 9. A new 8000 time step run could be performed with the same rho\_final = 1 to complete the conformational change at the same transition rate. Note that for restarted runs, the name of the TMD statistics file should be changed to prevent it being overwritten.

For more information about TMD, see (*Schlitter1*) and (*Schlitter2*).

#### 17.221.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*.

This fix can ramp its rho parameter over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

This fix is not invoked during *energy minimization*.

## 17.221.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

All TMD fixes must be listed in the input script after all integrator fixes (nve, nvt, npt) are applied. This ensures that atoms are moved before their positions are corrected to comply with the constraint.

Atoms that have a TMD fix applied should not be part of a group to which a SHAKE fix is applied. This is because LAMMPS assumes there are not multiple competing holonomic constraints applied to the same atoms.

To read gzipped target files, you must compile LAMMPS with the `-DLAMMPS_GZIP` option. See the [Build settings](#) doc page for details.

## 17.221.6 Related commands

none

## 17.221.7 Default

none

---

**(Schlitter1)** Schlitter, Swegat, Mulders, “Distance-type reaction coordinates for modelling activated processes”, *J Molecular Modeling*, 7, 171-177 (2001).

**(Schlitter2)** Schlitter and Klahn, “The free energy of a reaction coordinate at multiple constraints: a concise formulation”, *Molecular Physics*, 101, 3439-3443 (2003).

## 17.222 fix ttm command

### 17.223 fix ttm/grid command

### 17.224 fix ttm/mod command

#### 17.224.1 Syntax

```
fix ID group-ID ttm seed C_e rho_e kappa_e gamma_p gamma_s v_0 Nx Ny Nz keyword value ...
fix ID group-ID ttm/mod seed init_file Nx Ny Nz keyword value ...
```

- ID, group-ID are documented in [fix](#) command
- style = `ttm` or `ttm/grid` or `ttm/mod`
- seed = random number seed to use for white noise (positive integer)
- remaining arguments for `fix ttm` or `fix ttm/grid`

`C_e` = electronic specific heat (energy/(electron\*temperature) units)

`rho_e` = electronic density (electrons/volume units)

`kappa_e` = electronic thermal conductivity (energy/(time\*distance\*temperature) units)

`gamma_p` = friction coefficient due to electron-ion interactions (mass/time units)

```

gamma_s = friction coefficient due to electronic stopping (mass/time units)
v_0 = electronic stopping critical velocity (velocity units)
Nx = number of thermal solve grid points in the x-direction (positive integer)
Ny = number of thermal solve grid points in the y-direction (positive integer)
Nz = number of thermal solve grid points in the z-direction (positive integer)

```

- remaining arguments for fix ttm/mod:

```

init_file = file with the parameters to TTM
Nx = number of thermal solve grid points in the x-direction (positive integer)
Ny = number of thermal solve grid points in the y-direction (positive integer)
Nz = number of thermal solve grid points in the z-direction (positive integer)

```

- zero or more keyword/value(s) pairs may be appended
- keyword = *set* or *infile* or *outfile*

```

set value = Tinit
  Tinit = initial electronic temperature at all grid points (temperature units)
infile value = file.in with grid values for electronic temperatures
outfile values = Nout file.out
  Nout = dump grid temperatures every this many timesteps
  file.out = filename to write grid temperatures to

```

## 17.224.2 Examples

```

fix 2 all ttm 699489 1.0 1.0 10 0.1 0.0 2.0 1 12 1 infile initial outfile 1000 T.out
fix 3 all ttm/grid 123456 1.0 1.0 1.0 1.0 1.0 5.0 5 5 5 infile Te.in
fix 4 all ttm/mod 34277 parameters.txt 5 5 5 infile T_init outfile 10 T_out

```

Example input scripts using these commands can be found in examples/ttm.

## 17.224.3 Description

Use a two-temperature model (TTM) to represent heat transfer through and between electronic and atomic subsystems. LAMMPS models the atomic subsystem as usual with a molecular dynamics model and the classical force field specified by the user. The electronic subsystem is modeled as a continuum, or a background “gas”, on a regular grid which overlays the simulation domain. Energy can be transferred spatially within the grid representing the electrons. Energy can also be transferred between the electronic and atomic subsystems. The algorithm underlying this fix was derived by D. M. Duffy and A. M. Rutherford and is discussed in two J Physics: Condensed Matter papers: ([Duffy](#)) and ([Rutherford](#)). They used this algorithm in cascade simulations where a primary knock-on atom (PKA) was initialized with a high velocity to simulate a radiation event.

The description in this sub-section applies to all 3 fix styles: *ttm*, *ttm/grid*, and *ttm/mod*.

Fix *ttm/grid* distributes the regular grid across processors consistent with the sub-domains of atoms owned by each processor, but is otherwise identical to fix *ttm*. Note that fix *ttm* stores a copy of the grid on each processor, which is acceptable when the overall grid is reasonably small. For larger grids you should use fix *ttm/grid* instead.

Fix *ttm/mod* adds options to account for external heat sources (e.g. at a surface) and for specifying parameters that allow the electronic heat capacity to depend strongly on electronic temperature. It is more expensive computationally than fix *ttm* because it treats the thermal diffusion equation as non-linear. More details on fix *ttm/mod* are given below.

Heat transfer between the electronic and atomic subsystems is carried out via an inhomogeneous Langevin thermostat. Only atoms in the fix group contribute to and are affected by this heat transfer.

This thermostating differs from the regular Langevin thermostat (*fix langevin*) in three important ways. First, the Langevin thermostat is applied uniformly to all atoms in the user-specified group for a single target temperature, whereas the TTM fixes apply Langevin thermostating locally to atoms within the volumes represented by the user-specified grid points with a target temperature specific to that grid point. Second, the Langevin thermostat couples the temperature of the atoms to an infinite heat reservoir, whereas the heat reservoir for the TTM fixes is finite and represents the local electrons. Third, the TTM fixes allow users to specify not just one friction coefficient, but rather two independent friction coefficients: one for the electron-ion interactions (*gamma\_p*), and one for electron stopping (*gamma\_s*).

When the friction coefficient due to electron stopping, *gamma\_s*, is non-zero, electron stopping effects are included for atoms moving faster than the electron stopping critical velocity, *v\_0*. For further details about this algorithm, see (*Duffy*) and (*Rutherford*).

Energy transport within the electronic subsystem is solved according to the heat diffusion equation with added source terms for heat transfer between the subsystems:

$$C_e \rho_e \frac{\partial T_e}{\partial t} = \nabla \cdot (\kappa_e \nabla T_e) - g_p (T_e - T_a) + g_s T'_a$$

where *C\_e* is the specific heat, *rho\_e* is the density, *kappa\_e* is the thermal conductivity, *T* is temperature, the “e” and “a” subscripts represent electronic and atomic subsystems respectively, *g\_p* is the coupling constant for the electron-ion interaction, and *g\_s* is the electron stopping coupling parameter. *C\_e*, *rho\_e*, and *kappa\_e* are specified as parameters to the fix. The other quantities are derived. The form of the heat diffusion equation used here is almost the same as that in equation 6 of (*Duffy*), with the exception that the electronic density is explicitly represented, rather than being part of the specific heat parameter.

Currently, the TTM fixes assume that none of the user-supplied parameters will vary with temperature. Note that (*Duffy*) used a tanh() functional form for the temperature dependence of the electronic specific heat, but ignored temperature dependencies of any of the other parameters. See more discussion below for fix ttm/mod.

---

**Note:** These fixes do not perform time integration of the atoms in the fix group, they only rescale their velocities. Thus a time integration fix such as *fix nve* should be used in conjunction with these fixes. These fixes should not normally be used on atoms that have their temperature controlled by another thermostating fix, e.g. *fix nvt* or *fix langevin*.

---



---

**Note:** These fixes require use of an orthogonal 3d simulation box with periodic boundary conditions in all dimensions. They also require that the size and shape of the simulation box do not vary dynamically, e.g. due to use of the *fix npt* command. Likewise, the size/shape of processor sub-domains cannot vary due to dynamic load-balancing via use of the *fix balance* command. It is possible however to load balance before the simulation starts using the *balance* command, so that each processor has a different size sub-domain.

---

Periodic boundary conditions are also used in the heat equation solve for the electronic subsystem. This varies from the approach of (*Rutherford*) where the atomic subsystem was embedded within a larger continuum representation of the electronic subsystem.

The *set* keyword specifies a *Tinit* temperature value to initialize the value stored on all grid points.

The *infile* keyword specifies an input file of electronic temperatures for each grid point to be read in to initialize the grid. By default the temperatures are all zero when the grid is created. The input file is a text file which may have comments starting with the ‘#’ character. Each line contains four numeric columns: ix,iy,iz,Temperature. Empty or comment-only lines will be ignored. The number of lines must be equal to the number of user-specified grid points (Nx by Ny by Nz). The ix,iy,iz are grid point indices ranging from 0 to nxnodes-1 inclusive in each dimension. The lines can appear in any order. For example, the initial electronic temperatures on a 1 by 2 by 3 grid could be specified in the file as follows:

```
# UNITS: metal COMMENT: initial electron temperature
0 0 0 1.0
0 0 1 1.0
0 0 2 1.0
0 1 0 2.0
0 1 1 2.0
0 1 2 2.0
```

where the electronic temperatures along the  $y=0$  plane have been set to 1.0, and the electronic temperatures along the  $y=1$  plane have been set to 2.0. If all the grid point values are not specified, LAMMPS will generate an error. LAMMPS will check if a “UNITS:” tag is in the first line and stop with an error, if there is a mismatch with the current units used.

**Note:** The electronic temperature at each grid point must be a non-zero positive value, both initially, and as the temperature evolves over time. Thus you must use either the *set* or *infile* keyword or be restarting a simulation that used this fix previously.

The *outfile* keyword has 2 values. The first value *Nout* triggers output of the electronic temperatures for each grid point every *Nout* timesteps. The second value is the filename for output which will be suffixed by the timestep. The format of each output file is exactly the same as the input temperature file. It will contain a comment in the first line reporting the date the file was created, the LAMMPS units setting in use, grid size and the current timestep.

Note that the atomic temperature for atoms in each grid cell can also be computed and output by the *fix ave/chunk* command using the *compute chunk/atom* command to create a 3d array of chunks consistent with the grid used by this fix.

### Additional details for fix ttm/mod

Fix *ttm/mod* uses the heat diffusion equation with possible external heat sources (e.g. laser heating in ablation simulations):

$$C_e \rho_e \frac{\partial T_e}{\partial t} = \nabla \cdot (\kappa_e \nabla T_e) - g_p (T_e - T_a) + g_s T'_a + \theta(x - x_{surface}) I_0 \exp(-x/l_{skin})$$

where theta is the Heaviside step function,  $I_0$  is the (absorbed) laser pulse intensity for ablation simulations,  $l_{skin}$  is the depth of skin-layer, and all other designations have the same meaning as in the former equation. The duration of the pulse is set by the parameter *tau* in the *init\_file*.

Fix *ttm/mod* also allows users to specify the dependencies of  $C_e$  and  $\kappa_e$  on the electronic temperature. The specific heat is expressed as

$$C_e = C_0 + (a_0 + a_1 X + a_2 X^2 + a_3 X^3 + a_4 X^4) \exp(-(AX)^2)$$

where  $X = T_e/1000$ , and the thermal conductivity is defined as  $\kappa_e = D_e \rho_e C_e$ , where  $D_e$  is the thermal diffusion coefficient.

Electronic pressure effects are included in the TTM model to account for the blast force acting on ions because of electronic pressure gradient (see (*Chen*), (*Norman*)). The total force acting on an ion is:

$$\vec{F}_i = -\partial U / \partial \vec{r}_i + \vec{F}_{langevin} - \nabla P_e / n_{ion}$$

where  $F_{langevin}$  is a force from Langevin thermostat simulating electron-phonon coupling, and  $\nabla P_e / n_{ion}$  is the electron blast force.

The electronic pressure is taken to be  $P_e = B \rho_e C_e T_e$

The current fix ttm/mod implementation allows TTM simulations with a vacuum. The vacuum region is defined as the grid cells with zero electronic temperature. The numerical scheme does not allow energy exchange with such cells. Since the material can expand to previously unoccupied region in some simulations, the vacuum border can be allowed to move. It is controlled by the *surface\_movement* parameter in the *init\_file*. If it is set to 1, then “vacuum” cells can be changed to “electron-filled” cells with the temperature  $T_{e\_min}$  if atoms move into them (currently only implemented for the case of 1-dimensional motion of flat surface normal to the X axis). The initial borders of vacuum can be set in the *init\_file* via *lsurface* and *rsurface* parameters. In this case, electronic pressure gradient is calculated as

$$\nabla_x P_e = \left[ \frac{C_e T_e(x) \lambda}{(x + \lambda)^2} + \frac{x}{x + \lambda} \frac{(C_e T_e)_{x+\Delta x} - (C_e T_e)_x}{\Delta x} \right]$$

where lambda is the electron mean free path (see (*Norman*), (*Pisarev*))

The fix ttm/mod parameter file *init\_file* has the following syntax. Every line with an odd number is considered as a comment and ignored. The lines with the even numbers are treated as follows:

```
a_0, energy/(temperature*electron) units
a_1, energy/(temperature^2*electron) units
a_2, energy/(temperature^3*electron) units
a_3, energy/(temperature^4*electron) units
a_4, energy/(temperature^5*electron) units
C_0, energy/(temperature*electron) units
A, 1/temperature units
rho_e, electrons/volume units
D_e, length^2/time units
gamma_p, mass/time units
gamma_s, mass/time units
v_0, length/time units
I_0, energy/(time*length^2) units
lsurface, electron grid units (positive integer)
rsurface, electron grid units (positive integer)
l_skin, length units
tau, time units
B, dimensionless
lambda, length units
n_ion, ions/volume units
surface_movement: 0 to disable tracking of surface motion, 1 to enable
T_e_min, temperature units
```

#### 17.224.4 Restart, fix\_modify, output, run start/stop, minimize info

These fixes write the state of the electronic subsystem and the energy exchange between the subsystems to *binary restart files*. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion. Note that the restart script must define the same size grid as the original script.

Because the state of the random number generator is not saved in the restart files, this means you cannot do “exact” restarts with this fix, where the simulation continues on the same as if no restart had taken place. However, in a statistical sense, a restarted simulation should produce the same behavior.

None of the *fix\_modify* options are relevant to these fixes.

These fixes compute 2 output quantities stored in a vector of length 2, which can be accessed by various *output commands*. The first quantity is the total energy of the electronic subsystem. The second quantity is the energy transferred

from the electronic to the atomic subsystem on that timestep. Note that the velocity verlet integrator applies the fix ttm forces to the atomic subsystem as two half-step velocity updates: one on the current timestep and one on the subsequent timestep. Consequently, the change in the atomic subsystem energy is lagged by half a timestep relative to the change in the electronic subsystem energy. As a result of this, users may notice slight fluctuations in the sum of the atomic and electronic subsystem energies reported at the end of the timestep.

The vector values calculated are “extensive”.

No parameter of the fixes can be used with the *start/stop* keywords of the *run* command. The fixes are not invoked during *energy minimization*.

### 17.224.5 Restrictions

All these fixes are part of the EXTRA-FIX package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

As mentioned above, these fixes require 3d simulations and orthogonal simulation boxes periodic in all 3 dimensions.

### 17.224.6 Related commands

*fix langevin*, *fix dt/reset*

### 17.224.7 Default

none

---

**(Duffy)** D M Duffy and A M Rutherford, J. Phys.: Condens. Matter, 19, 016207-016218 (2007).

**(Rutherford)** A M Rutherford and D M Duffy, J. Phys.: Condens. Matter, 19, 496201-496210 (2007).

**(Chen)** J Chen, D Tzou and J Beraun, Int. J. Heat Mass Transfer, 49, 307-316 (2006).

**(Norman)** G E Norman, S V Starikov, V V Stegailov et al., Contrib. Plasma Phys., 53, 129-139 (2013).

**(Pisarev)** V V Pisarev and S V Starikov, J. Phys.: Condens. Matter, 26, 475401 (2014).

## 17.225 fix tune/kspace command

### 17.225.1 Syntax

<code>fix ID group-ID tune/kspace N</code>
--

- ID, group-ID are documented in *fix* command
- tune/kspace = style name of this fix command
- N = invoke this fix every N steps



## 17.225.2 Examples

```
fix 2 all tune/kpace 100
```

## 17.225.3 Description

This fix tests each kspace style (Ewald, PPPM, and MSM), and automatically selects the fastest style to use for the remainder of the run. If the fastest style is Ewald or PPPM, the fix also adjusts the Coulombic cutoff towards optimal speed. Future versions of this fix will automatically select other kspace parameters to use for maximum simulation speed. The kspace parameters may include the style, cutoff, grid points in each direction, order, Ewald parameter, MSM parallelization cut-point, MPI tasks to use, etc.

The rationale for this fix is to provide the user with as-fast-as-possible simulations that include long-range electrostatics (kspace) while meeting the user-prescribed accuracy requirement. A simple heuristic could never capture the optimal combination of parameters for every possible run-time scenario. But by performing short tests of various kspace parameter sets, this fix allows parameters to be tailored specifically to the user's machine, MPI ranks, use of threading or accelerators, the simulated system, and the simulation details. In addition, it is possible that parameters could be evolved with the simulation on-the-fly, which is useful for systems that are dynamically evolving (e.g. changes in box size/shape or number of particles).

When this fix is invoked, LAMMPS will perform short timed tests of various parameter sets to determine the optimal parameters. Tests are performed on-the-fly, with a new test initialized every N steps. N should be chosen large enough so that adequate CPU time lapses between tests, thereby providing statistically significant timings. But N should not be chosen to be so large that an unfortunate parameter set test takes an inordinate amount of wall time to complete. An N of 100 for most problems seems reasonable. Once an optimal parameter set is found, that set is used for the remainder of the run.

This fix uses heuristics to guide its selection of parameter sets to test, but the actual timed results will be used to decide which set to use in the simulation.

It is not necessary to discard trajectories produced using sub-optimal parameter sets, or a mix of various parameter sets, since the user-prescribed accuracy will have been maintained throughout. However, some users may prefer to use this fix only to discover the optimal parameter set for a given setup that can then be used on subsequent production runs.

This fix starts with kspace parameters that are set by the user with the *kpace\_style* and *kpace\_modify* commands. The prescribed accuracy will be maintained by this fix throughout the simulation.

None of the *fix\_modify* options are relevant to this fix.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.225.4 Restrictions

This fix is part of the KSPACE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Do not set "neigh\_modify once yes" or else this fix will never be called. RENEIGHBORING is required.

This fix is not compatible with a hybrid pair style, long-range dispersion, TIP4P water support, or long-range point dipole support.

## 17.225.5 Related commands

*kpspace\_style*, *boundary kpspace\_modify*, *pair\_style lj/cut/coul/long*, *pair\_style lj/charmm/coul/long*, *pair\_style lj/long*, *pair\_style lj/long/coul/long*, *pair\_style buck/coul/long*

## 17.225.6 Default

## 17.226 fix vector command

### 17.226.1 Syntax

```
fix ID group-ID vector Nevery value1 value2 ...
```

- ID, group-ID are documented in *fix* command
- vector = style name of this fix command
- Nevery = use input values every this many timesteps
- one or more input values can be listed
- value = c\_ID, c\_ID[N], f\_ID, f\_ID[N], v\_name

```
c_ID = global scalar calculated by a compute with ID
c_ID[I] = Ith component of global vector calculated by a compute with ID
f_ID = global scalar calculated by a fix with ID
f_ID[I] = Ith component of global vector calculated by a fix with ID
v_name = value calculated by an equal-style variable with name
v_name[I] = Ith component of vector-style variable with name
```

### 17.226.2 Examples

```
fix 1 all vector 100 c_myTemp
fix 1 all vector 5 c_myTemp v_integral
```

### 17.226.3 Description

Use one or more global values as inputs every few timesteps, and simply store them. For a single specified value, the values are stored as a global vector of growing length. For multiple specified values, they are stored as rows in a global array, whose number of rows is growing. The resulting vector or array can be used by other *output commands*.

One way to use this command is to accumulate a vector that is time-integrated using the *variable trap()* function. For example the velocity auto-correlation function (VACF) can be time-integrated, to yield a diffusion coefficient, as follows:

```
compute      2 all vacf
fix          5 all vector 1 c_2[4]
variable     diff equal dt*trap(f_5)
thermo_style custom step v_diff
```

The group specified with this command is ignored. However, note that specified values may represent calculations performed by computes and fixes which store their own “group” definitions.

Each listed value can be the result of a *compute* or *fix* or the evaluation of an equal-style or vector-style *variable*. In each case, the compute, fix, or variable must produce a global quantity, not a per-atom or local quantity. And the global quantity must be a scalar, not a vector or array.

*Computes* that produce global quantities are those which do not have the word *atom* in their style name. Only a few *fixes* produce global quantities. See the doc pages for individual fixes for info on which ones produce such values. *Variables* of style *equal* or *vector* are the only ones that can be used with this fix. Variables of style *atom* cannot be used, since they produce per-atom values.

The *Nevery* argument specifies on what timesteps the input values will be used in order to be stored. Only timesteps that are a multiple of *Nevery*, including timestep 0, will contribute values.

Note that if you perform multiple runs, using the “pre no” option of the *run* command to avoid initialization on subsequent runs, then you need to use the *stop* keyword with the first *run* command with a timestep value that encompasses all the runs. This is so that the vector or array stored by this fix can be allocated to a sufficient size.

---

If a value begins with “c\_”, a compute ID must follow which has been previously defined in the input script. If no bracketed term is appended, the global scalar calculated by the compute is used. If a bracketed term is appended, the *l*th element of the global vector calculated by the compute is used.

Note that there is a *compute reduce* command which can sum per-atom quantities into a global scalar or vector which can thus be accessed by fix vector. Or it can be a compute defined not in your input script, but by *thermodynamic output* or other fixes such as *fix nvt* or *fix temp/rescale*. See the doc pages for these commands which give the IDs of these computes. Users can also write code for their own compute styles and *add them to LAMMPS*.

If a value begins with “f\_”, a fix ID must follow which has been previously defined in the input script. If no bracketed term is appended, the global scalar calculated by the fix is used. If a bracketed term is appended, the *l*th element of the global vector calculated by the fix is used.

Note that some fixes only produce their values on certain timesteps, which must be compatible with *Nevery*, else an error will result. Users can also write code for their own fix styles and *add them to LAMMPS*.

If a value begins with “v\_”, a variable name must follow which has been previously defined in the input script. An equal-style or vector-style variable can be referenced; the latter requires a bracketed term to specify the *l*th element of the vector calculated by the variable. See the *variable* command for details. Note that variables of style *equal* and *vector* define a formula which can reference individual atom properties or thermodynamic keywords, or they can invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of specifying quantities to be stored by fix vector.

---

#### 17.226.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix produces a global vector or global array which can be accessed by various *output commands*. The values can only be accessed on timesteps that are multiples of *Nevery*.

A vector is produced if only a single input value is specified. An array is produced if multiple input values are specified. The length of the vector or the number of rows in the array grows by 1 every *Nevery* timesteps.

If the fix produces a vector, then the entire vector will be either “intensive” or “extensive”, depending on whether the values stored in the vector are “intensive” or “extensive”. If the fix produces an array, then all elements in the array must be the same, either “intensive” or “extensive”. If a compute or fix provides the value stored, then the compute or

fix determines whether the value is intensive or extensive; see the page for that compute or fix for further info. Values produced by a variable are treated as intensive.

This fix can allocate storage for stored values accumulated over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this. If using the *run pre no* command option, this is required to allow the fix to allocate sufficient storage for stored values.

This fix is not invoked during *energy minimization*.

## 17.226.5 Restrictions

none

## 17.226.6 Related commands

*compute, variable*

## 17.226.7 Default

none

# 17.227 fix viscosity command

## 17.227.1 Syntax

```
fix ID group-ID viscosity N vdim pdim Nbin keyword value ...
```

- ID, group-ID are documented in *fix* command
- viscosity = style name of this fix command
- N = perform momentum exchange every N steps
- vdim = *x* or *y* or *z* = which momentum component to exchange
- pdim = *x* or *y* or *z* = direction of momentum transfer
- Nbin = # of layers in pdim direction (must be even number)
- zero or more keyword/value pairs may be appended
- keyword = *swap* or *target*
  - swap* value = Nswap = number of swaps to perform every N steps
  - target* value = V or INF = target velocity of swap partners (velocity units)

## 17.227.2 Examples

```
fix 1 all viscosity 100 x z 20
fix 1 all viscosity 50 x z 20 swap 2 vtarget 1.5
```

## 17.227.3 Description

Use the Muller-Plathe algorithm described in [this paper](#) to exchange momenta between two particles in different regions of the simulation box every  $N$  steps. This induces a shear velocity profile in the system. As described below this enables a viscosity of the fluid to be calculated. This algorithm is sometimes called a reverse non-equilibrium MD (reverse NEMD) approach to computing viscosity. This is because the usual NEMD approach is to impose a shear velocity profile on the system and measure the response via an off-diagonal component of the stress tensor, which is proportional to the momentum flux. In the Muller-Plathe method, the momentum flux is imposed, and the shear velocity profile is the system's response.

The simulation box is divided into  $N_{bin}$  layers in the  $pdim$  direction, where the layer 1 is at the low end of that dimension and the layer  $N_{bin}$  is at the high end. Every  $N$  steps,  $N_{swap}$  pairs of atoms are chosen in the following manner. Only atoms in the `fix` group are considered.  $N_{swap}$  atoms in layer 1 with positive velocity components in the  $vdim$  direction closest to the target value  $V$  are selected. Similarly,  $N_{swap}$  atoms in the “middle” layer (see below) with negative velocity components in the  $vdim$  direction closest to the negative of the target value  $V$  are selected. The two sets of  $N_{swap}$  atoms are paired up and their  $vdim$  momenta components are swapped within each pair. This resets their velocities, typically in opposite directions. Over time, this induces a shear velocity profile in the system which can be measured using commands such as the following, which writes the profile to the file `tmp.profile`:

```
compute layers all chunk/atom bin/1d z lower 0.05 units reduced
fix f1 all ave/chunk 100 10 1000 layers vx file tmp.profile
```

Note that by default,  $N_{swap} = 1$  and  $vtarget = INF$ , though this can be changed by the optional `swap` and `vtarget` keywords. When  $vtarget = INF$ , one or more atoms with the most positive and negative velocity components are selected. Setting these parameters appropriately, in conjunction with the swap rate  $N$ , allows the momentum flux rate to be adjusted across a wide range of values, and the momenta to be exchanged in large chunks or more smoothly.

The “middle” layer for momenta swapping is defined as the  $N_{bin}/2 + 1$  layer. Thus if  $N_{bin} = 20$ , the two swapping layers are 1 and 11. This should lead to a symmetric velocity profile since the two layers are separated by the same distance in both directions in a periodic sense. This is why  $N_{bin}$  is restricted to being an even number.

As described below, the total momentum transferred by these velocity swaps is computed by the `fix` and can be output. Dividing this quantity by time and the cross-sectional area of the simulation box yields a momentum flux. The ratio of momentum flux to the slope of the shear velocity profile is proportional to the viscosity of the fluid, in appropriate units. See the [Muller-Plathe paper](#) for details.

---

**Note:** If your system is periodic in the direction of the momentum flux, then the flux is going in 2 directions. This means the effective momentum flux in one direction is reduced by a factor of 2. You will see this in the equations for viscosity in the Muller-Plathe paper. LAMMPS is simply tallying momentum which does not account for whether or not your system is periodic; you must use the value appropriately to yield a viscosity for your system.

---

**Note:** After equilibration, if the velocity profile you observe is not linear, then you are likely swapping momentum too frequently and are not in a regime of linear response. In this case you cannot accurately infer a viscosity and should try increasing the `Nevery` parameter.

---

An alternative method for calculating a viscosity is to run a NEMD simulation, as described on the [Howto nemd](#) doc page. NEMD simulations deform the simulation box via the `fix deform` command.

Some features or combination of settings in LAMMPS do not support non-orthogonal boxes. Using `fix viscosity` keeps the box orthogonal; thus it does not suffer from these limitations.

#### 17.227.4 Restart, `fix_modify`, `output`, `run start/stop`, `minimize info`

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global scalar which can be accessed by various *output commands*. The scalar is the cumulative momentum transferred between the bottom and middle of the simulation box (in the *pdim* direction) is stored as a scalar quantity by this fix. This quantity is zeroed when the fix is defined and accumulates thereafter, once every N steps. The units of the quantity are momentum = mass\*velocity. The scalar value calculated by this fix is “intensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.227.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Swaps conserve both momentum and kinetic energy, even if the masses of the swapped atoms are not equal. Thus you should not need to thermostat the system. If you do use a thermostat, you may want to apply it only to the non-swapped dimensions (other than *vdim*).

LAMMPS does not check, but you should not use this fix to swap velocities of atoms that are in constrained molecules, e.g. via *fix shake* or *fix rigid*. This is because application of the constraints will alter the amount of transferred momentum. You should, however, be able to use flexible molecules. See the *Maginn paper* for an example of using this algorithm in a computation of alcohol molecule properties.

When running a simulation with large, massive particles or molecules in a background solvent, you may want to only exchange momenta between solvent particles.

#### 17.227.6 Related commands

*fix ave/chunk*, *fix thermal/conductivity*

#### 17.227.7 Default

The option defaults are `swap = 1` and `vtarget = INF`.

---

(Muller-Plathe) Muller-Plathe, Phys Rev E, 59, 4894-4898 (1999).

(Maginn) Kelkar, Rafferty, Maginn, Siepmann, Fluid Phase Equilibria, 260, 218-231 (2007).

## 17.228 fix viscous command

### 17.228.1 Syntax

```
fix ID group-ID viscous gamma keyword values ...
```

- ID, group-ID are documented in *fix* command
- viscous = style name of this fix command
- gamma = damping coefficient (force/velocity units)
- zero or more keyword/value pairs may be appended

keyword = *scale*

scale values = type ratio

type = atom type (1-N)

ratio = factor to scale the damping coefficient by

### 17.228.2 Examples

```
fix 1 flow viscous 0.1
fix 1 damp viscous 0.5 scale 3 2.5
```

### 17.228.3 Description

Add a viscous damping force to atoms in the group that is proportional to the velocity of the atom. The added force can be thought of as a frictional interaction with implicit solvent, i.e. the no-slip Stokes drag on a spherical particle. In granular simulations this can be useful for draining the kinetic energy from the system in a controlled fashion. If used without additional thermostating (to add kinetic energy to the system), it has the effect of slowly (or rapidly) freezing the system; hence it can also be used as a simple energy minimization technique.

The damping force  $F_i$  is given by  $F_i = -\gamma v_i$ . The larger the coefficient, the faster the kinetic energy is reduced. If the optional keyword *scale* is used,  $\gamma$  can be scaled up or down by the specified factor for atoms of that type. It can be used multiple times to adjust  $\gamma$  for several atom types.

---

**Note:** You should specify gamma in force/velocity units. This is not the same as mass/time units, at least for some of the LAMMPS *units* options like “real” or “metal” that are not self-consistent.

---

In a Brownian dynamics context,  $\gamma = \frac{k_B T}{D}$ , where  $k_B$  = Boltzmann’s constant,  $T$  = temperature, and  $D$  = particle diffusion coefficient.  $D$  can be written as  $\frac{k_B T}{3\pi\eta d}$ , where  $\eta$  = dynamic viscosity of the frictional fluid and  $d$  = diameter of particle. This means  $\gamma = 3\pi\eta d$ , and thus is proportional to the viscosity of the fluid and the particle diameter.

In the current implementation, rather than have the user specify a viscosity,  $\gamma$  is specified directly in force/velocity units. If needed,  $\gamma$  can be adjusted for atoms of different sizes (i.e.  $\sigma$ ) by using the *scale* keyword.

Note that Brownian dynamics models also typically include a randomized force term to thermostat the system at a chosen temperature. The *fix langevin* command does this. It has the same viscous damping term as *fix viscous* and adds a random force to each atom. The random force term is proportional to the square root of the chosen thermostating temperature. Thus if you use *fix langevin* with a target  $T = 0$ , its random force term is zero, and you are essentially performing the same operation as *fix viscous*. Also note that the gamma of *fix viscous* is related to the damping parameter of *fix langevin*, however the former is specified in units of force/velocity and the latter in units of time, so that it can more easily be used as a thermostat.

#### 17.228.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is modifying forces. Default is the outermost level.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command. This fix should only be used with damped dynamics minimizers that allow for non-conservative forces. See the *min\_style* command for details.

#### 17.228.5 Restrictions

none

#### 17.228.6 Related commands

*fix langevin*, *fix viscous/sphere*, *fix damping/cundall*

#### 17.228.7 Default

none

### 17.229 fix viscous/sphere command

#### 17.229.1 Syntax

```
fix ID group-ID viscous/sphere gamma keyword values ...
```

- ID, group-ID are documented in *fix* command
- viscous/sphere = style name of this fix command
- gamma = damping coefficient (torque/angular velocity units)
- zero or more keyword/value pairs may be appended

keyword = *scale*

*scale* values = *type ratio* or *v\_name*

*type* = atom type (1-N)

*ratio* = factor to scale the damping coefficients by

*v\_name* = reference to atom style variable *name*



## 17.229.2 Examples

```
fix 1 flow viscous/sphere 0.1
fix 1 damp viscous/sphere 0.5 scale 3 2.5
fix 1 damp viscous/sphere 0.5 scale v_radscale
```

## 17.229.3 Description

Add a viscous damping torque to finite-size spherical particles in the group that is proportional to the angular velocity of the atom. In granular simulations this can be useful for draining the rotational kinetic energy from the system in a controlled fashion. If used without additional thermostating (to add kinetic energy to the system), it has the effect of slowly (or rapidly) freezing the system; hence it can also be used as a simple energy minimization technique.

The damping torque  $T_i$  is given by  $T_i = -\gamma\omega_i$ . The larger the coefficient, the faster the rotational kinetic energy is reduced.

If the optional keyword *scale* is used,  $\gamma$  can be scaled up or down by the specified factor for atoms. This factor can be set for different atom types and thus the *scale* keyword used multiple times followed by the atom type and the associated scale factor. Alternately the scaling factor can be computed for each atom (e.g. based on its radius) by using an *atom-style variable*.

---

**Note:** You should specify gamma in torque/angular velocity units. This is not the same as mass/time units, at least for some of the LAMMPS *units* options like “real” or “metal” that are not self-consistent.

---

In the current implementation, rather than have the user specify a viscosity,  $\gamma$  is specified directly in torque/angular velocity units. If needed,  $\gamma$  can be adjusted for atoms of different sizes (i.e.  $\sigma$ ) by using the *scale* keyword.

---

## 17.229.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is modifying torques. Default is the outermost level.

The torques due to this fix are imposed during an energy minimization, invoked by the *minimize* command. This fix should only be used with damped dynamics minimizers that allow for non-conservative forces. See the *min\_style* command for details.

## 17.229.5 Restrictions

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This fix requires that atoms store torque and angular velocity (omega) and a radius as defined by the *atom\_style sphere* command.

All particles in the group must be finite-size spheres. They cannot be point particles.

## 17.229.6 Related commands

*fix viscous, fix damping/cundall*

## 17.229.7 Default

none

## 17.230 fix wall/lj93 command

Accelerator Variants: *wall/lj93/kk*

## 17.231 fix wall/lj126 command

## 17.232 fix wall/lj1043 command

## 17.233 fix wall/colloid command

## 17.234 fix wall/harmonic command

## 17.235 fix wall/morse command

### 17.235.1 Syntax

```
fix ID group-ID style face args ... keyword value ...
```

- ID, group-ID are documented in *fix* command
- style = *wall/lj93* or *wall/lj126* or *wall/lj1043* or *wall/colloid* or *wall/harmonic* or *wall/morse*
- one or more face/arg pairs may be appended
- face = *xlo* or *xhi* or *ylo* or *yhi* or *zlo* or *zhi*
- args for styles *lj93* or *lj126* or *lj1043* or *colloid* or *harmonic*

args = coord epsilon sigma cutoff

coord = position of wall = EDGE or constant or variable

EDGE = current lo or hi edge of simulation box

constant = number like 0.0 or -30.0 (distance units)

variable = *equal-style variable* like v\_x or v\_wiggle

epsilon = strength factor for wall-particle interaction (energy or energy/distance<sup>↔</sup>  
↔2 units)

epsilon can be a variable (see below)

sigma = size factor for wall-particle interaction (distance units)

sigma can be a variable (see below)

cutoff = distance from wall at which wall-particle interaction is cut off (distance<sub>↔</sub>  
↔units)

- args for style *morse*

```
args = coord D_0 alpha r_0 cutoff
coord = position of wall = EDGE or constant or variable
      EDGE = current lo or hi edge of simulation box
      constant = number like 0.0 or -30.0 (distance units)
      variable = equal-style variable like v_x or v_wiggle
D_0 = depth of the potential (energy units)
      D_0 can be a variable (see below)
alpha = width factor for wall-particle interaction (1/distance units)
      alpha can be a variable (see below)
r_0 = distance of the potential minimum from the face of region (distance units)
      r_0 can be a variable (see below)
cutoff = distance from wall at which wall-particle interaction is cut off (distance_
↪units)
```

- zero or more keyword/value pairs may be appended

- keyword = *units* or *fld*

```
units value = lattice or box
  lattice = the wall position is defined in lattice units
  box = the wall position is defined in simulation box units
fld value = yes or no
  yes = invoke the wall constraint to be compatible with implicit FLD
  no = invoke the wall constraint in the normal way
pbc value = yes or no
  yes = allow periodic boundary in a wall dimension
  no = require non-periodic boundaries in any wall dimension
```

## 17.235.2 Examples

```
fix wallhi all wall/lj93 xlo -1.0 1.0 1.0 2.5 units box
fix wallhi all wall/lj93 xhi EDGE 1.0 1.0 2.5
fix wallhi all wall/morse xhi EDGE 1.0 1.0 1.0 2.5 units box
fix wallhi all wall/lj126 v_wiggle 23.2 1.0 1.0 2.5
fix zwalls all wall/colloid zlo 0.0 1.0 1.0 0.858 zhi 40.0 1.0 1.0 0.858
```

## 17.235.3 Description

Bound the simulation domain on one or more of its faces with a flat wall that interacts with the atoms in the group by generating a force on the atom in a direction perpendicular to the wall. The energy of wall-particle interactions depends on the style.

For style *wall/lj93*, the energy  $E$  is given by the 9/3 potential:

$$E = \epsilon \left[ \frac{2}{15} \left( \frac{\sigma}{r} \right)^9 - \left( \frac{\sigma}{r} \right)^3 \right] \quad r < r_c$$

For style *wall/lj126*, the energy  $E$  is given by the 12/6 potential:

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_c$$

For style *wall/lj1043*, the energy E is given by the 10/4/3 potential:

$$E = 2\pi\epsilon \left[ \frac{2}{5} \left( \frac{\sigma}{r} \right)^{10} - \left( \frac{\sigma}{r} \right)^4 - \frac{\sqrt{(2)}\sigma^3}{3 \left( r + \left( 0.61/\sqrt{(2)} \right) \sigma \right)^3} \right] \quad r < r_c$$

For style *wall/colloid*, the energy E is given by an integrated form of the *pair\_style colloid* potential:

$$E = \epsilon \left[ \frac{\sigma^6}{7560} \left( \frac{6R - D}{D^7} + \frac{D + 8R}{(D + 2R)^7} \right) - \frac{1}{6} \left( \frac{2R(D + R) + D(D + 2R) [\ln D - \ln(D + 2R)]}{D(D + 2R)} \right) \right] \quad r < r_c$$

For style *wall/harmonic*, the energy E is given by a harmonic spring potential:

$$E = \epsilon (r - r_c)^2 \quad r < r_c$$

For style *wall/morse*, the energy E is given by a Morse potential:

$$E = D_0 \left[ e^{-2\alpha(r-r_0)} - 2e^{-\alpha(r-r_0)} \right] \quad r < r_c$$

In all cases,  $r$  is the distance from the particle to the wall at position *coord*, and  $r_c$  is the *cutoff* distance at which the particle and wall no longer interact. The energy of the wall potential is shifted so that the wall-particle interaction energy is 0.0 at the cutoff distance.

Up to 6 walls or faces can be specified in a single command: *xlo, xhi, ylo, yhi, zlo, zhi*. A *lo* face interacts with particles near the lower side of the simulation box in that dimension. A *hi* face interacts with particles near the upper side of the simulation box in that dimension.

The position of each wall can be specified in one of 3 ways: as the EDGE of the simulation box, as a constant value, or as a variable. If EDGE is used, then the corresponding boundary of the current simulation box is used. If a numeric constant is specified then the wall is placed at that position in the appropriate dimension (x, y, or z). In both the EDGE and constant cases, the wall will never move. If the wall position is a variable, it should be specified as *v\_name*, where name is an *equal\_style variable* name. In this case the variable is evaluated each timestep and the result becomes the current position of the reflecting wall. Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent wall position. See examples below.

For the *wall/lj93* and *wall/lj126* and *wall/lj1043* styles,  $\epsilon$  and  $\sigma$  are the usual Lennard-Jones parameters, which determine the strength and size of the particle as it interacts with the wall. Epsilon has energy units. Note that this  $\epsilon$  and  $\sigma$  may be different than any  $\epsilon$  or  $\sigma$  values defined for a pair style that computes particle-particle interactions.

The *wall/lj93* interaction is derived by integrating over a 3d half-lattice of Lennard-Jones 12/6 particles. The *wall/lj126* interaction is effectively a harder, more repulsive wall interaction. The *wall/lj1043* interaction is yet a different form of wall interaction, described in Magda et al in (*Magda*).

For the *wall/colloid* style,  $R$  is the radius of the colloid particle,  $D$  is the distance from the surface of the colloid particle to the wall ( $r-R$ ), and  $\sigma$  is the size of a constituent LJ particle inside the colloid particle and wall. Note that the cutoff distance  $R_c$  in this case is the distance from the colloid particle center to the wall. The prefactor  $\epsilon$  can be thought of as an effective Hamaker constant with energy units for the strength of the colloid-wall interaction. More specifically, the  $\epsilon$  prefactor is  $4\pi^2 \rho_{wall} \rho_{colloid} \epsilon \sigma^6$ , where  $\epsilon$  and  $\sigma$  are the LJ parameters for the constituent LJ particles.  $\rho_{wall}$  and  $\rho_{colloid}$  are the number density of the constituent particles, in the wall and colloid respectively, in units of 1/volume.

The *wall/colloid* interaction is derived by integrating over constituent LJ particles of size  $\sigma$  within the colloid particle and a 3d half-lattice of Lennard-Jones 12/6 particles of size  $\sigma$  in the wall. As mentioned in the preceding paragraph, the density of particles in the wall and colloid can be different, as specified by the  $\epsilon$  prefactor.

For the *wall/harmonic* style,  $\epsilon$  is effectively the spring constant  $K$ , and has units (energy/distance<sup>2</sup>). The input parameter  $\sigma$  is ignored. The minimum energy position of the harmonic spring is at the *cutoff*. This is a repulsive-only spring since the interaction is truncated at the *cutoff*.

For the *wall/morse* style, the three parameters are in this order:  $D_0$  the depth of the potential,  $\alpha$  the width parameter, and  $r_0$  the location of the minimum.  $D_0$  has energy units,  $\alpha$  inverse distance units, and  $r_0$  distance units.

For any wall, the  $\epsilon$  and/or  $\sigma$  and/or  $\alpha$  parameter can be specified as an *equal-style variable*, in which case it should be specified as `v_name`, where name is the variable name. As with a variable wall position, the variable is evaluated each timestep and the result becomes the current epsilon or sigma of the wall. Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent wall interaction.

---

**Note:** For all of the styles, you must insure that  $r$  is always  $> 0$  for all particles in the group, or LAMMPS will generate an error. This means you cannot start your simulation with particles at the wall position *coord* ( $r = 0$ ) or with particles on the wrong side of the wall ( $r < 0$ ). For the *wall/lj93* and *wall/lj126* styles, the energy of the wall/particle interaction (and hence the force on the particle) blows up as  $r \rightarrow 0$ . The *wall/colloid* style is even more restrictive, since the energy blows up as  $D = r - R \rightarrow 0$ . This means the finite-size particles of radius  $R$  must be a distance larger than  $R$  from the wall position *coord*. The *harmonic* style is a softer potential and does not blow up as  $r \rightarrow 0$ , but you must use a large enough  $\epsilon$  that particles always remain on the correct side of the wall ( $r > 0$ ).

---

The *units* keyword determines the meaning of the distance units used to define a wall position, but only when a numeric constant or variable is used. It is not relevant when *EDGE* is used to specify a face position. In the variable case, the variable is assumed to produce a value compatible with the *units* setting you specify.

A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacings.

The *fld* keyword can be used with a *yes* setting to invoke the wall constraint before pairwise interactions are computed. This allows an implicit FLD model using *pair\_style lubricateU* to include the wall force in its calculations. If the setting is *no*, wall forces are imposed after pairwise interactions, in the usual manner.

The *pbw* keyword can be used with a *yes* setting to allow walls to be specified in a periodic dimension. See the *boundary* command for options on simulation box boundaries. The default for *pbw* is *no*, which means the system must be non-periodic when using a wall. But you may wish to use a periodic box. E.g. to allow some particles to interact with the wall via the fix group-ID, and others to pass through it and wrap around a periodic box. In this case you should insure that the wall is sufficiently far enough away from the box boundary. If you do not, then particles may interact with both the wall and with periodic images on the other side of the box, which is probably not what you want.

---

Here are examples of variable definitions that move the wall position in a time-dependent fashion using equal-style variables. The wall interaction parameters (epsilon, sigma) could be varied with additional variable definitions.

```
variable ramp equal ramp(0,10)
fix 1 all wall xlo v_ramp 1.0 1.0 2.5

variable linear equal vdisplace(0,20)
fix 1 all wall xlo v_linear 1.0 1.0 2.5

variable wiggle equal swiggle(0.0,5.0,3.0)
fix 1 all wall xlo v_wiggle 1.0 1.0 2.5

variable wiggle equal cwiggle(0.0,5.0,3.0)
fix 1 all wall xlo v_wiggle 1.0 1.0 2.5
```

The *ramp(lo,hi)* function adjusts the wall position linearly from *lo* to *hi* over the course of a run. The *vdisplace(c0,velocity)* function does something similar using the equation  $position = c0 + velocity * delta$ , where *delta* is the elapsed time.

The *swiggle(c0,A,period)* function causes the wall position to oscillate sinusoidally according to this equation, where  $omega = 2 PI / period$ :

$$position = c0 + A \sin(omega * delta)$$

The *cwiggle(c0,A,period)* function causes the wall position to oscillate sinusoidally according to this equation, which will have an initial wall velocity of 0.0, and thus may impose a gentler perturbation on the particles:

$$position = c0 + A (1 - \cos(omega * delta))$$

---

### 17.235.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the energy of interaction between atoms and all the specified walls to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify virial* option is supported by this fix to add the contribution due to the interaction between atoms and all the specified walls to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global scalar energy and a global vector of forces, which can be accessed by various *output commands*. Note that the scalar energy is the sum of interactions with all defined walls. If you want the energy on a per-wall basis, you need to use multiple fix wall commands. The length of the vector is equal to the number of walls defined by the fix. Each vector value is the normal force on a specific wall. Note that an outward force on a wall will be a negative value for *lo* walls and a positive value for *hi* walls. The scalar and vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the atom/wall interaction energy to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 17.235.5 Restrictions

none

## 17.235.6 Related commands

*fix wall/reflect*, *fix wall/gran*, *fix wall/region*

## 17.235.7 Default

The option defaults units = lattice, fld = no, and pbc = no.

---

(Magda) Magda, Tirrell, Davis, J Chem Phys, 83, 1888-1901 (1985); erratum in JCP 84, 2901 (1986).

## 17.236 fix wall/body/polygon command

### 17.236.1 Syntax

```
fix ID group-ID wall/body/polygon k_n c_n c_t wallstyle args keyword values ...
```

- ID, group-ID are documented in *fix* command
- wall/body/polygon = style name of this fix command
- k\_n = normal repulsion strength (force/distance or pressure units)
- c\_n = normal damping coefficient (force/distance or pressure units)
- c\_t = tangential damping coefficient (force/distance or pressure units)
- wallstyle = *xplane* or *yplane* or *zylinder*
- args = list of arguments for a particular style
  - xplane* or *yplane* args = lo hi  
lo,hi = position of lower and upper plane (distance units), either can be NULL)
  - zylinder* args = radius  
radius = cylinder radius (distance units)
- zero or more keyword/value pairs may be appended to args
- keyword = *wiggle*
  - wiggle* values = dim amplitude period  
dim = x or y or z  
amplitude = size of oscillation (distance units)  
period = time of oscillation (time units)

## 17.236.2 Examples

```
fix 1 all wall/body/polygon 1000.0 20.0 5.0 xplane -10.0 10.0
```

## 17.236.3 Description

This fix is for use with 2d models of body particles of style *rounded/polygon*. It bounds the simulation domain with wall(s). All particles in the group interact with the wall when they are close enough to touch it. The nature of the interaction between the wall and the polygon particles is the same as that between the polygon particles themselves, which is similar to a Hookean potential. See the *Howto body* page for more details on using body particles.

The parameters *k<sub>n</sub>*, *c<sub>n</sub>*, *c<sub>t</sub>* have the same meaning and units as those specified with the *pair\_style body/rounded/polygon* command.

The *wallstyle* can be planar or cylindrical. The 2 planar options specify a pair of walls in a dimension. Wall positions are given by *lo* and *hi*. Either of the values can be specified as NULL if a single wall is desired. For a *zcylinder* wallstyle, the cylinder's axis is at  $x = y = 0.0$ , and the radius of the cylinder is specified.

Optionally, the wall can be moving, if the *wiggle* keyword is appended.

For the *wiggle* keyword, the wall oscillates sinusoidally, similar to the oscillations of particles which can be specified by the *fix move* command. This is useful in packing simulations of particles. The arguments to the *wiggle* keyword specify a dimension for the motion, as well as its *amplitude* and *period*. Note that if the dimension is in the plane of the wall, this is effectively a shearing motion. If the dimension is perpendicular to the wall, it is more of a shaking motion. A *zcylinder* wall can only be wiggled in the z dimension.

Each timestep, the position of a wiggled wall in the appropriate *dim* is set according to this equation:

$$\text{position} = \text{coord} + A - A \cos(\omega * \text{delta})$$

where *coord* is the specified initial position of the wall, *A* is the *amplitude*, *omega* is  $2 \text{PI} / \text{period}$ , and *delta* is the time elapsed since the fix was specified. The velocity of the wall is set to the derivative of this expression.

## 17.236.4 Restart, fix\_modify, output, run start/stop, minimize info

None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.236.5 Restrictions

This fix is part of the BODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Any dimension (xy) that has a wall must be non-periodic.



## 17.236.6 Related commands

*atom\_style body, pair\_style body/rounded/polygon*

## 17.236.7 Default

none

## 17.237 fix wall/body/polyhedron command

### 17.237.1 Syntax

```
fix ID group-ID wall/body/polyhedron k_n c_n c_t wallstyle args keyword values ...
```

- ID, group-ID are documented in *fix* command
- wall/body/polyhedron = style name of this fix command
- k\_n = normal repulsion strength (force/distance units or pressure units - see discussion below)
- c\_n = normal damping coefficient (force/distance units or pressure units - see discussion below)
- c\_t = tangential damping coefficient (force/distance units or pressure units - see discussion below)
- wallstyle = *xplane* or *yplane* or *zplane*
- args = list of arguments for a particular style  
*xplane* or *yplane* or *zplane* args = lo hi  
lo,hi = position of lower and upper plane (distance units), either can be NULL)
- zero or more keyword/value pairs may be appended to args
- keyword = *wiggle*  
*wiggle* values = dim amplitude period  
dim = x or y or z  
amplitude = size of oscillation (distance units)  
period = time of oscillation (time units)

### 17.237.2 Examples

```
fix 1 all wall/body/polyhedron 1000.0 20.0 5.0 xplane -10.0 10.0
```

### 17.237.3 Description

This fix is for use with 3d models of body particles of style *rounded/polyhedron*. It bounds the simulation domain with wall(s). All particles in the group interact with the wall when they are close enough to touch it. The nature of the interaction between the wall and the polygon particles is the same as that between the polygon particles themselves, which is similar to a Hookean potential. See the *Howto body* page for more details on using body particles.

The parameters *k\_n*, *c\_n*, *c\_t* have the same meaning and units as those specified with the *pair\_style body/rounded/polyhedron* command.

The *wallstyle* can be planar or cylindrical. The 3 planar options specify a pair of walls in a dimension. Wall positions are given by *lo* and *hi*. Either of the values can be specified as NULL if a single wall is desired.

Optionally, the wall can be moving, if the *wiggle* keyword is appended.

For the *wiggle* keyword, the wall oscillates sinusoidally, similar to the oscillations of particles which can be specified by the *fix move* command. This is useful in packing simulations of particles. The arguments to the *wiggle* keyword specify a dimension for the motion, as well as its *amplitude* and *period*. Note that if the dimension is in the plane of the wall, this is effectively a shearing motion. If the dimension is perpendicular to the wall, it is more of a shaking motion.

Each timestep, the position of a wiggled wall in the appropriate *dim* is set according to this equation:

$$\text{position} = \text{coord} + A - A \cos(\omega * \text{delta})$$

where *coord* is the specified initial position of the wall, *A* is the *amplitude*, *omega* is  $2 \text{PI} / \text{period}$ , and *delta* is the time elapsed since the fix was specified. The velocity of the wall is set to the derivative of this expression.

### 17.237.4 Restart, fix\_modify, output, run start/stop, minimize info

None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.237.5 Restrictions

This fix is part of the BODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Any dimension (xyz) that has a wall must be non-periodic.

### 17.237.6 Related commands

*atom\_style body*, *pair\_style body/rounded/polyhedron*

## 17.237.7 Default

none

## 17.238 fix wall/ees command

## 17.239 fix wall/region/ees command

### 17.239.1 Syntax

```
fix ID group-ID style args
```

- ID, group-ID are documented in *fix* command
- style = *wall/ees* or *wall/region/ees*

args for style *wall/ees*: one or more *face parameters* groups may be appended

face = *xlo* or *xhi* or *ylo* or *yhi* or *zlo* or *zhi*

parameters = coord epsilon sigma cutoff

coord = position of wall = EDGE or constant or variable

EDGE = current lo or hi edge of simulation box

constant = number like 0.0 or -30.0 (distance units)

variable = *equal-style variable* like *v\_x* or *v\_wiggle*

epsilon = strength factor for wall-particle interaction (energy or energy/  
↪ distance<sup>2</sup> units)

epsilon can be a variable (see below)

sigma = size factor for wall-particle interaction (distance units)

sigma can be a variable (see below)

cutoff = distance from wall at which wall-particle interaction is cut off  
↪ (distance units)

args for style *wall/region/ees*: *region-ID epsilon sigma cutoff*

region-ID = region whose boundary will act as wall

epsilon = strength factor for wall-particle interaction (energy or energy/  
↪ distance<sup>2</sup> units)

sigma = size factor for wall-particle interaction (distance units)

cutoff = distance from wall at which wall-particle interaction is cut off  
↪ (distance units)

### 17.239.2 Examples

```
fix wallhi all wall/ees xlo -1.0 1.0 1.0 2.5 units box
fix wallhi all wall/ees xhi EDGE 1.0 1.0 2.5
fix wallhi all wall/ees v_wiggle 23.2 1.0 1.0 2.5
fix zwalls all wall/ees zlo 0.0 1.0 1.0 0.858 zhi 40.0 1.0 1.0 0.858

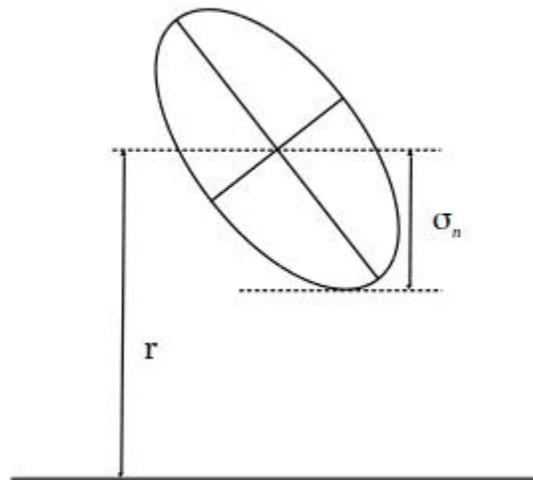
fix ees_cube all wall/region/ees myCube 1.0 1.0 2.5
```

### 17.239.3 Description

Fix *wall/ees* bounds the simulation domain on one or more of its faces with a flat wall that interacts with the ellipsoidal atoms in the group by generating a force on the atom in a direction perpendicular to the wall and a torque parallel with the wall. The energy of wall-particle interactions  $E$  is given by:

$$E = \epsilon \left[ \frac{2\sigma_{LJ}^{12} (7r^5 + 14r^3\sigma_n^2 + 3r\sigma_n^4)}{945 (r^2 - \sigma_n^2)^7} - \frac{\sigma_{LJ}^6 (2r\sigma_n^3 + \sigma_n^2 (r^2 - \sigma_n^2) \log \left[ \frac{r - \sigma_n}{r + \sigma_n} \right])}{12\sigma_n^5 (r^2 - \sigma_n^2)} \right] \quad \sigma_n < r < r_c$$

Introduced by Babadi and Ejtehad in (*Babadi*). Here,  $r$  is the distance from the particle to the wall at position *coord*, and  $R_c$  is the *cutoff* distance at which the particle and wall no longer interact. Also,  $\sigma_n$  is the distance between center of ellipsoid and the nearest point of its surface to the wall as shown below.



Details of using this command and specifications are the same as *fix/wall* command. You can also find an example in *USER/ees/* under *examples/* directory.

The prefactor  $\epsilon$  can be thought of as an effective Hamaker constant with energy units for the strength of the ellipsoid-wall interaction. More specifically, the  $\epsilon$  prefactor is

$$8\pi^2 \rho_{wall} \rho_{ellipsoid} \epsilon \sigma_a \sigma_b \sigma_c$$

where  $\epsilon$  is the LJ energy parameter for the constituent LJ particles and  $\sigma_a$ ,  $\sigma_b$ , and  $\sigma_c$  are the radii of the ellipsoidal particles.  $\rho_{wall}$  and  $\rho_{ellipsoid}$  are the number density of the constituent particles, in the wall and ellipsoid respectively, in units of 1/volume.

---

**Note:** You must insure that  $r$  is always bigger than  $\sigma_n$  for all particles in the group, or LAMMPS will generate an error. This means you cannot start your simulation with particles touching the wall position *coord* ( $r = \sigma_n$ ) or with particles penetrating the wall ( $0 = < r < \sigma_n$ ) or with particles on the wrong side of the wall ( $r < 0$ ).

---

Fix *wall/region/ees* treats the surface of the geometric region defined by the *region-ID* as a bounding wall which interacts with nearby ellipsoidal particles according to the EES potential introduced above.

Other details of this command are the same as for the *fix wall/region* command. One may also find an example of using this *fix* in the *examples/PACKAGES/ees/* directory.

---

## 17.239.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about these fixes are written to *binary restart files*.

The *fix\_modify energy* option is supported by these fixes to add the energy of interaction between atoms and all the specified walls or region wall to the global potential energy of the system as part of *thermodynamic output*. The default settings for these fixes are *fix\_modify energy no*.

The *fix\_modify respa* option is supported by these fixes. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

These fixes computes a global scalar and a global vector of forces, which can be accessed by various *output commands*. See the *fix wall* command for a description of the scalar and vector.

No parameter of these fixes can be used with the *start/stop* keywords of the *run* command.

The forces due to these fixes are imposed during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the atom/wall interaction energy to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

## 17.239.5 Restrictions

These fixes are part of the EXTRA-FIX package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These fixes requires that atoms be ellipsoids as defined by the *atom\_style ellipsoid* command.

## 17.239.6 Related commands

*fix wall*, *pair resquared*

## 17.239.7 Default

none

---

(Babadi) Babadi and Ejtehadi, EPL, 77 (2007) 23002.

## 17.240 fix wall/gran command

### 17.240.1 Syntax

```
fix ID group-ID wall/gran fstyle fstyle_params wallstyle args keyword values ...
```

- ID, group-ID are documented in *fix* command
- wall/gran = style name of this fix command
- fstyle = style of force interactions between particles and wall

```
possible choices: hooke, hooke/history, hertz/history, granular
```

- `fstyle_params` = parameters associated with force interaction style

For *hooke*, *hooke/history*, and *hertz/history*, `fstyle_params` are:

```
Kn = elastic constant for normal particle repulsion (force/distance units or
↪pressure units - see discussion below)
Kt = elastic constant for tangential contact (force/distance units or
↪pressure units - see discussion below)
gamma_n = damping coefficient for collisions in normal direction (1/time
↪units or 1/time-distance units - see discussion below)
gamma_t = damping coefficient for collisions in tangential direction (1/time
↪units or 1/time-distance units - see discussion below)
xmu = static yield criterion (unitless value between 0.0 and 1.0e4)
dampflag = 0 or 1 if tangential damping force is excluded or included
optional keyword = limit_damping, limit damping to prevent attractive
↪interaction
```

For *granular*, `fstyle_params` are set using the same syntax as for the `pair_coeff` command of `pair_style granular`

- `wallstyle` = *xplane* or *yplane* or *zplane* or *zcylinder*
- `args` = list of arguments for a particular style
  - xplane* or *yplane* or *zplane* `args` = lo hi
    - lo,hi = position of lower and upper plane (distance units), either can be NULL)
  - zcylinder* `args` = radius
    - radius = cylinder radius (distance units)
- zero or more keyword/value pairs may be appended to `args`
- `keyword` = *wiggle* or *shear* or *contacts*
  - wiggle* `values` = dim amplitude period
    - dim = x or y or z
    - amplitude = size of oscillation (distance units)
    - period = time of oscillation (time units)
  - shear* `values` = dim vshear
    - dim = x or y or z
    - vshear = magnitude of shear velocity (velocity units)
  - contacts* `value` = none
    - generate contact information for each particle

## 17.240.2 Examples

```
fix 1 all wall/gran hooke 200000.0 NULL 50.0 NULL 0.5 0 xplane -10.0 10.0
fix 1 all wall/gran hooke/history 200000.0 NULL 50.0 NULL 0.5 0 zplane 0.0 NULL
fix 2 all wall/gran hooke 100000.0 20000.0 50.0 30.0 0.5 1 zcylinder 15.0 wiggle z 3.0 2.
↪0
fix 3 all wall/gran/region granular hooke 1000.0 50.0 tangential linear_nohistory 1.0 0.
↪4 damping velocity region myBox
fix 4 all wall/gran/region granular jkr 1e5 1500.0 0.3 10.0 tangential mindlin NULL 1.0
↪0.5 rolling sds 500.0 200.0 0.5 twisting marshall region myCone
fix 5 all wall/gran/region granular dmt 1e5 0.2 0.3 10.0 tangential mindlin NULL 1.0 0.5
```

(continues on next page)

(continued from previous page)

```
↪rolling sds 500.0 200.0 0.5 twisting marshall damping tsuji region myCone
fix 6 all wall/gran hooke 200000.0 NULL 50.0 NULL 0.5 0 xplane -10.0 10.0 contacts
```

### 17.240.3 Description

Bound the simulation domain of a granular system with a frictional wall. All particles in the group interact with the wall when they are close enough to touch it.

The nature of the wall/particle interactions are determined by the *fstyle* setting. It can be any of the styles defined by the *pair\_style gran/\** or the more general *pair\_style granular* commands. Currently the options are *hooke*, *hooke/history*, or *hertz/history* for the former, and *granular* with all the possible options of the associated *pair\_coeff* command for the latter. The equation for the force between the wall and particles touching it is the same as the corresponding equation on the *pair\_style gran/\** and *pair\_style granular* doc pages, in the limit of one of the two particles going to infinite radius and mass (flat wall). Specifically,  $\delta = \text{radius} - r = \text{overlap of particle with wall}$ ,  $m_{\text{eff}} = \text{mass of particle}$ , and the effective radius of contact  $= R_i R_j / (R_i + R_j)$  is set to the radius of the particle.

The parameters *Kn*, *Kt*, *gamma\_n*, *gamma\_t*, *xmu*, *dampflag*, and the optional keyword *limit\_damping* have the same meaning and units as those specified with the *pair\_style gran/\** commands. This means a NULL can be used for either *Kt* or *gamma\_t* as described on that page. If a NULL is used for *Kt*, then a default value is used where  $Kt = 2/7 Kn$ . If a NULL is used for *gamma\_t*, then a default value is used where  $gamma_t = 1/2 gamma_n$ .

All the model choices for cohesion, tangential friction, rolling friction and twisting friction supported by the *pair\_style granular* through its *pair\_coeff* command are also supported for walls. These are discussed in greater detail on the doc page for *pair\_style granular*.

Note that you can choose a different force styles and/or different values for the wall/particle coefficients than for particle/particle interactions. E.g. if you wish to model the wall as a different material.

---

**Note:** As discussed on the page for *pair\_style gran/\**, versions of LAMMPS before 9Jan09 used a different equation for Hertzian interactions. This means Hertzian wall/particle interactions have also changed. They now include a  $\sqrt{\text{radius}}$  term which was not present before. Also the previous versions used *Kn* and *Kt* from the pairwise interaction and hardwired *dampflag* to 1, rather than letting them be specified directly. This means you can set the values of the wall/particle coefficients appropriately in the current code to reproduce the results of a previous Hertzian monodisperse calculation. For example, for the common case of a monodisperse system with particles of diameter 1, *Kn*, *Kt*, *gamma\_n*, and *gamma\_s* should be set  $\sqrt{2.0}$  larger than they were previously.

---

The effective mass  $m_{\text{eff}}$  in the formulas listed on the *pair\_style granular* page is the mass of the particle for particle/wall interactions (mass of wall is infinite). If the particle is part of a rigid body, its mass is replaced by the mass of the rigid body in those formulas. This is determined by searching for a *fix rigid* command (or its variants).

The *wallstyle* can be planar or cylindrical. The 3 planar options specify a pair of walls in a dimension. Wall positions are given by *lo* and *hi*. Either of the values can be specified as NULL if a single wall is desired. For a *zylinder* wallstyle, the cylinder's axis is at  $x = y = 0.0$ , and the radius of the cylinder is specified.

Optionally, the wall can be moving, if the *wiggle* or *shear* keywords are appended. Both keywords cannot be used together.

For the *wiggle* keyword, the wall oscillates sinusoidally, similar to the oscillations of particles which can be specified by the *fix move* command. This is useful in packing simulations of granular particles. The arguments to the *wiggle* keyword specify a dimension for the motion, as well as its *amplitude* and *period*. Note that if the dimension is in the plane of the wall, this is effectively a shearing motion. If the dimension is perpendicular to the wall, it is more of a shaking motion. A *zylinder* wall can only be wiggled in the *z* dimension.

Each timestep, the position of a wiggled wall in the appropriate *dim* is set according to this equation:

$\text{position} = \text{coord} + A - A \cos(\omega * \text{delta})$

where *coord* is the specified initial position of the wall, *A* is the *amplitude*, *omega* is  $2 \text{PI} / \text{period}$ , and *delta* is the time elapsed since the fix was specified. The velocity of the wall is set to the derivative of this expression.

For the *shear* keyword, the wall moves continuously in the specified dimension with velocity *vshear*. The dimension must be tangential to walls with a planar *wallstyle*, e.g. in the *y* or *z* directions for an *xplane* wall. For *zcylinder* walls, a dimension of *z* means the cylinder is moving in the *z*-direction along its axis. A dimension of *x* or *y* means the cylinder is spinning around the *z*-axis, either in the clockwise direction for *vshear* > 0 or counter-clockwise for *vshear* < 0. In this case, *vshear* is the tangential velocity of the wall at whatever *radius* has been defined.

#### 17.240.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the shear friction state of atoms interacting with the wall to *binary restart files*, so that a simulation can continue correctly if granular potentials with shear “history” effects are being used. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

If the *contacts* option is used, this fix generates a per-atom array with 8 columns as output, containing the contact information for owned particles (*nlocal* on each processor). All columns in this per-atom array will be zero if no contact has occurred. The values of these columns are listed in the following table:

Index	Value	Units
1	1.0 if particle is in contact with wall, 0.0 otherwise	
2	Force $f_x$ exerted by the wall	force units
3	Force $f_y$ exerted by the wall	force units
4	Force $f_z$ exerted by the wall	force units
5	<i>x</i> -coordinate of contact point on wall	distance units
6	<i>y</i> -coordinate of contact point on wall	distance units
7	<i>z</i> -coordinate of contact point on wall	distance units
8	Radius <i>r</i> of atom	distance units

None of the *fix\_modify* options are relevant to this fix. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.240.5 Restrictions

This fix is part of the GRANULAR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Any dimension (xyz) that has a granular wall must be non-periodic.

#### 17.240.6 Related commands

*fix move*, *fix wall/gran/region*, *pair\_style gran/\* pair\_style granular*



## 17.240.7 Default

none

## 17.241 fix wall/gran/region command

### 17.241.1 Syntax

```
fix ID group-ID wall/gran/region fstyle fstyle_params wallstyle regionID keyword values .
↪...
```

- ID, group-ID are documented in *fix* command
- wall/region = style name of this fix command
- fstyle = style of force interactions between particles and wall

```
possible choices: hooke, hooke/history, hertz/history, granular
```

- fstyle\_params = parameters associated with force interaction style

For *hooke*, *hooke/history*, and *hertz/history*, *fstyle\_params* are:

Kn = elastic constant for normal particle repulsion (force/distance units or ↪  
↪pressure units - see discussion below)

Kt = elastic constant for tangential contact (force/distance units or ↪  
↪pressure units - see discussion below)

gamma\_n = damping coefficient for collisions in normal direction (1/time ↪  
↪units or 1/time-distance units - see discussion below)

gamma\_t = damping coefficient for collisions in tangential direction (1/time ↪  
↪units or 1/time-distance units - see discussion below)

xmu = static yield criterion (unitless value between 0.0 and 1.0e4)

dampflag = 0 or 1 if tangential damping force is excluded or included

For *granular*, *fstyle\_params* are set using the same syntax as for the *pair\_coeff* ↪

↪command of *pair\_style granular*

- wallstyle = region (see *fix wall/gran* for options for other kinds of walls)
- region-ID = region whose boundary will act as wall
- keyword = *contacts*

*contacts* value = none

generate contact information for each particle

### 17.241.2 Examples

```
fix wall all wall/gran/region hooke/history 1000.0 200.0 200.0 100.0 0.5 1 region myCone
fix 3 all wall/gran/region granular hooke 1000.0 50.0 tangential linear_nohistory 1.0 0.
↪4 damping velocity region myBox
fix 4 all wall/gran/region granular jkr 1e5 1500.0 0.3 10.0 tangential mindlin NULL 1.0 ↪
↪0.5 rolling sds 500.0 200.0 0.5 twisting marshall region myCone
fix 5 all wall/gran/region granular dmt 1e5 0.2 0.3 10.0 tangential mindlin NULL 1.0 0.5 ↪
↪rolling sds 500.0 200.0 0.5 twisting marshall damping tsuji region myCone
```

(continues on next page)

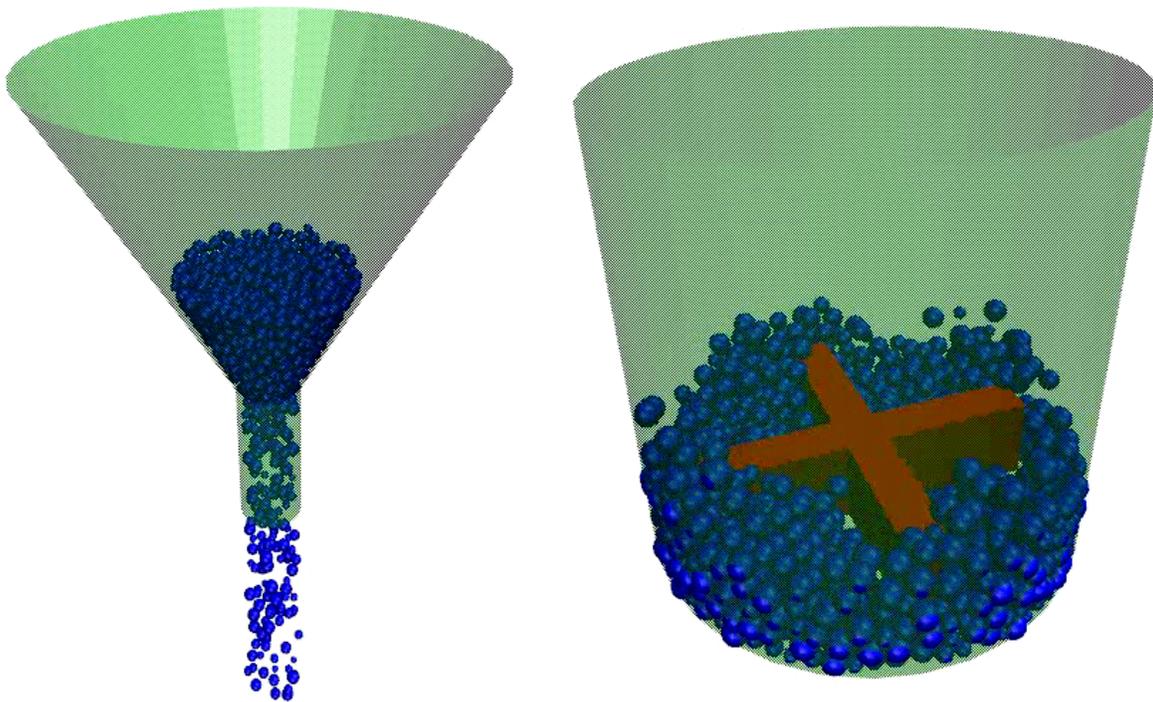
(continued from previous page)

```
fix wall all wall/gran/region hooke/history 1000.0 200.0 200.0 100.0 0.5 1 region myCone_
↪contacts
```

### 17.241.3 Description

Treat the surface of the geometric region defined by the *region-ID* as a bounding frictional wall which interacts with nearby finite-size granular particles when they are close enough to touch the wall. See the *fix wall/region* and *fix wall/gran* commands for related kinds of walls for non-granular particles and simpler wall geometries, respectively.

Here are snapshots of example models using this command. Corresponding input scripts can be found in [examples/granregion](#). Movies of these simulations are [here on the Movies page](#) of the LAMMPS website.



The distance between a particle and the region boundary is the distance to the nearest point on the region surface. The force the wall exerts on the particle is along the direction between that point and the particle center, which is the direction normal to the surface at that point. Note that if the region surface is comprised of multiple “faces”, then each face can exert a force on the particle if it is close enough. E.g. for *region\_style block*, a particle in the interior, near a corner of the block, could feel wall forces from 1, 2, or 3 faces of the block.

Regions are defined using the *region* command. Note that the region volume can be interior or exterior to the bounding surface, which will determine in which direction the surface interacts with particles, i.e. the direction of the surface normal. The exception to this is if one or more *open* options are specified for the region command, in which case particles interact with both the interior and exterior surfaces of regions.

Regions can either be primitive shapes (block, sphere, cylinder, etc) or combinations of primitive shapes specified via the *union* or *intersect* region styles. These latter styles can be used to construct particle containers with complex shapes.

Regions can also move dynamically via the *region* command keywords (*move*) and *rotate*, or change their shape by use of variables as inputs to the *region* command. If such a region is used with this *fix*, then the region surface will move in time in the corresponding manner.

---

**Note:** As discussed on the [region](#) command doc page, regions in LAMMPS do not get wrapped across periodic boundaries. It is up to you to ensure that the region location with respect to periodic or non-periodic boundaries is specified appropriately via the [region](#) and [boundary](#) commands when using a region as a wall that bounds particle motion.

---

**Note:** For primitive regions with sharp corners and/or edges (e.g. a block or cylinder), wall/particle forces are computed accurately for both interior and exterior regions. For *union* and *intersect* regions, additional sharp corners and edges may be present due to the intersection of the surfaces of 2 or more primitive volumes. These corners and edges can be of two types: concave or convex. Concave points/edges are like the corners of a cube as seen by particles in the interior of a cube. Wall/particle forces around these features are computed correctly. Convex points/edges are like the corners of a cube as seen by particles exterior to the cube, i.e. the points jut into the volume where particles are present. LAMMPS does NOT compute the location of these convex points directly, and hence wall/particle forces in the cutoff volume around these points suffer from inaccuracies. The basic problem is that the outward normal of the surface is not continuous at these points. This can cause particles to feel no force (they don't "see" the wall) when in one location, then move a distance epsilon, and suddenly feel a large force because they now "see" the wall. In a worst-case scenario, this can blow particles out of the simulation box. Thus, as a general rule you should not use the `fix wall/gran/region` command with *union* or *intersect* regions that have convex points or edges resulting from the union/intersection (convex points/edges in the union/intersection due to a single sub-region are still OK).

---

**Note:** Similarly, you should not define *union* or *intersect* regions for use with this command that share an overlapping common face that is part of the overall outer boundary (interior boundary is OK), even if the face is smooth. E.g. two regions of style `block` in a *union* region, where the two blocks overlap on one or more of their faces. This is because LAMMPS discards points that are part of multiple sub-regions when calculating wall/particle interactions, to avoid double-counting the interaction. Having two coincident faces could cause the face to become invisible to the particles. The solution is to make the two faces differ by epsilon in their position.

---

The nature of the wall/particle interactions are determined by the *fstyle* setting. It can be any of the styles defined by the [pair\\_style gran/\\*](#) or the more general [pair\\_style granular](#) commands. Currently the options are *hooke*, *hooke/history*, or *hertz/history* for the former, and *granular* with all the possible options of the associated [pair\\_coeff](#) command for the latter. The equation for the force between the wall and particles touching it is the same as the corresponding equation on the [pair\\_style gran/\\*](#) and [pair\\_style granular](#) doc pages, but the effective radius is calculated using the radius of the particle and the radius of curvature of the wall at the contact point.

Specifically,  $\delta = \text{radius} - r = \text{overlap of particle with wall}$ ,  $m_{\text{eff}} = \text{mass of particle}$ , and  $R_i R_j / (R_i + R_j)$  is the effective radius, with  $R_j$  replaced by the radius of curvature of the wall at the contact point. The radius of curvature can be negative for a concave wall section, e.g. the interior of cylinder. For a flat wall,  $\delta = \text{radius} - r = \text{overlap of particle with wall}$ ,  $m_{\text{eff}} = \text{mass of particle}$ , and the effective radius of contact is just the radius of the particle.

The parameters *Kn*, *Kt*, *gamma\_n*, *gamma\_t*, *xmu*, *dampflag*, and the optional keyword *limit\_damping* have the same meaning and units as those specified with the [pair\\_style gran/\\*](#) commands. This means a NULL can be used for either *Kt* or *gamma\_t* as described on that page. If a NULL is used for *Kt*, then a default value is used where  $Kt = 2/7 Kn$ . If a NULL is used for *gamma\_t*, then a default value is used where  $\gamma_t = 1/2 \gamma_n$ .

All the model choices for cohesion, tangential friction, rolling friction and twisting friction supported by the [pair\\_style granular](#) through its [pair\\_coeff](#) command are also supported for walls. These are discussed in greater detail on the doc page for [pair\\_style granular](#).

Note that you can choose a different force styles and/or different values for the 6 wall/particle coefficients than for particle/particle interactions. E.g. if you wish to model the wall as a different material.

### 17.241.4 Restart, fix\_modify, output, run start/stop, minimize info

Similar to *fix wall/gran* command, this fix writes the shear friction state of atoms interacting with the wall to *binary restart files*, so that a simulation can continue correctly if granular potentials with shear “history” effects are being used. This fix also includes info about a moving region in the restart file. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

---

**Note:** Information about region definitions is NOT included in restart files, as discussed on the *read\_restart* doc page. So you must re-define your region and if it is a moving region, define its motion attributes in a way that is consistent with the simulation that wrote the restart file. In particular, if you want to change the region motion attributes (e.g. its velocity), then you should ensure the position/orientation of the region at the initial restart timestep is the same as it was on the timestep the restart file was written. If this is not possible, you may need to ignore info in the restart file by defining a new *fix wall/gran/region* command in your restart script, e.g. with a different fix ID. Or if you want to keep the shear history info but discard the region motion information, you can use the same fix ID for *fix wall/gran/region*, but assign it a region with a different region ID.

---

If the *contacts* option is used, this fix generates a per-atom array with 8 columns as output, containing the contact information for owned particles (nlocal on each processor). All columns in this per-atom array will be zero if no contact has occurred. The values of these columns are listed in the following table:

Index	Value	Units
1	1.0 if particle is in contact with wall, 0.0 otherwise	
2	Force $f_x$ exerted by the wall	force units
3	Force $f_y$ exerted by the wall	force units
4	Force $f_z$ exerted by the wall	force units
5	$x$ -coordinate of contact point on wall	distance units
6	$y$ -coordinate of contact point on wall	distance units
7	$z$ -coordinate of contact point on wall	distance units
8	Radius $r$ of atom	distance units

None of the *fix\_modify* options are relevant to this fix. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.241.5 Restrictions

This fix is part of the GRANULAR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 17.241.6 Related commands

*fix\_move*, *fix wall/gran*, *fix wall/region*, *pair\_style granular*, *region*

## 17.241.7 Default

none

## 17.242 fix wall/piston command

### 17.242.1 Syntax

```
fix ID group-ID wall/piston face ... keyword value ...
```

- ID, group-ID are documented in *fix* command
- wall/piston = style name of this fix command
- face = *zlo*
- zero or more keyword/value pairs may be appended
- keyword = *pos* or *vel* or *ramp* or *units*

*pos* args = *z*

*z* = *z* coordinate at which the piston begins (distance units)

*vel* args = *vz*

*vz* = final velocity of the piston (velocity units)

*ramp* = use a linear velocity ramp from 0 to *vz*

*temp* args = target damp seed extent

target = target velocity for region immediately ahead of the piston

damp = damping parameter (time units)

seed = random number seed for langevin kicks

extent = extent of thermostatted region (distance units)

*units* value = *lattice* or *box*

*lattice* = the wall position is defined in lattice units

*box* = the wall position is defined in simulation box units

### 17.242.2 Examples

```
fix xwalls all wall/piston zlo
fix walls all wall/piston zlo pos 1.0 vel 10.0 units box
fix top all wall/piston zlo vel 10.0 ramp
```

### 17.242.3 Description

Bound the simulation with a moving wall which reflect particles in the specified group and drive the system with an effective infinite-mass piston capable of driving shock waves.

A momentum mirror technique is used, which means that if an atom (or the wall) moves such that an atom is outside the wall on a timestep by a distance  $\delta$  (e.g. due to *fix nve*), then it is put back inside the face by the same  $\delta$ , and the velocity relative to the moving wall is flipped in *z*. For instance, a stationary particle hit with a piston wall with velocity *vz*, will end the timestep with a velocity of  $2*vz$ .

Currently the *face* keyword can only be *zlo*. This creates a piston moving in the positive *z* direction. Particles with *z* coordinate less than the wall position are reflected to a *z* coordinate greater than the wall position. If the piston velocity is *vpz* and the particle velocity before reflection is *vzi*, the particle velocity after reflection is  $-vzi + 2*vpz$ .

The initial position of the wall can be specified by the *pos* keyword.

The final velocity of the wall can be specified by the *vel* keyword

The *ramp* keyword will cause the wall/piston to adjust the velocity linearly from zero velocity to *vel* over the course of the run. If the *ramp* keyword is omitted then the wall/piston moves at a constant velocity defined by *vel*.

The *temp* keyword will cause the region immediately in front of the wall/piston to be thermostatted with a Langevin thermostat. This region moves with the piston. The damping and kicking are measured in the reference frame of the piston. So, a temperature of zero would mean all particles were moving at exactly the speed of the wall/piston.

The *units* keyword determines the meaning of the distance units used to define a wall position, but only when a numeric constant is used.

A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacings.

---

#### 17.242.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

#### 17.242.5 Restrictions

This fix style is part of the SHOCK package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The face that has the wall/piston must be boundary type ‘s’ (shrink-wrapped). The opposing face can be any boundary type other than periodic.

A wall/piston should not be used with rigid bodies such as those defined by a “fix rigid” command. This is because the wall/piston displaces atoms directly rather than exerting a force on them.

#### 17.242.6 Related commands

*fix wall/reflect* command, *fix append/atoms* command

#### 17.242.7 Default

The keyword defaults are *pos* = 0, *vel* = 0, *units* = lattice.

## 17.243 fix wall/reflect command

Accelerator Variants: *wall/reflect/kk*

### 17.243.1 Syntax

```
fix ID group-ID wall/reflect face arg ... keyword value ...
```

- ID, group-ID are documented in *fix* command
- wall/reflect = style name of this fix command
- one or more face/arg pairs may be appended
- face = *xlo* or *xhi* or *ylo* or *yhi* or *zlo* or *zhi*  
 arg = EDGE or constant or variable  
 EDGE = current lo edge of simulation box  
 constant = number like 0.0 or 30.0 (distance units)  
 variable = *equal-style variable* like v\_x or v\_wiggle
- zero or more keyword/value pairs may be appended
- keyword = *units*  
*units* value = *lattice* or *box*  
*lattice* = the wall position is defined in lattice units  
*box* = the wall position is defined in simulation box units

### 17.243.2 Examples

```
fix xwalls all wall/reflect xlo EDGE xhi EDGE
fix walls all wall/reflect xlo 0.0 ylo 10.0 units box
fix top all wall/reflect zhi v_pressdown
```

### 17.243.3 Description

Bound the simulation with one or more walls which reflect particles in the specified group when they attempt to move through them.

Reflection means that if an atom moves outside the wall on a timestep by a distance delta (e.g. due to *fix nve*), then it is put back inside the face by the same delta, and the sign of the corresponding component of its velocity is flipped.

When used in conjunction with *fix nve* and *run\_style verlet*, the resultant time-integration algorithm is equivalent to the primitive splitting algorithm (PSA) described by *Bond*. Because each reflection event divides the corresponding timestep asymmetrically, energy conservation is only satisfied to  $O(dt)$ , rather than to  $O(dt^2)$  as it would be for velocity-Verlet integration without reflective walls.

Up to 6 walls or faces can be specified in a single command: *xlo*, *xhi*, *ylo*, *yhi*, *zlo*, *zhi*. A *lo* face reflects particles that move to a coordinate less than the wall position, back in the *hi* direction. A *hi* face reflects particles that move to a coordinate higher than the wall position, back in the *lo* direction.

The position of each wall can be specified in one of 3 ways: as the EDGE of the simulation box, as a constant value, or as a variable. If EDGE is used, then the corresponding boundary of the current simulation box is used. If a numeric constant is specified then the wall is placed at that position in the appropriate dimension (x, y, or z). In both the EDGE

and constant cases, the wall will never move. If the wall position is a variable, it should be specified as `v_name`, where name is an *equal-style variable* name. In this case the variable is evaluated each timestep and the result becomes the current position of the reflecting wall. Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent wall position.

The *units* keyword determines the meaning of the distance units used to define a wall position, but only when a numeric constant or variable is used. It is not relevant when EDGE is used to specify a face position. In the variable case, the variable is assumed to produce a value compatible with the *units* setting you specify.

A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacings.

---

Here are examples of variable definitions that move the wall position in a time-dependent fashion using equal-style variables.

```
variable ramp equal ramp(0,10)
fix 1 all wall/reflect xlo v_ramp

variable linear equal vdisplace(0,20)
fix 1 all wall/reflect xlo v_linear

variable wiggle equal swiggle(0.0,5.0,3.0)
fix 1 all wall/reflect xlo v_wiggle

variable wiggle equal cwiggle(0.0,5.0,3.0)
fix 1 all wall/reflect xlo v_wiggle
```

The *ramp(lo,hi)* function adjusts the wall position linearly from *lo* to *hi* over the course of a run. The *vdisplace(c0,velocity)* function does something similar using the equation  $position = c0 + velocity * delta$ , where *delta* is the elapsed time.

The *swiggle(c0,A,period)* function causes the wall position to oscillate sinusoidally according to this equation, where  $omega = 2 PI / period$ :

$$position = c0 + A \sin(omega * delta)$$

The *cwiggle(c0,A,period)* function causes the wall position to oscillate sinusoidally according to this equation, which will have an initial wall velocity of 0.0, and thus may impose a gentler perturbation on the particles:

$$position = c0 + A (1 - \cos(omega * delta))$$

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---



### 17.243.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix. No global or per-atom quantities are stored by this fix for access by various *output commands*. No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.243.5 Restrictions

Any dimension (xyz) that has a reflecting wall must be non-periodic.

A reflecting wall should not be used with rigid bodies such as those defined by a “fix rigid” command. This is because the wall/reflect displaces atoms directly rather than exerts a force on them. For rigid bodies, use a soft wall instead, such as *fix wall/lj93*. LAMMPS will flag the use of a rigid fix with fix wall/reflect with a warning, but will not generate an error.

### 17.243.6 Related commands

*fix wall/lj93*, *fix oneway*

### 17.243.7 Default

The default for the units keyword is lattice.

---

**(Bond)** Bond and Leimkuhler, SIAM J Sci Comput, 30, p 134 (2007).

## 17.244 fix wall/reflect/stochastic command

### 17.244.1 Syntax

```
fix ID group-ID wall/reflect/stochastic rstyle seed face args ... keyword value ...
```

- ID, group-ID are documented in *fix* command
- wall/reflect/stochastic = style name of this fix command
- rstyle = diffusive or maxwell or ccl
- seed = random seed for stochasticity (positive integer)
- one or more face/args pairs may be appended
- face = *xlo* or *xhi* or *ylo* or *yhi* or *zlo* or *zhi*

```
args = pos temp velx vely velz accomx accomy accomz
pos = EDGE or constant
EDGE = current lo or hi edge of simulation box
constant = number like 0.0 or 30.0 (distance units)
temp = wall temperature (temperature units)
velx,vely,velz = wall velocity in x,y,z directions (velocity units)
accomx,accomy,accomz = accommodation coeffs in x,y,z directions (unitless)
```

(continues on next page)

(continued from previous page)

```

not specified for rstyle = diffusive
single accom coeff specified for rstyle maxwell
all 3 coeffs specified for rstyle cll

```

- zero or more keyword/value pairs may be appended
- keyword = *units*

*units* value = *lattice* or *box*

*lattice* = the wall position is defined in lattice units

*box* = the wall position is defined in simulation box units

## 17.244.2 Examples

```

fix zwalls all wall/reflect/stochastic diffusive 23424 zlo EDGE 300 0.1 0.1 0 zhi EDGE_
↪200 0.1 0.1 0
fix ywalls all wall/reflect/stochastic maxwell 345533 ylo 5.0 300 0.1 0.0 0.0 0.8 yhi 10.
↪0 300 0.1 0.0 0.0 0.8
fix xwalls all wall/reflect/stochastic cercignanilampis 2308 xlo 0.0 300 0.0 0.1 0.9 0.8_
↪0.7 xhi EDGE 300 0.0 0.1 0 0.9 0.8 0.7 units box

```

## 17.244.3 Description

Bound the simulation with one or more walls which reflect particles in the specified group when they attempt to move through them.

Reflection means that if an atom moves outside the wall on a timestep (e.g. due to the *fix nve* command), then it is put back inside the wall with a changed velocity.

This fix models treats the wall as a moving solid boundary with a finite temperature, which can exchange energy with particles that collide with it. This is different than the simpler *fix wall/reflect* command which models mirror reflection. For this fix, the post collision velocity of each particle is treated stochastically. The randomness can come from many sources: thermal motion of the wall atoms, surface roughness, etc. Three stochastic reflection models are currently implemented.

For rstyle *diffusive*, particles are reflected diffusively. Their velocity distribution corresponds to an equilibrium distribution of particles at the wall temperature. No accommodation coefficients are specified.

For rstyle *maxwell*, particle reflection is Maxwellian which means partially diffusive and partially specular (*Maxwell*). A single accommodation coeff is specified which must be between 0.0 and 1.0 inclusive. It determines the fraction of the collision which is diffusive versus specular. An accommodation coefficient of 1.0 is fully diffusive; a coefficient of 0.0 is fully specular.

For rstyle *cll*, particle collisions are computed by the Cercignani/Lampis model. See *CL* and *To* for details. Three accommodations coefficient are specified. Each must be between 0.0 and 1.0 inclusive. Two are velocity accommodation coefficients; one is a normal kinetic energy accommodation. The normal coeff is the one corresponding to the normal of the wall itself. For example if the wall is *ylo* or *yhi*, *accomx* and *accomz* are the tangential velocity accommodation coefficients, and *accomy* is the normal kinetic energy accommodation coefficient.

The optional *units* keyword determines the distance units used to define a wall position. A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacings.

## 17.244.4 Restrictions

This fix has the same limitations as the *fix wall/reflect* command. Any dimension (xyz) that has a wall must be non-periodic. It should not be used with rigid bodies such as those defined by the *fix rigid* command. The wall velocity must lie on the same plane as the wall itself.

This fix is part of the EXTRA-FIX package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 17.244.5 Related commands

*fix wall/reflect*

## 17.244.6 Default

The default for the units keyword is lattice.

---

**(Maxwell)** J.C. Maxwell, Philos. Tans. Royal Soc. London, 157: 49-88 (1867).

**(Cercignani)** C. Cercignani and M. Lampis. Trans. Theory Stat. Phys. 1, 2, 101 (1971).

**(To)** Q.D. To, V.H. Vu, G. Lauriat, and C. Leonard. J. Math. Phys. 56, 103101 (2015).

## 17.245 fix wall/region command

### 17.245.1 Syntax

```
fix ID group-ID wall/region region-ID style args ... cutoff
```

- ID, group-ID are documented in *fix* command
- wall/region = style name of this fix command
- region-ID = region whose boundary will act as wall
- style = *lj93* or *lj126* or *lj1043* or *colloid* or *harmonic* or *morse*
- args for styles *lj93* or *lj126* or *lj1043* or *colloid* or *harmonic* =  
 epsilon = strength factor for wall-particle interaction (energy or energy/distance<sup>↔</sup>  
 ↔2 units)  
 sigma = size factor for wall-particle interaction (distance units)
- args for style *morse* =  
 D\_0 = depth of the potential (energy units)  
 alpha = width parameter (1/distance units)  
 r\_0 = distance of the potential minimum from wall position (distance units)
- cutoff = distance from wall at which wall-particle interaction is cut off (distance units)

## 17.245.2 Examples

```
fix wall all wall/region mySphere lj93 1.0 1.0 2.5
fix wall all wall/region mySphere harmonic 1.0 0.0 2.5
fix wall all wall/region box_top morse 1.0 1.0 1.5 3.0
```

## 17.245.3 Description

Treat the surface of the geometric region defined by the *region-ID* as a bounding wall which interacts with nearby particles according to the specified style.

The distance between a particle and the surface is the distance to the nearest point on the surface and the force the wall exerts on the particle is along the direction between that point and the particle, which is the direction normal to the surface at that point. Note that if the region surface is comprised of multiple “faces”, then each face can exert a force on the particle if it is close enough. E.g. for *region\_style block*, a particle in the interior, near a corner of the block, could feel wall forces from 1, 2, or 3 faces of the block.

Regions are defined using the *region* command. Note that the region volume can be interior or exterior to the bounding surface, which will determine in which direction the surface interacts with particles, i.e. the direction of the surface normal. The surface of the region only exerts forces on particles “inside” the region; if a particle is “outside” the region it will generate an error, because it has moved through the wall.

Regions can either be primitive shapes (block, sphere, cylinder, etc) or combinations of primitive shapes specified via the *union* or *intersect* region styles. These latter styles can be used to construct particle containers with complex shapes. Regions can also change over time via the *region* command keywords (move) and *rotate*. If such a region is used with this fix, then the of region surface will move over time in the corresponding manner.

---

**Note:** As discussed on the *region* command doc page, regions in LAMMPS do not get wrapped across periodic boundaries. It is up to you to insure that periodic or non-periodic boundaries are specified appropriately via the *boundary* command when using a region as a wall that bounds particle motion. This also means that if you embed a region in your simulation box and want it to repulse particles from its surface (using the “side out” option in the *region* command), that its repulsive force will not be felt across a periodic boundary.

---

---

**Note:** For primitive regions with sharp corners and/or edges (e.g. a block or cylinder), wall/particle forces are computed accurately for both interior and exterior regions. For *union* and *intersect* regions, additional sharp corners and edges may be present due to the intersection of the surfaces of 2 or more primitive volumes. These corners and edges can be of two types: concave or convex. Concave points/edges are like the corners of a cube as seen by particles in the interior of a cube. Wall/particle forces around these features are computed correctly. Convex points/edges are like the corners of a cube as seen by particles exterior to the cube, i.e. the points jut into the volume where particles are present. LAMMPS does NOT compute the location of these convex points directly, and hence wall/particle forces in the cutoff volume around these points suffer from inaccuracies. The basic problem is that the outward normal of the surface is not continuous at these points. This can cause particles to feel no force (they don’t “see” the wall) when in one location, then move a distance epsilon, and suddenly feel a large force because they now “see” the wall. In a worst-case scenario, this can blow particles out of the simulation box. Thus, as a general rule you should not use the *fix wall/gran/region* command with *union* or *intersect* regions that have convex points or edges resulting from the union/intersection (convex points/edges in the union/intersection due to a single sub-region are still OK).

---

---

**Note:** Similarly, you should not define *union* or *intersert* regions for use with this command that share an overlapping common face that is part of the overall outer boundary (interior boundary is OK), even if the face is smooth. E.g. two regions of style *block* in a *union* region, where the two blocks overlap on one or more of their faces. This is because

---

LAMMPS discards points that are part of multiple sub-regions when calculating wall/particle interactions, to avoid double-counting the interaction. Having two coincident faces could cause the face to become invisible to the particles. The solution is to make the two faces differ by epsilon in their position.

The energy of wall-particle interactions depends on the specified style.

For style *lj93*, the energy E is given by the 9/3 potential:

$$E = \epsilon \left[ \frac{2}{15} \left( \frac{\sigma}{r} \right)^9 - \left( \frac{\sigma}{r} \right)^3 \right] \quad r < r_c$$

For style *lj126*, the energy E is given by the 12/6 potential:

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_c$$

For style *wall/lj1043*, the energy E is given by the 10/4/3 potential:

$$E = 2\pi\epsilon \left[ \frac{2}{5} \left( \frac{\sigma}{r} \right)^{10} - \left( \frac{\sigma}{r} \right)^4 - \frac{\sqrt{2}\sigma^3}{3 \left( r + (0.61/\sqrt{2})\sigma \right)^3} \right] \quad r < r_c$$

For style *colloid*, the energy E is given by an integrated form of the *pair\_style colloid* potential:

$$E = \epsilon \left[ \frac{\sigma^6}{7560} \left( \frac{6R - D}{D^7} + \frac{D + 8R}{(D + 2R)^7} \right) - \frac{1}{6} \left( \frac{2R(D + R) + D(D + 2R) [\ln D - \ln(D + 2R)]}{D(D + 2R)} \right) \right] \quad r < r_c$$

For style *wall/harmonic*, the energy E is given by a harmonic spring potential (the distance parameter is ignored):

$$E = \epsilon (r - r_c)^2 \quad r < r_c$$

For style *wall/morse*, the energy E is given by the Morse potential:

$$E = D_0 \left[ e^{-2\alpha(r-r_0)} - 2e^{-\alpha(r-r_0)} \right] \quad r < r_c$$

Unlike other styles, this requires three parameters ( $D_0$ ,  $\alpha$ , and  $r_0$  in this order) instead of two like for the other wall styles.

In all cases,  $r$  is the distance from the particle to the region surface, and  $R_c$  is the *cutoff* distance at which the particle and surface no longer interact. The cutoff is always the last argument. The energy of the wall potential is shifted so that the wall-particle interaction energy is 0.0 at the cutoff distance.

For a full description of these wall styles, see `fix_style wall`

#### 17.245.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*.

The *fix\_modify energy* option is supported by this fix to add the energy of interaction between atoms and the region wall to the global potential energy of the system as part of *thermodynamic output*. The default setting for this fix is *fix\_modify energy no*.

The *fix\_modify virial* option is supported by this fix to add the contribution due to the interaction between atoms and the region wall to both the global pressure and per-atom stress of the system via the *compute pressure* and *compute*

*stress/atom* commands. The former can be accessed by *thermodynamic output*. The default setting for this fix is *fix\_modify virial no*.

The *fix\_modify respa* option is supported by this fix. This allows to set at which level of the *r-RESPA* integrator the fix is adding its forces. Default is the outermost level.

This fix computes a global scalar energy and a global 3-length vector of forces, which can be accessed by various *output commands*. The scalar energy is the sum of energy interactions for all particles interacting with the wall represented by the region surface. The 3 vector quantities are the x,y,z components of the total force acting on the wall due to the particles. The scalar and vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command.

The forces due to this fix are imposed during an energy minimization, invoked by the *minimize* command.

---

**Note:** If you want the atom/wall interaction energy to be included in the total potential energy of the system (the quantity being minimized), you MUST enable the *fix\_modify energy* option for this fix.

---

## 17.245.5 Restrictions

none

## 17.245.6 Related commands

*fix wall/lj93*, *fix wall/lj126*, *fix wall/lj1043*, *fix wall/colloid*, *fix wall/harmonic*, *fix wall/gran*

## 17.245.7 Default

none

# 17.246 fix wall/srd command

## 17.246.1 Syntax

<code>fix ID group-ID wall/srd face arg ... keyword value ...</code>
--

- ID, group-ID are documented in *fix* command
- wall/srd = style name of this fix command
- one or more face/arg pairs may be appended
- face = *xlo* or *xhi* or *ylo* or *yhi* or *zlo* or *zhi*

*xlo,ylo,zlo* arg = EDGE or constant or variable

EDGE = current lo edge of simulation box

constant = number like 0.0 or -30.0 (distance units)

variable = *equal-style variable* like v\_x or v\_wiggle

*xhi,yhi,zhi* arg = EDGE or constant or variable

EDGE = current hi edge of simulation box

constant = number like 50.0 or 100.3 (distance units)

variable = *equal-style variable* like v\_x or v\_wiggle

- zero or more keyword/value pairs may be appended
- keyword = *units*

```
units value = lattice or box
  lattice = the wall position is defined in lattice units
  box = the wall position is defined in simulation box units
```

## 17.246.2 Examples

```
fix xwalls all wall/srd xlo EDGE xhi EDGE
fix walls all wall/srd xlo 0.0 ylo 10.0 units box
fix top all wall/srd zhi v_pressdown
```

## 17.246.3 Description

Bound the simulation with one or more walls which interact with stochastic reaction dynamics (SRD) particles as slip (smooth) or no-slip (rough) flat surfaces. The wall interaction is actually invoked via the *fix srd* command, only on the group of SRD particles it defines, so the group setting for the *fix wall/srd* command is ignored.

A particle/wall collision occurs if an SRD particle moves outside the wall on a timestep. This alters the position and velocity of the SRD particle and imparts a force to the wall.

The *collision* and *Tsrd* settings specified via the *fix srd* command affect the SRD/wall collisions. A *slip* setting for the *collision* keyword means that the tangential component of the SRD particle momentum is preserved. Thus only a normal force is imparted to the wall. The normal component of the new SRD velocity is sampled from a Gaussian distribution at temperature *Tsrd*.

For a *noslip* setting of the *collision* keyword, both the normal and tangential components of the new SRD velocity are sampled from a Gaussian distribution at temperature *Tsrd*. Additionally, a new tangential direction for the SRD velocity is chosen randomly. This collision style imparts both a normal and tangential force to the wall.

Up to 6 walls or faces can be specified in a single command: *xlo, xhi, ylo, yhi, zlo, zhi*. A *lo* face reflects particles that move to a coordinate less than the wall position, back in the *hi* direction. A *hi* face reflects particles that move to a coordinate higher than the wall position, back in the *lo* direction.

The position of each wall can be specified in one of 3 ways: as the EDGE of the simulation box, as a constant value, or as a variable. If EDGE is used, then the corresponding boundary of the current simulation box is used. If a numeric constant is specified then the wall is placed at that position in the appropriate dimension (x, y, or z). In both the EDGE and constant cases, the wall will never move. If the wall position is a variable, it should be specified as *v\_name*, where name is an *equal-style variable* name. In this case the variable is evaluated each timestep and the result becomes the current position of the reflecting wall. Equal-style variables can specify formulas with various mathematical functions, and include *thermo\_style* command keywords for the simulation box parameters and timestep and elapsed time. Thus it is easy to specify a time-dependent wall position.

---

**Note:** Because the trajectory of the SRD particle is tracked as it collides with the wall, you must insure that *r* = distance of the particle from the wall, is always > 0 for SRD particles, or LAMMPS will generate an error. This means you cannot start your simulation with SRD particles at the wall position *coord* (*r* = 0) or with particles on the wrong side of the wall (*r* < 0).

---

**Note:** If you have 2 or more walls that come together at an edge or corner (e.g. walls in the x and y dimensions), then be sure to set the *overlap* keyword to *yes* in the *fix srd* command, since the walls effectively overlap when SRD particles

collide with them. LAMMPS will issue a warning if you do not do this.

---

**Note:** The walls of this fix only interact with SRD particles, as defined by the *fix srd* command. If you are simulating a mixture containing other kinds of particles, then you should typically use *another wall command* to act on the other particles. Since SRD particles will be colliding both with the walls and the other particles, it is important to insure that the other particle's finite extent does not overlap an SRD wall. If you do not do this, you may generate errors when SRD particles end up "inside" another particle or a wall at the beginning of a collision step.

---

The *units* keyword determines the meaning of the distance units used to define a wall position, but only when a numeric constant is used. It is not relevant when EDGE or a variable is used to specify a face position.

A *box* value selects standard distance units as defined by the *units* command, e.g. Angstroms for units = real or metal. A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacings.

---

Here are examples of variable definitions that move the wall position in a time-dependent fashion using equal-style *variables*.

```
variable ramp equal ramp(0,10)
fix 1 all wall/srd xlo v_ramp

variable linear equal vdisplace(0,20)
fix 1 all wall/srd xlo v_linear

variable wiggle equal swiggle(0.0,5.0,3.0)
fix 1 all wall/srd xlo v_wiggle

variable wiggle equal cwiggle(0.0,5.0,3.0)
fix 1 all wall/srd xlo v_wiggle
```

The *ramp(lo,hi)* function adjusts the wall position linearly from *lo* to *hi* over the course of a run. The *vdisplace(c0,velocity)* function does something similar using the equation  $position = c0 + velocity * delta$ , where *delta* is the elapsed time.

The *swiggle(c0,A,period)* function causes the wall position to oscillate sinusoidally according to this equation, where  $omega = 2 PI / period$ :

$$position = c0 + A \sin(omega * delta)$$

The *cwiggle(c0,A,period)* function causes the wall position to oscillate sinusoidally according to this equation, which will have an initial wall velocity of 0.0, and thus may impose a gentler perturbation on the particles:

$$position = c0 + A (1 - \cos(omega * delta))$$

---



## 17.246.4 Restart, fix\_modify, output, run start/stop, minimize info

No information about this fix is written to *binary restart files*. None of the *fix\_modify* options are relevant to this fix.

This fix computes a global array of values which can be accessed by various *output commands*. The number of rows in the array is equal to the number of walls defined by the fix. The number of columns is 3, for the x,y,z components of force on each wall.

Note that an outward normal force on a wall will be a negative value for *lo* walls and a positive value for *hi* walls. The array values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

## 17.246.5 Restrictions

Any dimension (xyz) that has an SRD wall must be non-periodic.

## 17.246.6 Related commands

*fix srd*

## 17.246.7 Default

none

## 17.247 fix widom command

### 17.247.1 Syntax

```
fix ID group-ID widom N M type seed T keyword values ...
```

- ID, group-ID are documented in *fix* command
- widom = style name of this fix command
- N = invoke this fix every N steps
- M = number of Widom insertions to attempt every N steps
- type = atom type for inserted atoms (must be 0 if mol keyword used)
- seed = random # seed (positive integer)
- T = temperature of the system (temperature units)
- zero or more keyword/value pairs may be appended to args

keyword = *mol*, *region*, *full\_energy*, *charge*, *intra\_energy*

*mol* value = template-ID

template-ID = ID of molecule template specified in a separate *molecule* command

*region* value = region-ID

region-ID = ID of region where Widom insertions are allowed

*full\_energy* = compute the entire system energy when performing Widom insertions

*charge* value = charge of inserted atoms (charge units)  
*intra\_energy* value = intramolecular energy (energy units)

## 17.247.2 Examples

```
fix 2 gas widom 1 50000 1 19494 2.0
fix 3 water widom 1000 100 0 29494 300.0 mol h2omol full_energy
```

## 17.247.3 Description

This fix performs Widom insertions of atoms or molecules at the given temperature as discussed in (*Frenkel*). Specific uses include computation of Henry constants of small molecules in microporous materials or amorphous systems.

Every N timesteps the fix attempts M number of Widom insertions of atoms or molecules.

If the *mol* keyword is used, only molecule insertions are performed. Conversely, if the *mol* keyword is not used, only atom insertions are performed.

This command may optionally use the *region* keyword to define an insertion volume. The specified region must have been previously defined with a *region* command. It must be defined with *side = in*. Insertion attempts occur only within the specified region. For non-rectangular regions, random trial points are generated within the rectangular bounding box until a point is found that lies inside the region. If no valid point is generated after 1000 trials, no insertion is performed. If an attempted insertion places the atom or molecule center-of-mass outside the specified region, a new attempted insertion is generated. This process is repeated until the atom or molecule center-of-mass is inside the specified region.

Note that neighbor lists are re-built every timestep that this fix is invoked, so you should not set N to be too small. See the *neighbor* command for details.

When an atom or molecule is to be inserted, its coordinates are chosen at a random position within the current simulation cell or region. Relative coordinates for atoms in a molecule are taken from the template molecule provided by the user. The center of mass of the molecule is placed at the insertion point. The orientation of the molecule is chosen at random by rotating about this point.

Individual atoms are inserted, unless the *mol* keyword is used. It specifies a *template-ID* previously defined using the *molecule* command, which reads a file that defines the molecule. The coordinates, atom types, charges, etc., as well as any bonding and special neighbor information for the molecule can be specified in the molecule file. See the *molecule* command for details. The only settings required to be in this file are the coordinates and types of atoms in the molecule.

Note that fix widom does not use configurational bias MC or any other kind of sampling of intramolecular degrees of freedom. Inserted molecules can have different orientations, but they will all have the same intramolecular configuration, which was specified in the molecule command input.

For atoms, inserted particles have the specified atom type. For molecules, they use the same atom types as in the template molecule supplied by the user.

The excess chemical potential  $\mu_{ex}$  is defined as:

$$\mu_{ex} = -kT \ln(\langle \exp(-(U_{N+1} - U_N)/k_B T) \rangle)$$

where  $k_B$  is the Boltzmann constant,  $T$  is the user-specified temperature,  $U_N$  and  $U_{N+1}$  is the potential energy of the system with  $N$  and  $N + 1$  particles.

The *full\_energy* option means that the fix calculates the total potential energy of the entire simulated system, instead of just the energy of the part that is changed. By default, this option is off, in which case only partial energies are computed to determine the energy difference due to the proposed change.

The *full\_energy* option is needed for systems with complicated potential energy calculations, including the following:

- long-range electrostatics (kspace)
- many-body pair styles
- hybrid pair styles
- eam pair styles
- tail corrections
- need to include potential energy contributions from other fixes

In these cases, LAMMPS will automatically apply the *full\_energy* keyword and issue a warning message.

When the *mol* keyword is used, the *full\_energy* option also includes the intramolecular energy of inserted and deleted molecules, whereas this energy is not included when *full\_energy* is not used. If this is not desired, the *intra\_energy* keyword can be used to define an amount of energy that is subtracted from the final energy when a molecule is inserted, and subtracted from the initial energy when a molecule is deleted. For molecules that have a non-zero intramolecular energy, this will ensure roughly the same behavior whether or not the *full\_energy* option is used.

Some fixes have an associated potential energy. Examples of such fixes include: *efield*, *gravity*, *addforce*, *restrain*, and *wall fixes*. For that energy to be included in the total potential energy of the system (the quantity used when performing Widom insertions), you MUST enable the *fix\_modify energy* option for that fix. The doc pages for individual *fix* commands specify if this should be done.

Use the *charge* option to insert atoms with a user-specified point charge. Note that doing so will cause the system to become non-neutral. LAMMPS issues a warning when using long-range electrostatics (kspace) with non-neutral systems. See the *compute group/group* documentation for more details about simulating non-neutral systems with kspace on.

#### 17.247.4 Restart, fix\_modify, output, run start/stop, minimize info

This fix writes the state of the fix to *binary restart files*. This includes information about the random number generator seed, the next timestep for Widom insertions etc. See the *read\_restart* command for info on how to re-specify a fix in an input script that reads a restart file, so that the operation of the fix continues in an uninterrupted fashion.

---

**Note:** For this to work correctly, the timestep must **not** be changed after reading the restart with *reset\_timestep*. The fix will try to detect it and stop with an error.

---

None of the *fix\_modify* options are relevant to this fix.

This fix computes a global vector of length 3, which can be accessed by various *output commands*. The vector values are the following global cumulative quantities:

- 1 = average excess chemical potential on each timestep
- 2 = average difference in potential energy on each timestep
- 3 = volume of the insertion region

The vector values calculated by this fix are “extensive”.

No parameter of this fix can be used with the *start/stop* keywords of the *run* command. This fix is not invoked during *energy minimization*.

### 17.247.5 Restrictions

This fix is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

Do not set “neigh\_modify once yes” or else this fix will never be called. RENEIGHBORING is **required**.

Can be run in parallel, but aspects of the GCMC part will not scale well in parallel. Only usable for 3D simulations.

### 17.247.6 Related commands

*fix gcmc fix atom/swap, neighbor, fix deposit, fix evaporate,*

### 17.247.7 Default

The option defaults are mol = no, intra\_energy = 0.0 and full\_energy = no, except for the situations where full\_energy is required, as listed above.

---

**(Frenkel)** Frenkel and Smit, Understanding Molecular Simulation, Academic Press, London, 2002.

## 18.1 compute ackland/atom command

### 18.1.1 Syntax

```
compute ID group-ID ackland/atom keyword/value
```

- ID, group-ID are documented in *compute* command
- ackland/atom = style name of this compute command
- zero or more keyword/value pairs may be appended
- keyword = *legacy*  
*legacy* args = *yes* or *no* = use (*yes*) or do not use (*no*) legacy Ackland algorithm.  
→ implementation

### 18.1.2 Examples

```
compute 1 all ackland/atom  
compute 1 all ackland/atom legacy yes
```

### 18.1.3 Description

Defines a computation that calculates the local lattice structure according to the formulation given in (*Ackland*). Historically, LAMMPS had two, slightly different implementations of the algorithm from the paper. With the *legacy* keyword, it is possible to switch between the pre-2015 (*legacy yes*) and post-2015 implementation (*legacy no*). The post-2015 variant is the default.

In contrast to the *centro-symmetry parameter* this method is stable against temperature boost, because it is based not on the distance between particles but the angles. Therefore statistical fluctuations are averaged out a little more. A comparison with the Common Neighbor Analysis metric is made in the paper.

The result is a number which is mapped to the following different lattice structures:

- 0 = UNKNOWN
- 1 = BCC
- 2 = FCC
- 3 = HCP

- 4 = ICO

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (i.e. each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently or to have multiple compute/dump commands, each of which computes this quantity.-

### 18.1.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

### 18.1.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The per-atom vector values will be unitless since they are the integers defined above.

### 18.1.6 Related commands

*compute centro/atom*

### 18.1.7 Default

The keyword *legacy* defaults to *no*.

---

(Ackland) Ackland, Jones, Phys Rev B, 73, 054104 (2006).

## 18.2 compute adf command

### 18.2.1 Syntax

```
compute ID group-ID adf Nbin itype1 jtype1 ktype1 Rjinner1 Rjouter1 Rkinner1 Rkouter1 ...
```

- ID, group-ID are documented in *compute* command
- adf = style name of this compute command
- Nbin = number of ADF bins
- itypeN = central atom type for Nth ADF histogram (see asterisk form below)
- jtypeN = J atom type for Nth ADF histogram (see asterisk form below)
- ktypeN = K atom type for Nth ADF histogram (see asterisk form below)
- RjinnerN = inner radius of J atom shell for Nth ADF histogram (distance units)
- RjouterN = outer radius of J atom shell for Nth ADF histogram (distance units)
- RkinnerN = inner radius of K atom shell for Nth ADF histogram (distance units)
- RkouterN = outer radius of K atom shell for Nth ADF histogram (distance units)

- zero or one keyword/value pairs may be appended
- keyword = *ordinate*

*ordinate* value = *degree* or *radian* or *cosine*

Choose the ordinate parameter for the histogram

## 18.2.2 Examples

```
compute 1 fluid adf 32 1 1 1 0.0 1.2 0.0 1.2 &
                    1 1 2 0.0 1.2 0.0 1.5 &
                    1 2 2 0.0 1.5 0.0 1.5 &
                    2 1 1 0.0 1.2 0.0 1.2 &
                    2 1 2 0.0 1.5 2.0 3.5 &
                    2 2 2 2.0 3.5 2.0 3.5
compute 1 fluid adf 32 1*2 1*2 1*2 0.5 3.5
compute 1 fluid adf 32
```

## 18.2.3 Description

Define a computation that calculates one or more angular distribution functions (ADF) for a group of particles. Each ADF is calculated in histogram form by measuring the angle formed by a central atom and two neighbor atoms and binning these angles into *Nbin* bins. Only neighbors for which  $Rinner < R < Router$  are counted, where *Rinner* and *Router* are specified separately for the first and second neighbor atom in each requested ADF.

---

**Note:** If you have a bonded system, then the settings of *special\_bonds* command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the *special\_bonds* command, and means those pairwise interactions do not appear in the neighbor list. Because this fix uses a neighbor list, it also means those pairs will not be included in the ADF. This does not apply when using long-range coulomb interactions (*coul/long*, *coul/msm*, *coul/wolf* or similar. One way to get around this would be to set *special\_bond* scaling factors to very tiny numbers that are not exactly zero (e.g. 1.0e-50). Another workaround is to write a dump file, and use the *rerun* command to compute the ADF for snapshots in the dump file. The rerun script can use a *special\_bonds* command that includes all pairs in the neighbor list.

---

**Note:** If you request any outer cutoff  $Router > force$  cutoff, or if no pair style is defined, e.g. the *rerun* command is being used to post-process a dump file of snapshots you must insure ghost atom information out to the largest value of  $Router + skin$  is communicated, via the *comm\_modify cutoff* command, else the ADF computation cannot be performed, and LAMMPS will give an error message. The *skin* value is what is specified with the *neighbor* command.

---

The *itypeN*, *jtypeN*, *ktypeN* settings can be specified in one of two ways. An explicit numeric value can be used, as in the first example above. Or a wild-card asterisk can be used to specify a range of atom types as in the second example above. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If N = the number of atom types, then an asterisk with no numeric values means all types from 1 to N. A leading asterisk means all types from 1 to n (inclusive). A trailing asterisk means all types from n to N (inclusive). A middle asterisk means all types from m to n (inclusive).

If *itypeN*, *jtypeN*, and *ktypeN* are single values, as in the first example above, this means that the ADF is computed where atoms of type *itypeN* are the central atom, and neighbor atoms of type *jtypeN* and *ktypeN* are forming the angle. If any of *itypeN*, *jtypeN*, or *ktypeN* represent a range of values via the wild-card asterisk, as in the second example above, this means that the ADF is computed where atoms of any of the range of types represented by *itypeN* are the

central atom, and the angle is formed by two neighbors, one neighbor in the range of types represented by *jtypeN* and another neighbor in the range of types represented by *ktypeN*.

If no *itypeN*, *jtypeN*, *ktypeN* settings are specified, then LAMMPS will generate a single ADF for all atoms in the group. The inner cutoff is set to zero and the outer cutoff is set to the force cutoff. If no *pair\_style* is specified, there is no force cutoff and LAMMPS will give an error message. Note that in most cases, generating an ADF for all atoms is not a good thing. Such an ADF is both uninformative and extremely expensive to compute. For example, with liquid water with a 10 Å force cutoff, there are 80,000 angles per atom. In addition, most of the interesting angular structure occurs for neighbors that are the closest to the central atom, involving just a few dozen angles.

Angles for each ADF are generated by double-looping over the list of neighbors of each central atom I, just as they would be in the force calculation for a three-body potential such as *Stillinger-Weber*. The angle formed by central atom I and neighbor atoms J and K is included in an ADF if the following criteria are met:

- atoms I,J,K are all in the specified compute group
- the distance between atoms I,J is between *Rjinner* and *Rjouter*
- the distance between atoms I,K is between *Rkinner* and *Rkouter*
- the type of the I atom matches *itypeN* (one or a range of types)
- atoms I,J,K are distinct
- the type of the J atom matches *jtypeN* (one or a range of types)
- the type of the K atom matches *ktypeN* (one or a range of types)

Each unique angle satisfying the above criteria is counted only once, regardless of whether either or both of the neighbor atoms making up the angle appear in both the J and K lists. It is OK if a particular angle is included in more than one individual histogram, due to the way the *itypeN*, *jtypeN*, *ktypeN* arguments are specified.

The first ADF value for a bin is calculated from the histogram count by dividing by the total number of triples satisfying the criteria, so that the integral of the ADF w.r.t. angle is 1, i.e. the ADF is a probability density function.

The second ADF value is reported as a cumulative sum of all bins up to the current bins, averaged over atoms of type *itypeN*. It represents the number of angles per central atom with angle less than or equal to the angle of the current bin, analogous to the coordination number radial distribution function.

The *ordinate* optional keyword determines whether the bins are of uniform angular size from zero to 180 (*degree*), zero to Pi (*radian*), or the cosine of the angle uniform in the range [-1,1] (*cosine*). *cosine* has the advantage of eliminating the *acos()* function call, which speeds up the compute by 2-3x, and it is also preferred on physical grounds, because for uniformly distributed particles in 3D, the angular probability density w.r.t *dtheta* is  $\sin(\theta)/2$ , while for  $d(\cos(\theta))$ , it is  $1/2$ . Regardless of which ordinate is chosen, the first column of ADF values is normalized w.r.t. the range of that ordinate, so that the integral is 1.

The simplest way to output the results of the compute adf calculation to a file is to use the *fix ave/time* command, for example:

```
compute myADF all adf 32 2 2 2 0.5 3.5 0.5 3.5
fix 1 all ave/time 100 1 100 c_myADF[*] file tmp.adf mode vector
```



## 18.2.4 Output info

This compute calculates a global array with the number of rows =  $Nbins$  and the number of columns =  $1 + 2 \times Ntriples$ , where  $Ntriples$  is the number of I,J,K triples specified. The first column has the bin coordinate (angle-related ordinate at midpoint of bin). Each subsequent column has the two ADF values for a specific set of  $(itypeN, jtypeN, ktypeN)$  interactions, as described above. These values can be used by any command that uses a global values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The array values calculated by this compute are all “intensive”.

The first column of array values is the angle-related ordinate, either the angle in degrees or radians, or the cosine of the angle. Each subsequent pair of columns gives the first and second kinds of ADF for a specific set of  $(itypeN, jtypeN, ktypeN)$ . The values in the first ADF column are normalized numbers  $\geq 0.0$ , whose integral w.r.t. the ordinate is 1, i.e. the first ADF is a normalized probability distribution. The values in the second ADF column are also numbers  $\geq 0.0$ . They are the cumulative density distribution of angles per atom. By definition, this ADF is monotonically increasing from zero to a maximum value equal to the average total number of angles per atom satisfying the ADF criteria.

## 18.2.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

The ADF is not computed for neighbors outside the force cutoff, since processors (in parallel) don’t know about atom coordinates for atoms further away than that distance. If you want an ADF for larger distances, you can use the [rerun](#) command to post-process a dump file and set the cutoff for the potential to be longer in the rerun script. Note that in the rerun context, the force cutoff is arbitrary, since you are not running dynamics and thus are not changing your model.

## 18.2.6 Related commands

*compute rdf, fix ave/time, compute\_modify*

## 18.2.7 Default

The keyword default is ordinate = degree.

# 18.3 compute angle command

## 18.3.1 Syntax

```
compute ID group-ID angle
```

- ID, group-ID are documented in [compute](#) command
- angle = style name of this compute command

## 18.3.2 Examples

```
compute 1 all angle
```

## 18.3.3 Description

Define a computation that extracts the angle energy calculated by each of the angle sub-styles used in the *angle\_style hybrid* command. These values are made accessible for output or further processing by other commands. The group specified for this command is ignored.

This compute is useful when using *angle\_style hybrid* if you want to know the portion of the total energy contributed by one or more of the hybrid sub-styles.

## 18.3.4 Output info

This compute calculates a global vector of length  $N$ , where  $N$  is the number of sub\_styles defined by the *angle\_style hybrid* command, which can be accessed by indices 1 through  $N$ . These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector values are “extensive” and will be in energy *units*.

## 18.3.5 Restrictions

none

## 18.3.6 Related commands

*compute pe*, *compute pair*

## 18.3.7 Default

none

# 18.4 compute angle/local command

## 18.4.1 Syntax

```
compute ID group-ID angle/local value1 value2 ... keyword args ...
```

- ID, group-ID are documented in *compute* command
- angle/local = style name of this compute command
- one or more values may be appended
- value = *theta* or *eng* or *v\_name*

```

theta = tabulate angles
eng = tabulate angle energies
v_name = equal-style variable with name (see below)

```

- zero or more keyword/args pairs may be appended
- keyword = *set*

```

set args = theta name
  theta = only currently allowed arg
  name = name of variable to set with theta

```

## 18.4.2 Examples

```

compute 1 all angle/local theta
compute 1 all angle/local eng theta
compute 1 all angle/local theta v_cos set theta t

```

## 18.4.3 Description

Define a computation that calculates properties of individual angle interactions. The number of datums generated, aggregated across all processors, equals the number of angles in the system, modified by the group parameter as explained below.

The value *theta* is the angle for the three atoms in the interaction.

The value *eng* is the interaction energy for the angle.

The value *v\_name* can be used together with the *set* keyword to compute a user-specified function of the angle *theta*. The *name* specified for the *v\_name* value is the name of an *equal-style variable* which should evaluate a formula based on a variable which will store the angle *theta*. This other variable must be an *internal-style variable* defined in the input script; its initial numeric value can be anything. It must be an internal-style variable, because this command resets its value directly. The *set* keyword is used to identify the name of this other variable associated with *theta*.

Note that the value of *theta* for each angle which stored in the internal variable is in radians, not degrees.

As an example, these commands can be added to the bench/in.rhodo script to compute the cosine and cosine-squared of every angle in the system and output the statistics in various ways:

```

variable t internal 0.0
variable cos equal cos(v_t)
variable cossq equal cos(v_t)*cos(v_t)

compute 1 all property/local aatom1 aatom2 aatom3 atype
compute 2 all angle/local eng theta v_cos v_cossq set theta t
dump 1 all local 100 tmp.dump c_1[*] c_2[*]

compute 3 all reduce ave c_2[*]
thermo_style custom step temp press c_3[*]

fix 10 all ave/histo 10 10 100 -1 1 20 c_2[3] mode vector file tmp.histo

```

The *dump local* command will output the potential energy ( $\phi$ ), the angle ( $\theta$ ),  $\cos(\theta)$ , and  $\cos^2(\theta)$  for every angle  $\theta$  in the system. The *thermo\_style* command will print the average of those quantities via the *compute reduce* command with thermo output. And the *fix ave/histo* command will histogram the  $\cos(\theta)$  values and write them to a file.

The local data stored by this command is generated by looping over all the atoms owned on a processor and their angles. An angle will only be included if all three atoms in the angle are in the specified compute group. Any angles that have been broken (see the *angle\_style* command) by setting their angle type to 0 are not included. Angles that have been turned off (see the *fix shake* or *delete\_bonds* commands) by setting their angle type negative are written into the file, but their energy will be 0.0.

Note that as atoms migrate from processor to processor, there will be no consistent ordering of the entries within the local vector or array from one timestep to the next. The only consistency that is guaranteed is that the ordering on a particular timestep will be the same for local vectors or arrays generated by other compute commands. For example, angle output from the *compute property/local* command can be combined with data from this command and output by the *dump local* command in a consistent way.

Here is an example of how to do this:

```
compute 1 all property/local atype aatom1 aatom2 aatom3
compute 2 all angle/local theta eng
dump 1 all local 1000 tmp.dump index c_1[1] c_1[2] c_1[3] c_1[4] c_2[1] c_2[2]
```

#### 18.4.4 Output info

This compute calculates a local vector or local array depending on the number of values. The length of the vector or number of rows in the array is the number of angles. If a single value is specified, a local vector is produced. If two or more values are specified, a local array is produced where the number of columns = the number of values. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The output for *theta* will be in degrees. The output for *eng* will be in energy *units*.

#### 18.4.5 Restrictions

none

#### 18.4.6 Related commands

*dump local*, *compute property/local*

#### 18.4.7 Default

none

## 18.5 compute angmom/chunk command

### 18.5.1 Syntax

```
compute ID group-ID angmom/chunk chunkID
```

- ID, group-ID are documented in *compute* command
- angmom/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command

### 18.5.2 Examples

```
compute 1 fluid angmom/chunk molchunk
```

### 18.5.3 Description

Define a computation that calculates the angular momentum of multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

This compute calculates the 3 components of the angular momentum vector for each chunk, due to the velocity/momentum of the individual atoms in the chunk around the center-of-mass of the chunk. The calculation includes all effects due to atoms passing through periodic boundaries.

Note that only atoms in the specified group contribute to the calculation. The *compute chunk/atom* command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

---

**Note:** The coordinates of an atom contribute to the chunk’s angular momentum in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g. to 0) before invoking this compute by using the *set image* command.

---

The simplest way to output the results of the compute angmom/chunk calculation to a file is to use the *fix ave/time* command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk all angmom/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk[*] file tmp.out mode vector
```

## 18.5.4 Output info

This compute calculates a global array where the number of rows = the number of chunks *Nchunk* as calculated by the specified *compute chunk/atom* command. The number of columns = 3 for the three (*x*, *y*, *z*) components of the angular momentum for each chunk. These values can be accessed by any command that uses global array values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The array values are “intensive”. The array values will be in mass-velocity-distance *units*.

## 18.5.5 Restrictions

none

## 18.5.6 Related commands

*variable angmom()* function

## 18.5.7 Default

none

# 18.6 compute ave/sphere/atom command

Accelerator Variants: *ave/sphere/atom/kk*

## 18.6.1 Syntax

```
compute ID group-ID ave/sphere/atom keyword values ...
```

- ID, group-ID are documented in *compute* command
- ave/sphere/atom = style name of this compute command
- one or more keyword/value pairs may be appended

```
keyword = cutoff  
cutoff value = distance cutoff
```

## 18.6.2 Examples

```
compute 1 all ave/sphere/atom  
  
compute 1 all ave/sphere/atom cutoff 5.0  
comm_modify cutoff 5.0
```

### 18.6.3 Description

Define a computation that calculates the local mass density and temperature for each atom based on its neighbors inside a spherical cutoff. If an atom has  $M$  neighbors, then its local mass density is calculated as the sum of its mass and its  $M$  neighbor masses, divided by the volume of the cutoff sphere (or circle in 2d). The local temperature of the atom is calculated as the temperature of the collection of  $M + 1$  atoms, after subtracting the center-of-mass velocity of the  $M + 1$  atoms from each of the  $M + 1$  atom's velocities. This is effectively the thermal velocity of the neighborhood of the central atom, similar to *compute temp/com*.

The optional keyword *cutoff* defines the distance cutoff used when searching for neighbors. The default value is the cutoff specified by the pair style. If no pair style is defined, then a cutoff must be defined using this keyword. If the specified cutoff is larger than that of the pair\_style plus neighbor skin (or no pair style is defined), the *comm\_modify cutoff* option must also be set to match that of the *cutoff* keyword.

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (i.e. each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently.

---

**Note:** If you have a bonded system, then the settings of *special\_bonds* command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the *special\_bonds* command, and means those pairwise interactions do not appear in the neighbor list. Because this compute uses the neighbor list, it also means those pairs will not be included in the order parameter. This difficulty can be circumvented by writing a dump file, and using the *rerun* command to compute the order parameter for snapshots in the dump file. The rerun script can use a *special\_bonds* command that includes all pairs in the neighbor list.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 18.6.4 Output info

This compute calculates a per-atom array with two columns: mass density in density *units* and temperature in temperature *units*.

These values can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

### 18.6.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 18.6.6 Related commands

*comm\_modify*

### 18.6.7 Default

The option defaults are *cutoff* = pair style cutoff.

## 18.7 compute basal/atom command

### 18.7.1 Syntax

```
compute ID group-ID basal/atom
```

- ID, group-ID are documented in *compute* command
- basal/atom = style name of this compute command

### 18.7.2 Examples

```
compute 1 all basal/atom
```

### 18.7.3 Description

Defines a computation that calculates the hexagonal close-packed “c” lattice vector for each atom in the group. It does this by calculating the normal unit vector to the basal plane for each atom. The results enable efficient identification and characterization of twins and grains in hexagonal close-packed structures.

The output of the compute is thus the 3 components of a unit vector associated with each atom. The components are set to 0.0 for atoms not in the group.

Details of the calculation are given in (*Barrett*).

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (i.e. each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently or to have multiple compute/dump commands, each of which computes this quantity.

An example input script that uses this compute is provided in examples/PACKAGES/basal.



## 18.7.4 Output info

This compute calculates a per-atom array with three columns, which can be accessed by indices 1–3 by any command that uses per-atom values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The per-atom vector values are unitless since the three columns represent components of a unit vector.

## 18.7.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The output of this compute will be meaningless unless the atoms are on (or near) hcp lattice sites, since the calculation assumes a well-defined basal plane.

## 18.7.6 Related commands

*compute centro/atom*, *compute ackland/atom*

## 18.7.7 Default

none

---

(Barrett) Barrett, Tschopp, El Kadiri, Scripta Mat. 66, p.666 (2012).

# 18.8 compute body/local command

## 18.8.1 Syntax

```
compute ID group-ID body/local input1 input2 ...
```

- ID, group-ID are documented in *compute* command
- body/local = style name of this compute command
- one or more keywords may be appended
- keyword = *id* or *type* or *integer*
  - id* = atom ID of the body particle
  - type* = atom type of the body particle
  - integer* = 1,2,3,etc = index of fields defined by body style

## 18.8.2 Examples

```
compute 1 all body/local type 1 2 3
compute 1 all body/local 3 6
```

## 18.8.3 Description

Define a computation that calculates properties of individual body sub-particles. The number of data generated, aggregated across all processors, equals the number of body sub-particles plus the number of non-body particles in the system, modified by the `group` parameter as explained below. See the [Howto body](#) page for more details on using body particles.

The local data stored by this command is generated by looping over all the atoms. An atom will only be included if it is in the group. If the atom is a body particle, then its  $N$  sub-particles will be looped over, and it will contribute  $N$  data to the count of data. If it is not a body particle, it will contribute 1 datum.

For both body particles and non-body particles, the `id` keyword will store the ID of the particle.

For both body particles and non-body particles, the `type` keyword will store the type of the particle.

The *integer* keywords mean different things for body and non-body particles. If the atom is not a body particle, only its  $x$ ,  $y$ ,  $z$  coordinates can be referenced, using the *integer* keywords 1,2,3. Note that this means that if you want to access more fields than this for body particles, then you cannot include non-body particles in the group.

For a body particle, the *integer* keywords refer to fields calculated by the body style for each sub-particle. The body style, as specified by the *atom\_style body*, determines how many fields exist and what they are. See the [Howto\\_body](#) doc page for details of the different styles.

Here is an example of how to output body information using the *dump local* command with this compute. If fields 1, 2, and 3 for the body sub-particles are  $(x, y, z)$  coordinates, then the dump file will be formatted similar to the output of a *dump atom or custom* command.

```
compute 1 all body/local type 1 2 3
dump 1 all local 1000 tmp.dump index c_1[1] c_1[2] c_1[3] c_1[4]
```

## 18.8.4 Output info

This compute calculates a local vector or local array depending on the number of keywords. The length of the vector or number of rows in the array is the number of data as described above. If a single keyword is specified, a local vector is produced. If two or more keywords are specified, a local array is produced where the number of columns = the number of keywords. The vector or array can be accessed by any command that uses local values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The *units* for output values depend on the body style.

## 18.8.5 Restrictions

none

## 18.8.6 Related commands

*dump local*

## 18.8.7 Default

none

# 18.9 compute bond command

## 18.9.1 Syntax

```
compute ID group-ID bond
```

- ID, group-ID are documented in *compute* command
- bond = style name of this compute command

## 18.9.2 Examples

```
compute 1 all bond
```

## 18.9.3 Description

Define a computation that extracts the bond energy calculated by each of the bond sub-styles used in the *bond\_style hybrid* command. These values are made accessible for output or further processing by other commands. The group specified for this command is ignored.

This compute is useful when using *bond\_style hybrid* if you want to know the portion of the total energy contributed by one or more of the hybrid sub-styles.

## 18.9.4 Output info

This compute calculates a global vector of length  $N$ , where  $N$  is the number of sub\_styles defined by the *bond\_style hybrid* command, which can be accessed by indices 1 through  $N$ . These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector values are “extensive” and will be in energy *units*.

## 18.9.5 Restrictions

none

## 18.9.6 Related commands

*compute pe, compute pair*

## 18.9.7 Default

none

# 18.10 compute bond/local command

## 18.10.1 Syntax

```
compute ID group-ID bond/local value1 value2 ... keyword args ...
```

- ID, group-ID are documented in *compute* command
- bond/local = style name of this compute command
- one or more values may be appended
- value = *dist* or *dx* or *dy* or *dz* or *engpot* or *force* or *fx* or *fy* or *fz* or *engvib* or *engrot* or *engtrans* or *omega* or *velvib* or *v\_name*

*dist* = bond distance

*engpot* = bond potential energy

*force* = bond force

*dx, dy, dz* = components of pairwise distance

*fx, fy, fz* = components of bond force

*engvib* = bond kinetic energy of vibration

*engrot* = bond kinetic energy of rotation

*engtrans* = bond kinetic energy of translation

*omega* = magnitude of bond angular velocity

*velvib* = vibrational velocity along the bond length

*v\_name* = equal-style variable with name (see below)

- zero or more keyword/args pairs may be appended
- keyword = *set*

*set args* = *dist* name

*dist* = only currently allowed arg

name = name of variable to set with distance (*dist*)

## 18.10.2 Examples

```
compute 1 all bond/local engpot
compute 1 all bond/local dist engpot force

compute 1 all bond/local dist fx fy fz

compute 1 all bond/local dist v_distsq set dist d
```

## 18.10.3 Description

Define a computation that calculates properties of individual bond interactions. The number of datums generated, aggregated across all processors, equals the number of bonds in the system, modified by the group parameter as explained below.

All these properties are computed for the pair of atoms in a bond, whether the 2 atoms represent a simple diatomic molecule, or are part of some larger molecule.

The value *dist* is the current length of the bond. The values *dx*, *dy*, and *dz* are the xyz components of the *distance* between the pair of atoms. This value is always the distance from the atom of lower to the one with the higher id.

The value *engpot* is the potential energy for the bond, based on the current separation of the pair of atoms in the bond.

The value *force* is the magnitude of the force acting between the pair of atoms in the bond.

The values *fx*, *fy*, and *fz* are the xyz components of *force* between the pair of atoms in the bond.

The remaining properties are all computed for motion of the two atoms relative to the center of mass (COM) velocity of the 2 atoms in the bond.

The value *engvib* is the vibrational kinetic energy of the two atoms in the bond, which is simply  $\frac{1}{2}m_1v_1^2 + \frac{1}{2}m_2v_2^2$ , where  $v_1$  and  $v_2$  are the magnitude of the velocity of the two atoms along the bond direction, after the COM velocity has been subtracted from each.

The value *engrot* is the rotational kinetic energy of the two atoms in the bond, which is simply  $\frac{1}{2}m_1v_1^2 + \frac{1}{2}m_2v_2^2$ , where  $v_1$  and  $v_2$  are the magnitude of the velocity of the two atoms perpendicular to the bond direction, after the COM velocity has been subtracted from each.

The value *engtrans* is the translational kinetic energy associated with the motion of the COM of the system itself, namely  $\frac{1}{2}(m_1 + m_2)V_{cm}^2$ , where  $V_{cm}$  = magnitude of the velocity of the COM.

Note that these three kinetic energy terms are simply a partitioning of the summed kinetic energy of the two atoms themselves. That is, the total kinetic energy is  $\frac{1}{2}m_1v_1^2 + \frac{1}{2}m_2v_2^2 = \text{engvib} + \text{engrot} + \text{engtrans}$ , where  $v_1$  and  $v_2$  are the magnitude of the velocities of the two atoms, without any adjustment for the COM velocity.

The value *omega* is the magnitude of the angular velocity of the two atoms around their COM position.

The value *velvib* is the magnitude of the relative velocity of the two atoms in the bond towards each other. A negative value means the two atoms are moving toward each other; a positive value means they are moving apart.

The value *v\_name* can be used together with the *set* keyword to compute a user-specified function of the bond distance. The *name* specified for the *v\_name* value is the name of an *equal-style variable* which should evaluate a formula based on a variable which will store the bond distance. This other variable must be an *internal-style variable* defined in the input script; its initial numeric value can be anything. It must be an internal-style variable, because this command resets its value directly. The *set* keyword is used to identify the name of this other variable associated with theta.

As an example, these commands can be added to the bench/in.rhodo script to compute the length<sup>2</sup> of every bond in the system and output the statistics in various ways:

```
variable d internal 0.0
variable dsq equal v_d*v_d

compute 1 all property/local batom1 batom2 btype
compute 2 all bond/local engpot dist v_dsq set dist d
dump 1 all local 100 tmp.dump c_1[*] c_2[*]

compute 3 all reduce ave c_2[*]
thermo_style custom step temp press c_3[*]

fix 10 all ave/histo 10 10 100 0 6 20 c_2[3] mode vector file tmp.histo
```

The *dump local* command will output the energy, length, and length<sup>2</sup> for every bond in the system. The *thermo\_style* command will print the average of those quantities via the *compute reduce* command with thermo output, and the *fix ave/histo* command will histogram the length<sup>2</sup> values and write them to a file.

---

The local data stored by this command is generated by looping over all the atoms owned on a processor and their bonds. A bond will only be included if both atoms in the bond are in the specified compute group. Any bonds that have been broken (see the *bond\_style* command) by setting their bond type to 0 are not included. Bonds that have been turned off (see the *fix shake* or *delete\_bonds* commands) by setting their bond type negative are written into the file, but their energy will be 0.0.

Note that as atoms migrate from processor to processor, there will be no consistent ordering of the entries within the local vector or array from one timestep to the next. The only consistency that is guaranteed is that the ordering on a particular timestep will be the same for local vectors or arrays generated by other compute commands. For example, bond output from the *compute property/local* command can be combined with data from this command and output by the *dump local* command in a consistent way.

Here is an example of how to do this:

```
compute 1 all property/local btype batom1 batom2
compute 2 all bond/local dist engpot
dump 1 all local 1000 tmp.dump index c_1[*] c_2[*]
```

### 18.10.4 Output info

This compute calculates a local vector or local array depending on the number of values. The length of the vector or number of rows in the array is the number of bonds. If a single value is specified, a local vector is produced. If two or more values are specified, a local array is produced where the number of columns = the number of values. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The output for *dist* will be in distance *units*. The output for *velvib* will be in velocity *units*. The output for *omega* will be in velocity/distance *units*. The output for *engtrans*, *engvib*, *engrot*, and *engpot* will be in energy *units*. The output for *force* will be in force *units*.

## 18.10.5 Restrictions

none

## 18.10.6 Related commands

*dump local, compute property/local*

## 18.10.7 Default

none

# 18.11 compute born/matrix command

## 18.11.1 Syntax

```
compute ID group-ID born/matrix keyword value ...
```

- ID, group-ID are documented in *compute* command
- born/matrix = style name of this compute command
- zero or more keywords or keyword/value pairs may be appended

keyword = *numdiff* or *pair* or *bond* or *angle* or *dihedral* or *improper*

*numdiff* values = delta virial-ID

delta = magnitude of strain (dimensionless)

virial-ID = ID of pressure compute for virial (string)

(*numdiff* cannot be used with any other keyword)

*pair* = compute pair-wise contributions

*bond* = compute bonding contributions

*angle* = compute angle contributions

*dihedral* = compute dihedral contributions

*improper* = compute improper contributions

## 18.11.2 Examples

```
compute 1 all born/matrix
compute 1 all born/matrix bond angle
compute 1 all born/matrix numdiff 1.0e-4 myvirial
```

### 18.11.3 Description

New in version 4May2022.

Define a compute that calculates  $\frac{\partial^2 U}{\partial \varepsilon_i \partial \varepsilon_j}$ , the second derivatives of the potential energy  $U$  with respect to the strain tensor  $\varepsilon$  elements. These values are related to:

$$C_{i,j}^B = \frac{1}{V} \frac{\partial^2 U}{\partial \varepsilon_i \partial \varepsilon_j}$$

also called the Born term of elastic constants in the stress-stress fluctuation formalism. This quantity can be used to compute the elastic constant tensor. Using the symmetric Voigt notation, the elastic constant tensor can be written as a 6x6 symmetric matrix:

$$C_{i,j} = \langle C_{i,j}^B \rangle + \frac{V}{k_B T} (\langle \sigma_i \sigma_j \rangle - \langle \sigma_i \rangle \langle \sigma_j \rangle) + \frac{N k_B T}{V} (\delta_{i,j} + (\delta_{1,i} + \delta_{2,i} + \delta_{3,i}) * (\delta_{1,j} + \delta_{2,j} + \delta_{3,j}))$$

In the above expression,  $\sigma$  stands for the virial stress tensor,  $\delta$  is the Kronecker delta and the usual notation apply for the number of particle, the temperature and volume respectively  $N$ ,  $T$  and  $V$ .  $k_B$  is the Boltzmann constant.

The Born term is a symmetric 6x6 matrix, as is the matrix of second derivatives of potential energy w.r.t strain, whose 21 independent elements are output in this order:

$$\begin{bmatrix} C_1 & C_7 & C_8 & C_9 & C_{10} & C_{11} \\ C_7 & C_2 & C_{12} & C_{13} & C_{14} & C_{15} \\ \vdots & C_{12} & C_3 & C_{16} & C_{17} & C_{18} \\ \vdots & C_{13} & C_{16} & C_4 & C_{19} & C_{20} \\ \vdots & \vdots & \vdots & C_{19} & C_5 & C_{21} \\ \vdots & \vdots & \vdots & \vdots & C_{21} & C_6 \end{bmatrix}$$

in this matrix the indices of  $C_k$  value are the corresponding element  $k$  in the global vector output by this compute. Each term comes from the sum of the derivatives of every contribution to the potential energy in the system as explained in (*VanWorkum*).

The output can be accessed using usual Lammmps routines:

```
compute 1 all born/matrix
compute 2 all pressure NULL virial
variable S1 equal -c_2[1]
variable S2 equal -c_2[2]
variable S3 equal -c_2[3]
variable S4 equal -c_2[4]
variable S5 equal -c_2[5]
variable S6 equal -c_2[6]
fix 1 all ave/time 1 1 1 v_S1 v_S2 v_S3 v_S4 v_S5 v_S6 c_1[*] file born.out
```

In this example, the file *born.out* will contain the information needed to compute the first and second terms of the elastic constant matrix in a post processing procedure. The other required quantities can be accessed using any other *LAMMPS* usual method. Several examples of this method are provided in the examples/ELASTIC\_T/BORN\_MATRIX directory described on the *Examples* doc page.

NOTE: In the above  $C_{i,j}$  computation, the fluctuation term involving the virial stress tensor  $\sigma$  is the covariance between each elements. In a solid the stress fluctuations can vary rapidly, while average fluctuations can be slow to converge. A detailed analysis of the convergence rate of all the terms in the elastic tensor is provided in the paper by Clavier et al. (*Clavier*).



Two different computation methods for the Born matrix are implemented in this compute and are mutually exclusive.

The first one is a direct computation from the analytical formula from the different terms of the potential used for the simulations (*VanWorkum*). However, the implementation of such derivations must be done for every potential form. This has not been done yet and can be very complicated for complex potentials. At the moment a warning message is displayed for every term that is not supporting the compute at the moment. This method is the default for now.

The second method uses finite differences of energy to numerically approximate the second derivatives (*Zhen*). This is useful when using interaction styles for which the analytical second derivatives have not been implemented. In this cases, the compute applies linear strain fields of magnitude *delta* to all the atoms relative to a point at the center of the box. The strain fields are in six different directions, corresponding to the six Cartesian components of the stress tensor defined by LAMMPS. For each direction it applies the strain field in both the positive and negative senses, and the new stress virial tensor of the entire system is calculated after each. The difference in these two virials divided by two times *delta*, approximates the corresponding components of the second derivative, after applying a suitable unit conversion.

---

**Note:** It is important to choose a suitable value for delta, the magnitude of strains that are used to generate finite difference approximations to the exact virial stress. For typical systems, a value in the range of 1 part in 1e5 to 1e6 will be sufficient. However, the best value will depend on a multitude of factors including the stiffness of the interatomic potential, the thermodynamic state of the material being probed, and so on. The only way to be sure that you have made a good choice is to do a sensitivity study on a representative atomic configuration, sweeping over a wide range of values of delta. If delta is too small, the output values will vary erratically due to truncation effects. If delta is increased beyond a certain point, the output values will start to vary smoothly with delta, due to growing contributions from higher order derivatives. In between these two limits, the numerical virial values should be largely independent of delta.

---

The keyword requires the additional arguments *delta* and *virial-ID*. *delta* gives the size of the applied strains. *virial-ID* gives the ID string of the pressure compute that provides the virial stress tensor, requiring that it use the virial keyword e.g.

```
compute myvirial all pressure NULL virial
compute 1 all born/matrix numdiff 1.0e-4 myvirial
```

#### Output info:

This compute calculates a global vector with 21 values that are the second derivatives of the potential energy with respect to strain. The values are in energy units. The values are ordered as explained above. These values can be used by any command that uses global values from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options.

The array values calculated by this compute are all “extensive”.

### 18.11.4 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. LAMMPS was built with that package. See the [Build package](#) page for more info.

The Born term can be decomposed as a product of two terms. The first one is a general term which depends on the configuration. The second one is specific to every interaction composing your force field (non-bonded, bonds, angle, ...). Currently not all LAMMPS interaction styles implement the *born\_matrix* method giving first and second order derivatives and LAMMPS will exit with an error if this compute is used with such interactions unless the *numdiff* option is also used. The *numdiff* option cannot be used with any other keyword. In this situation, LAMMPS will also exit with an error.

### 18.11.5 Default

none

---

**(Van Workum)** K. Van Workum et al., J. Chem. Phys. 125 144506 (2006)

**(Clavier)** G. Clavier, N. Desbiens, E. Bourasseau, V. Lachet, N. Brusselle-Dupend and B. Rousseau, Mol Sim, 43, 1413 (2017).

**(Zhen)** Y. Zhen, C. Chu, Computer Physics Communications 183(2012)261-265

## 18.12 compute centro/atom command

### 18.12.1 Syntax

```
compute ID group-ID centro/atom lattice keyword value ...
```

- ID, group-ID are documented in *compute* command
- centro/atom = style name of this compute command
- lattice = *fcc* or *bcc* or *N* = # of neighbors per atom to include
- zero or more keyword/value pairs may be appended
- keyword = *axes*

axes value = *no* or *yes*

*no* = do not calculate 3 symmetry axes

*yes* = calculate 3 symmetry axes

### 18.12.2 Examples

```
compute 1 all centro/atom fcc
```

```
compute 1 all centro/atom 8
```

### 18.12.3 Description

Define a computation that calculates the centro-symmetry parameter for each atom in the group, for either FCC or BCC lattices, depending on the choice of the *lattice* argument. In solid-state systems the centro-symmetry parameter is a useful measure of the local lattice disorder around an atom and can be used to characterize whether the atom is part of a perfect lattice, a local defect (e.g. a dislocation or stacking fault), or at a surface.

The value of the centro-symmetry parameter will be 0.0 for atoms not in the specified compute group.

This parameter is computed using the following formula from (*Kelchner*)

$$CS = \sum_{i=1}^{N/2} |\vec{R}_i + \vec{R}_{i+N/2}|^2$$

where the  $N$  nearest neighbors of each atom are identified and  $\vec{R}_i$  and  $\vec{R}_{i+N/2}$  are vectors from the central atom to a particular pair of nearest neighbors. There are  $N(N-1)/2$  possible neighbor pairs that can contribute to this formula.

The quantity in the sum is computed for each, and the  $N/2$  smallest are used. This will typically be for pairs of atoms in symmetrically opposite positions with respect to the central atom; hence the  $i + N/2$  notation.

$N$  is an input parameter, which should be set to correspond to the number of nearest neighbors in the underlying lattice of atoms. If the keyword *fcc* or *bcc* is used,  $N$  is set to 12 and 8 respectively. More generally,  $N$  can be set to a positive, even integer.

For an atom on a lattice site, surrounded by atoms on a perfect lattice, the centro-symmetry parameter will be 0. It will be near 0 for small thermal perturbations of a perfect lattice. If a point defect exists, the symmetry is broken, and the parameter will be a larger positive value. An atom at a surface will have a large positive parameter. If the atom does not have  $N$  neighbors (within the potential cutoff), then its centro-symmetry parameter is set to 0.0.

If the keyword *axes* has the setting *yes*, then this compute also estimates three symmetry axes for each atom's local neighborhood. The first two of these are the vectors joining the two pairs of neighbor atoms with smallest contributions to the centrosymmetry parameter, i.e. the two most symmetric pairs of atoms. The third vector is normal to the first two by the right-hand rule. All three vectors are normalized to unit length. For FCC crystals, the first two vectors will lie along a  $\langle 110 \rangle$  direction, while the third vector will lie along either a  $\langle 100 \rangle$  or  $\langle 111 \rangle$  direction. For HCP crystals, the first two vectors will lie along  $\langle 1000 \rangle$  directions, while the third vector will lie along  $\langle 0001 \rangle$ . This provides a simple way to measure local orientation in HCP structures. In general, the *axes* keyword can be used to estimate the orientation of symmetry axes in the neighborhood of any atom.

Only atoms within the cutoff of the pairwise neighbor list are considered as possible neighbors. Atoms not in the compute group are included in the  $N$  neighbors used in this calculation.

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (e.g., each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently or to have multiple compute/dump commands, each with a *centro/atom* style.

## 18.12.4 Output info

By default, this compute calculates the centrosymmetry value for each atom as a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

If the *axes* keyword setting is *yes*, then a per-atom array is calculated. The first column is the centrosymmetry parameter. The next three columns are the  $x$ ,  $y$ , and  $z$  components of the first symmetry axis, followed by the second, and third symmetry axes in columns 5–7 and 8–10.

The centrosymmetry values are unitless values  $\geq 0.0$ . Their magnitude depends on the lattice style due to the number of contributing neighbor pairs in the summation in the formula above. And it depends on the local defects surrounding the central atom, as described above. For the *axes yes* case, the vector components are also unitless, since they represent spatial directions.

Here are typical centro-symmetry values, from a nanoindentation simulation into gold (FCC). These were provided by Jon Zimmerman (Sandia):

```
Bulk lattice = 0
Dislocation core ~ 1.0 (0.5 to 1.25)
Stacking faults ~ 5.0 (4.0 to 6.0)
Free surface ~ 23.0
```

These values are **not** normalized by the square of the lattice parameter. If they were, normalized values would be:

```
Bulk lattice = 0
Dislocation core ~ 0.06 (0.03 to 0.075)
Stacking faults ~ 0.3 (0.24 to 0.36)
Free surface ~ 1.38
```

For BCC materials, the values for dislocation cores and free surfaces would be somewhat different, due to their being only 8 neighbors instead of 12.

## 18.12.5 Restrictions

none

## 18.12.6 Related commands

*compute cna/atom*

## 18.12.7 Default

The default value for the optional keyword is axes = no.

---

(**Kelchner**) Kelchner, Plimpton, Hamilton, Phys Rev B, 58, 11085 (1998).

# 18.13 compute chunk/atom command

## 18.13.1 Syntax

```
compute ID group-ID chunk/atom style args keyword values ...
```

- ID, group-ID are documented in *compute* command
- chunk/atom = style name of this compute command

style = *bin/1d* or *bin/2d* or *bin/3d* or *bin/sphere* or *bin/cylinder* or *type* or *↔molecule* or *c\_ID*, *c\_ID[I]*, *f\_ID*, *f\_ID[I]*, *v\_name*

*bin/1d* args = dim origin delta

dim = x or y or z

origin = *lower* or *center* or *upper* or coordinate value (distance units)

delta = thickness of spatial bins in dim (distance units)

*bin/2d* args = dim origin delta dim origin delta

dim = x or y or z

origin = *lower* or *center* or *upper* or coordinate value (distance units)

delta = thickness of spatial bins in dim (distance units)

*bin/3d* args = dim origin delta dim origin delta dim origin delta

dim = x or y or z

origin = *lower* or *center* or *upper* or coordinate value (distance units)

delta = thickness of spatial bins in dim (distance units)

*bin/sphere* args = xorig yorig zorig rmin rmax nsbin

xorig,yorig,zorig = center point of sphere

srmin,srmax = bin from sphere radius rmin to rmax

nsbin = # of spherical shell bins between rmin and rmax

*bin/cylinder* args = dim origin delta c1 c2 rmin rmax ncbn

dim = x or y or z = axis of cylinder axis

origin = *lower* or *center* or *upper* or coordinate value (distance units)

delta = thickness of spatial bins in dim (distance units)

```

c1,c2 = coords of cylinder axis in other 2 dimensions (distance units)
crmin,crmax = bin from cylinder radius rmin to rmax (distance units)
ncbin = # of concentric circle bins between rmin and rmax
type args = none
molecule args = none
c_ID, c_ID[I], f_ID, f_ID[I], v_name args = none
c_ID = per-atom vector calculated by a compute with ID
c_ID[I] = Ith column of per-atom array calculated by a compute with ID
f_ID = per-atom vector calculated by a fix with ID
f_ID[I] = Ith column of per-atom array calculated by a fix with ID
v_name = per-atom vector calculated by an atom-style variable with name

```

- zero or more keyword/values pairs may be appended
- keyword = *region* or *nchunk* or *limit* or *ids* or *compress* or *discard* or *bound* or *pbcs* or *units*

```

region value = region-ID
region-ID = ID of region atoms must be in to be part of a chunk
nchunk value = once or every
once = only compute the number of chunks once
every = re-compute the number of chunks whenever invoked
limit values = 0 or Nc max or Nc exact
0 = no limit on the number of chunks
Nc max = limit number of chunks to be <= Nc
Nc exact = set number of chunks to exactly Nc
ids value = once or nfreq or every
once = assign chunk IDs to atoms only once, they persist thereafter
nfreq = assign chunk IDs to atoms only once every Nfreq steps (if invoked by fix_
→ave/chunk which sets Nfreq)
every = assign chunk IDs to atoms whenever invoked
compress value = yes or no
yes = compress chunk IDs to eliminate IDs with no atoms
no = do not compress chunk IDs even if some IDs have no atoms
discard value = yes or no or mixed
yes = discard atoms with out-of-range chunk IDs by assigning a chunk ID = 0
no = keep atoms with out-of-range chunk IDs by assigning a valid chunk ID
mixed = keep or discard such atoms according to spatial binning rule
bound values = x/y/z lo hi
x/y/z = x or y or z to bound spatial bins in this dimension
lo = lower or coordinate value (distance units)
hi = upper or coordinate value (distance units)
pbcs value = no or yes
yes = use periodic distance for bin/sphere and bin/cylinder styles
units value = box or lattice or reduced

```

## 18.13.2 Examples

```
compute 1 all chunk/atom type
compute 1 all chunk/atom bin/1d z lower 0.02 units reduced
compute 1 all chunk/atom bin/2d z lower 1.0 y 0.0 2.5
compute 1 all chunk/atom molecule region sphere nchunk once ids once compress yes
compute 1 all chunk/atom bin/sphere 5 5 5 2.0 5.0 5 discard yes
compute 1 all chunk/atom bin/cylinder z lower 2 10 10 2.0 5.0 3 discard yes
compute 1 all chunk/atom c_cluster
```

## 18.13.3 Description

Define a computation that calculates an integer chunk ID from 1 to *Nchunk* for each atom in the group. Values of chunk IDs are determined by the *style* of chunk, which can be based on atom type or molecule ID or spatial binning or a per-atom property or value calculated by another *compute*, *fix*, or *atom-style variable*. Per-atom chunk IDs can be used by other computes with “chunk” in their style name, such as *compute com/chunk* or *compute msd/chunk*. Or they can be used by the *fix ave/chunk* command to sum and time average a variety of per-atom properties over the atoms in each chunk. Or they can simply be accessed by any command that uses per-atom values from a compute as input, as discussed on the *Howto output* doc page.

See the *Howto chunk* page for an overview of how this compute can be used with a variety of other commands to tabulate properties of a simulation. The page gives several examples of input script commands that can be used to calculate interesting properties.

Conceptually it is important to realize that this compute does two simple things. First, it sets the value of *Nchunk* = the number of chunks, which can be a constant value or change over time. Second, it assigns each atom to a chunk via a chunk ID. Chunk IDs range from 1 to *Nchunk* inclusive; some chunks may have no atoms assigned to them. Atoms that do not belong to any chunk are assigned a value of 0. Note that the two operations are not always performed together. For example, spatial bins can be setup once (which sets *Nchunk*), and atoms assigned to those bins many times thereafter (setting their chunk IDs).

All other commands in LAMMPS that use chunk IDs assume there are *Nchunk* number of chunks, and that every atom is assigned to one of those chunks, or not assigned to any chunk.

There are many options for specifying for how and when *Nchunk* is calculated, and how and when chunk IDs are assigned to atoms. The details depend on the chunk *style* and its *args*, as well as optional keyword settings. They can also depend on whether a *fix ave/chunk* command is using this compute, since that command requires *Nchunk* to remain static across windows of timesteps it specifies, while it accumulates per-chunk averages.

The details are described below.

---

The different chunk styles operate as follows. For each style, how it calculates *Nchunk* and assigns chunk IDs to atoms is explained. Note that using the optional keywords can change both of those actions, as described further below where the keywords are discussed.

---

The *binning* styles perform a spatial binning of atoms, and assign an atom the chunk ID corresponding to the bin number it is in. *Nchunk* is set to the number of bins, which can change if the simulation box size changes. This also depends on the setting of the *units* keyword (e.g., for *reduced* units the number of chunks may not change even if the box size does).

The *bin/1d*, *bin/2d*, and *bin/3d* styles define bins as 1d layers (slabs), 2d pencils, or 3d boxes. The *dim*, *origin*, and *delta* settings are specified 1, 2, or 3 times. For 2d or 3d bins, there is no restriction on specifying *dim = x* before *dim = y* or *z*, or *dim = y* before *dim = z*. Bins in a particular *dim* have a bin size in that dimension given by *delta*. In each

dimension, bins are defined relative to a specified *origin*, which may be the lower/upper edge of the simulation box (in that dimension), or its center point, or a specified coordinate value. Starting at the origin, sufficient bins are created in both directions to completely span the simulation box or the bounds specified by the optional *bounds* keyword.

For orthogonal simulation boxes, the bins are layers, pencils, or boxes aligned with the xyz coordinate axes. For triclinic (non-orthogonal) simulation boxes, the bin faces are parallel to the tilted faces of the simulation box. See the [Howto triclinic](#) page for a discussion of the geometry of triclinic boxes in LAMMPS. As described there, a tilted simulation box has edge vectors  $\vec{a}$ ,  $\vec{b}$ , and  $\vec{c}$ . In that nomenclature, bins in the  $x$  dimension have faces with normals in the  $\vec{b} \times \vec{c}$  direction, bins in  $y$  have faces normal to the  $\vec{a} \times \vec{c}$  direction, and bins in  $z$  have faces normal to the  $\vec{a} \times \vec{b}$  direction. Note that in order to define the size and position of these bins in an unambiguous fashion, the *units* option must be set to *reduced* when using a triclinic simulation box, as noted below.

The meaning of *origin* and *delta* for triclinic boxes is as follows. Consider a triclinic box with bins that are 1d layers or slabs in the  $x$  dimension. No matter how the box is tilted, an *origin* of 0.0 means start layers at the lower  $\vec{b} \times \vec{c}$  plane of the simulation box and an *origin* of 1.0 means to start layers at the upper  $\vec{b} \times \vec{c}$  face of the box. A *delta* value of 0.1 in *reduced* units means there will be 10 layers from 0.0 to 1.0, regardless of the current size or shape of the simulation box.

The *bin/sphere* style defines a set of spherical shell bins around the origin ( $xorig, yorig, zorig$ ), using *nsbin* bins with radii equally spaced between *srmin* and *srmax*. This is effectively a 1d vector of bins. For example, if *srmin* = 1.0 and *srmax* = 10.0 and *nsbin* = 9, then the first bin spans  $1.0 < r < 2.0$ , and the last bin spans  $9.0 < r < 10.0$ . The geometry of the bins is the same whether the simulation box is orthogonal or triclinic (i.e., the spherical shells are not tilted or scaled differently in different dimensions to transform them into ellipsoidal shells).

The *bin/cylinder* style defines bins for a cylinder oriented along the axis *dim* with the axis coordinates in the other two radial dimensions at ( $c1, c2$ ). For  $dim = x$ ,  $c_1/c_2 = y/z$ ; for  $dim = y$ ,  $c_1/c_2 = x/z$ ; for  $dim = z$ ,  $c_1/c_2 = x/y$ . This is effectively a 2d array of bins. The first dimension is along the cylinder axis, the second dimension is radially outward from the cylinder axis. The bin size and positions along the cylinder axis are specified by the *origin* and *delta* values, the same as for the *bin/1d*, *bin/2d*, and *bin/3d* styles. There are *ncbin* concentric circle bins in the radial direction from the cylinder axis with radii equally spaced between *crmin* and *crmax*. For example, if *crmin* = 1.0 and *crmax* = 10.0 and *ncbin* = 9, then the first bin spans  $1.0 < r < 2.0$  and the last bin spans  $9.0 < r < 10.0$ . The geometry of the bins in the radial dimensions is the same whether the simulation box is orthogonal or triclinic (i.e., the concentric circles are not tilted or scaled differently in the two different dimensions to transform them into ellipses).

The created bins (and hence the chunk IDs) are numbered consecutively from 1 to the number of bins = *Nchunk*. For *bin2d* and *bin3d*, the numbering varies most rapidly in the first dimension (which could be  $x$ ,  $y$ , or  $z$ ), next rapidly in the second dimension, and most slowly in the third dimension. For *bin/sphere*, the bin with smallest radii is chunk 1 and the bin with largest radii is chunk  $Nchunk = ncbin$ . For *bin/cylinder*, the numbering varies most rapidly in the dimension along the cylinder axis and most slowly in the radial direction.

Each time this compute is invoked, each atom is mapped to a bin based on its current position. Note that between reneighboring timesteps, atoms can move outside the current simulation box. If the box is periodic (in that dimension) the atom is remapping into the periodic box for purposes of binning. If the box is not periodic, the atom may have moved outside the bounds of all bins. If an atom is not inside any bin, the *discard* keyword is used to determine how a chunk ID is assigned to the atom.

---

The *type* style uses the atom type as the chunk ID. *Nchunk* is set to the number of atom types defined for the simulation (e.g., via the *create\_box* or *read\_data* commands).

---

The *molecule* style uses the molecule ID of each atom as its chunk ID. *Nchunk* is set to the largest chunk ID. Note that this excludes molecule IDs for atoms which are not in the specified group or optional region.

There is no requirement that all atoms in a particular molecule are assigned the same chunk ID (zero or non-zero), though you probably want that to be the case, if you wish to compute a per-molecule property. LAMMPS will issue a warning if that is not the case, but only the first time that *Nchunk* is calculated.

Note that atoms with a molecule ID = 0, which may be non-molecular solvent atoms, have an out-of-range chunk ID. These atoms are discarded (not assigned to any chunk) or assigned to *Nchunk*, depending on the value of the *discard* keyword.

---

The *compute/fix/variable* styles set the chunk ID of each atom based on a quantity calculated and stored by a compute, fix, or variable. In each case, it must be a per-atom quantity. In each case the referenced floating point values are converted to an integer chunk ID as follows. The floating point value is truncated (rounded down) to an integer value. If the integer value is  $\leq 0$ , then a chunk ID of 0 is assigned to the atom. If the integer value is  $> 0$ , it becomes the chunk ID to the atom. *Nchunk* is set to the largest chunk ID. Note that this excludes atoms which are not in the specified group or optional region.

If the style begins with “c\_”, a compute ID must follow which has been previously defined in the input script. If no bracketed integer is appended, the per-atom vector calculated by the compute is used. If a bracketed integer is appended, the *I*th column of the per-atom array calculated by the compute is used. Users can also write code for their own compute styles and *add them to LAMMPS*.

If the style begins with “f\_”, a fix ID must follow which has been previously defined in the input script. If no bracketed integer is appended, the per-atom vector calculated by the fix is used. If a bracketed integer is appended, the *I*th column of the per-atom array calculated by the fix is used. Note that some fixes only produce their values on certain timesteps, which must be compatible with the timestep on which this compute accesses the fix, else an error results. Users can also write code for their own fix styles and *add them to LAMMPS*.

If a value begins with “v\_”, a variable name for an *atom* or *atomfile* style *variable* must follow which has been previously defined in the input script. Variables of style *atom* can reference thermodynamic keywords and various per-atom attributes, or invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of generating per-atom quantities to treat as a chunk ID.

---

Normally, *Nchunk* = the number of chunks, is re-calculated every time this fix is invoked, though the value may or may not change. As explained below, the *nchunk* keyword can be set to *once* which means *Nchunk* will never change.

If a *fix ave/chunk* command uses this compute, it can also turn off the re-calculation of *Nchunk* for one or more windows of timesteps. The extent of the windows, during which *Nchunk* is held constant, are determined by the *Nevery*, *Nrepeat*, *Nfreq* values and the *ave* keyword setting that are used by the *fix ave/chunk* command.

Specifically, if *ave* = *one*, then for each span of *Nfreq* timesteps, *Nchunk* is held constant between the first timestep when averaging is done (within the *Nfreq*-length window), and the last timestep when averaging is done (multiple of *Nfreq*). If *ave* = *running* or *window*, then *Nchunk* is held constant forever, starting on the first timestep when the *fix ave/chunk* command invokes this compute.

Note that multiple *fix ave/chunk* commands can use the same compute chunk/atom compute. However, the time windows they induce for holding *Nchunk* constant must be identical, else an error will be generated.

---

The various optional keywords operate as follows. Note that some of them function differently or are ignored by different chunk styles. Some of them also have different default values, depending on the chunk style, as listed below.

The *region* keyword applies to all chunk styles. If used, an atom must be in both the specified group and the specified geometric *region* to be assigned to a chunk.

---

The *nchunk* keyword applies to all chunk styles. It specifies how often *Nchunk* is recalculated, which in turn can affect the chunk IDs assigned to individual atoms.

If *nchunk* is set to *once*, then *Nchunk* is only calculated once, the first time this compute is invoked. If *nchunk* is set to *every*, then *Nchunk* is re-calculated every time the compute is invoked. Note that, as described above, the use of this compute by the *fix ave/chunk* command can override the *every* setting.

---



The default values for *nchunk* are listed below and depend on the chunk style and other system and keyword settings. They attempt to represent typical use cases for the various chunk styles. The *nchunk* value can always be set explicitly if desired.

---

The *limit* keyword can be used to limit the calculated value of *Nchunk* = the number of chunks. The limit is applied each time *Nchunk* is calculated, which also limits the chunk IDs assigned to any atom. The *limit* keyword is used by all chunk styles except the *binning* styles, which ignore it. This is because the number of bins can be tailored using the *bound* keyword (described below) which effectively limits the size of *Nchunk*.

If *limit* is set to  $Nc = 0$ , then no limit is imposed on *Nchunk*, though the *compress* keyword can still be used to reduce *Nchunk*, as described below.

If  $Nc > 0$ , then the effect of the *limit* keyword depends on whether the *compress* keyword is also used with a setting of *yes*, and whether the *compress* keyword is specified before the *limit* keyword or after.

In all cases, *Nchunk* is first calculated in the usual way for each chunk style, as described above.

First, here is what occurs if *compress yes* is not set. If *limit* is set to *Nc max*, then *Nchunk* is reset to the smaller of *Nchunk* and *Nc*. If *limit* is set to *Nc exact*, then *Nchunk* is reset to *Nc*, whether the original *Nchunk* was larger or smaller than *Nc*. If *Nchunk* shrank due to the *limit* setting, then atom chunk IDs  $> Nchunk$  will be reset to 0 or *Nchunk*, depending on the setting of the *discard* keyword. If *Nchunk* grew, there will simply be some chunks with no atoms assigned to them.

If *compress yes* is set, and the *compress* keyword comes before the *limit* keyword, the compression operation is performed first, as described below, which resets *Nchunk*. The *limit* keyword is then applied to the new *Nchunk* value, exactly as described in the preceding paragraph. Note that in this case, all atoms will end up with chunk IDs  $\leq Nc$ , but their original values (e.g., molecule ID or compute/fix/variable) may have been  $> Nc$ , because of the compression operation.

If *compress yes* is set, and the *compress* keyword comes after the *limit* keyword, then the *limit* value of *Nc* is applied first to the uncompressed value of *Nchunk*, but only if  $Nc < Nchunk$  (whether *Nc max* or *Nc exact* is used). This effectively means all atoms with chunk IDs  $> Nc$  have their chunk IDs reset to 0 or *Nc*, depending on the setting of the *discard* keyword. The compression operation is then performed, which may shrink *Nchunk* further. If the new  $Nchunk < Nc$  and *limit* = *Nc exact* is specified, then *Nchunk* is reset to *Nc*, which results in extra chunks with no atoms assigned to them. Note that in this case, all atoms will end up with chunk IDs  $\leq Nc$ , and their original values (e.g., molecule ID or compute/fix/variable value) will also have been  $\leq Nc$ .

---

The *ids* keyword applies to all chunk styles. If the setting is *once* then the chunk IDs assigned to atoms the first time this compute is invoked will be permanent, and never be re-computed.

If the setting is *nfreq* and if a *fix ave/chunk* command is using this compute, then in each of the *Nchunk* = constant time windows (discussed above), the chunk ID's assigned to atoms on the first step of the time window will persist until the end of the time window.

If the setting is *every*, which is the default, then chunk IDs are re-calculated on any timestep this compute is invoked.

---

**Note:** If you want the persistent chunk-IDs calculated by this compute to be continuous when running from a *restart file*, then you should use the same ID for this compute, as in the original run. This is so that the fix this compute creates to store per-atom quantities will also have the same ID, and thus be initialized correctly with chunk IDs from the restart file.

---

The *compress* keyword applies to all chunk styles and affects how *Nchunk* is calculated, which in turn affects the chunk IDs assigned to each atom. It is useful for converting a “sparse” set of chunk IDs (with many IDs that have no atoms assigned to them), into a “dense” set of IDs, where every chunk has one or more atoms assigned to it.

---

Two possible use cases are as follows. If a large simulation box is mostly empty space, then the *binning* style may produce many bins with no atoms. If *compress* is set to *yes*, only bins with atoms will be contribute to *Nchunk*. Likewise, the *molecule* or *compute/fix/variable* styles may produce large *Nchunk* values. For example, the *compute cluster/atom* command assigns every atom an atom ID for one of the atoms it is clustered with. For a million-atom system with 5 clusters, there would only be 5 unique chunk IDs, but the largest chunk ID might be 1 million, resulting in *Nchunk* = 1 million. If *compress* is set to *yes*, *Nchunk* will be reset to 5.

If *compress* is set to *no*, which is the default, no compression is done. If it is set to *yes*, all chunk IDs with no atoms are removed from the list of chunk IDs, and the list is sorted. The remaining chunk IDs are renumbered from 1 to *Nchunk* where *Nchunk* is the new length of the list. The chunk IDs assigned to each atom reflect the new renumbering from 1 to *Nchunk*.

The original chunk IDs (before renumbering) can be accessed by the *compute property/chunk* command and its *id* keyword, or by the *fix ave/chunk* command which outputs the original IDs as one of the columns in its global output array. For example, using the “compute cluster/atom” command discussed above, the original 5 unique chunk IDs might be atom IDs (27,4982,58374,857838,1000000). After compression, these will be renumbered to (1,2,3,4,5). The original values (27,...,1000000) can be output to a file by the *fix ave/chunk* command, or by using the *fix ave/time* command in conjunction with the *compute property/chunk* command.

---

**Note:** The compression operation requires global communication across all processors to share their chunk ID values. It can require large memory on every processor to store them, even after they are compressed, if there are a large number of unique chunk IDs with atoms assigned to them. It uses a STL map to find unique chunk IDs and store them in sorted order. Each time an atom is assigned a compressed chunk ID, it must access the STL map. All of this means that compression can be expensive, both in memory and CPU time. The use of the *limit* keyword in conjunction with the *compress* keyword can affect these costs, depending on which keyword is used first. So use this option with care.

---

The *discard* keyword applies to all chunk styles. It affects what chunk IDs are assigned to atoms that do not match one of the valid chunk IDs from 1 to *Nchunk*. Note that it does not apply to atoms that are not in the specified group or optionally specified region. Those atoms are always assigned a chunk ID = 0.

If the calculated chunk ID for an atom is not within the range 1 to *Nchunk* then it is a “discard” atom. Note that *Nchunk* may have been shrunk by the *limit* keyword. Or the *compress* keyword may have eliminated chunk IDs that were valid before the compression took place, and are now not in the compressed list. Also note that for the *molecule* chunk style, if new molecules are added to the system, their chunk IDs may exceed a previously calculated *Nchunk*. Likewise, evaluation of a *compute/fix/variable* on a later timestep may return chunk IDs that are invalid for the previously calculated *Nchunk*.

All the chunk styles except the *binning* styles, must use *discard* set to either *yes* or *no*. If *discard* is set to *yes*, which is the default, then every “discard” atom has its chunk ID set to 0. If *discard* is set to *no*, every “discard” atom has its chunk ID set to *Nchunk*. I.e. it becomes part of the last chunk.

The *binning* styles use the *discard* keyword to decide whether to discard atoms outside the spatial domain covered by bins, or to assign them to the bin they are nearest to.

For the *bin/1d*, *bin/2d*, *bin/3d* styles the details are as follows. If *discard* is set to *yes*, an out-of-domain atom will have its chunk ID set to 0. If *discard* is set to *no*, the atom will have its chunk ID set to the first or last bin in that dimension. If *discard* is set to *mixed*, which is the default, it will only have its chunk ID set to the first or last bin if bins extend to the simulation box boundary in that dimension. This is the case if the *bound* keyword settings are *lower* and *upper*, which is the default. If the *bound* keyword settings are numeric values, then the atom will have its chunk ID set to 0 if it is outside the bounds of any bin. Note that in this case, it is possible that the first or last bin extends beyond the numeric *bounds* settings, depending on the specified *origin*. If this is the case, the chunk ID of the atom is only set to 0 if it is outside the first or last bin, not if it is simply outside the numeric *bounds* setting.

For the *bin/sphere* style the details are as follows. If *discard* is set to *yes*, an out-of-domain atom will have its chunk ID set to 0. If *discard* is set to *no* or *mixed*, the atom will have its chunk ID set to the first or last bin, i.e. the innermost

or outermost spherical shell. If the distance of the atom from the origin is less than  $rmin$ , it will be assigned to the first bin. If the distance of the atom from the origin is greater than  $rmax$ , it will be assigned to the last bin.

For the *bin/cylinder* style the details are as follows. If *discard* is set to *yes*, an out-of-domain atom will have its chunk ID set to 0. If *discard* is set to *no*, the atom will have its chunk ID set to the first or last bin in both the radial and axis dimensions. If *discard* is set to *mixed*, which is the default, the radial dimension is treated the same as for *discard* = *no*. But for the axis dimension, it will only have its chunk ID set to the first or last bin if bins extend to the simulation box boundary in the axis dimension. This is the case if the *bound* keyword settings are *lower* and *upper*, which is the default. If the *bound* keyword settings are numeric values, then the atom will have its chunk ID set to 0 if it is outside the bounds of any bin. Note that in this case, it is possible that the first or last bin extends beyond the numeric *bounds* settings, depending on the specified *origin*. If this is the case, the chunk ID of the atom is only set to 0 if it is outside the first or last bin, not if it is simply outside the numeric *bounds* setting.

If *discard* is set to *no* or *mixed*, the atom will have its chunk ID set to the first or last bin, i.e. the innermost or outermost spherical shell. If the distance of the atom from the origin is less than  $rmin$ , it will be assigned to the first bin. If the distance of the atom from the origin is greater than  $rmax$ , it will be assigned to the last bin.

---

The *bound* keyword only applies to the *bin/1d*, *bin/2d*, *bin/3d* styles and to the axis dimension of the *bin/cylinder* style; otherwise it is ignored. It can be used one or more times to limit the extent of bin coverage in a specified dimension, i.e. to only bin a portion of the box. If the *lo* setting is *lower* or the *hi* setting is *upper*, the bin extent in that direction extends to the box boundary. If a numeric value is used for *lo* and/or *hi*, then the bin extent in the *lo* or *hi* direction extends only to that value, which is assumed to be inside (or at least near) the simulation box boundaries, though LAMMPS does not check for this. Note that using the *bound* keyword typically reduces the total number of bins and thus the number of chunks *Nchunk*.

The *pbz* keyword only applies to the *bin/sphere* and *bin/cylinder* styles. If set to *yes*, the distance an atom is from the sphere origin or cylinder axis is calculated in a minimum image sense with respect to periodic dimensions, when determining which bin the atom is in. I.e. if  $x$  is a periodic dimension and the distance between the atom and the sphere center in the  $x$  dimension is greater than  $0.5 * \text{simulation box length in } x$ , then a box length is subtracted to give a distance  $< 0.5 * \text{simulation box length}$ . This allows the sphere or cylinder center to be near a box edge, and atoms on the other side of the periodic box will still be close to the center point/axis. Note that with a setting of *yes*, the outer sphere or cylinder radius must also be  $\leq 0.5 * \text{simulation box length}$  in any periodic dimension except for the cylinder axis dimension, or an error is generated.

The *units* keyword only applies to the *binning* styles; otherwise it is ignored. For the *bin/1d*, *bin/2d*, *bin/3d* styles, it determines the meaning of the distance units used for the bin sizes *delta* and for *origin* and *bounds* values if they are coordinate values. For the *bin/sphere* style it determines the meaning of the distance units used for *xorig*, *yorig*, *zorig* and the radii *srmin* and *srmax*. For the *bin/cylinder* style it determines the meaning of the distance units used for *delta*, *c1*, *c2* and the radii *crmin* and *crmax*.

For orthogonal simulation boxes, any of the 3 options may be used. For non-orthogonal (triclinic) simulation boxes, only the *reduced* option may be used.

A *box* value selects standard distance units as defined by the *units* command (e.g., Å for units = *real* or *metal*). A *lattice* value means the distance units are in lattice spacings. The *lattice* command must have been previously used to define the lattice spacing. A *reduced* value means normalized unitless values between 0 and 1, which represent the lower and upper faces of the simulation box respectively. Thus an *origin* value of 0.5 means the center of the box in any dimension. A *delta* value of 0.1 means 10 bins span the box in that dimension.

Note that for the *bin/sphere* style, the radii *srmin* and *srmax* are scaled by the lattice spacing or reduced value of the  $x$  dimension.

Note that for the *bin/cylinder* style, the radii *crmin* and *crmax* are scaled by the lattice spacing or reduced value of the first dimension perpendicular to the cylinder axis (e.g.,  $y$  for an  $x$ -axis cylinder,  $x$  for a  $y$ -axis cylinder, and  $x$  for a  $z$ -axis cylinder).

### 18.13.4 Output info

This compute calculates a per-atom vector (the chunk ID), which can be accessed by any command that uses per-atom values from a compute as input. It also calculates a global scalar (the number of chunks), which can be similarly accessed everywhere outside of a per-atom context. See the *Howto output* page for an overview of LAMMPS output options.

The per-atom vector values are unitless chunk IDs, ranging from 1 to *Nchunk* (inclusive) for atoms assigned to chunks, and 0 for atoms not belonging to a chunk. The scalar contains the value of *Nchunk*.

### 18.13.5 Restrictions

Even if the *nchunk* keyword is set to *once*, the chunk IDs assigned to each atom are not stored in a restart files. This means you cannot expect those assignments to persist in a restarted simulation. Instead you must re-specify this command and assign atoms to chunks when the restarted simulation begins.

### 18.13.6 Related commands

*fix ave/chunk*, *compute global/atom*

### 18.13.7 Default

The option defaults are as follows:

- region = none
- nchunk = every, if compress is yes, overriding other defaults listed here
- nchunk = once, for type style
- nchunk = once, for mol style if region is none
- nchunk = every, for mol style if region is set
- nchunk = once, for binning style if the simulation box size is static or units = reduced
- nchunk = every, for binning style if the simulation box size is dynamic and units is lattice or box
- nchunk = every, for compute/fix/variable style
- limit = 0
- ids = every
- compress = no
- discard = yes, for all styles except binning
- discard = mixed, for binning styles
- bound = lower and upper in all dimensions
- pbc = no
- units = lattice

## 18.14 compute chunk/spread/atom command

### 18.14.1 Syntax

```
compute ID group-ID chunk/spread/atom chunkID input1 input2 ...
```

- ID, group-ID are documented in *compute* command
- chunk/spread/atom = style name of this compute command
- chunkID = ID of *compute chunk/atom* command
- one or more inputs can be listed
- input = c\_ID, c\_ID[N], f\_ID, f\_ID[N]

```
c_ID = global vector calculated by a compute with ID
c_ID[I] = Ith column of global array calculated by a compute with ID, I can include
↳wildcard (see below)
f_ID = global vector calculated by a fix with ID
f_ID[I] = Ith column of global array calculated by a fix with ID, I can include
↳wildcard (see below)
```

### 18.14.2 Examples

```
compute 1 all chunk/spread/atom mychunk c_com[*] c_gyration
```

### 18.14.3 Description

Define a calculation that “spreads” one or more per-chunk values to each atom in the chunk. This can be useful in several scenarios:

- For creating a *dump file* where each atom lists info about the chunk it is in, e.g. for post-processing purposes.
- To access chunk value in *atom-style variables* that need info about the chunk each atom is in.
- To use the *fix ave/chunk* command to spatially average per-chunk values calculated by a per-chunk compute.

Examples are given below.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

For inputs that are computes, they must be a compute that calculates per-chunk values. These are computes whose style names end in “/chunk”.

For inputs that are fixes, they should be a fix that calculates per-chunk values. For example, *fix ave/chunk* or *fix ave/time* (assuming it is time-averaging per-chunk data).

For each atom, this compute accesses its chunk ID from the specified *chunkID* compute, then accesses the per-chunk value in each input. Those values are copied to this compute to become the output for that atom.

The values generated by this compute will be 0.0 for atoms not in the specified compute group *group-ID*. They will also be 0.0 if the atom is not in a chunk, as assigned by the *chunkID* compute. They will also be 0.0 if the current chunk ID for the atom is out-of-bounds with respect to the number of chunks stored by a particular input compute or fix.

---

**Note:** LAMMPS does not check that a compute or fix which calculates per-chunk values uses the same definition of chunks as this compute. It's up to you to be consistent. Likewise, for a fix input, LAMMPS does not check that it is per-chunk data. It only checks that the fix produces a global vector or array.

---

Each listed input is operated on independently.

If a bracketed index I is used, it can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If N = the number of columns in the array, then an asterisk with no numeric values means all indices from 1 to N. A leading asterisk means all indices from 1 to n (inclusive). A trailing asterisk means all indices from n to N (inclusive). A middle asterisk means all indices from m to n (inclusive).

Using a wildcard is the same as if the individual columns of the array had been listed one by one. E.g. these 2 compute chunk/spread/atom commands are equivalent, since the *compute com/chunk* command creates a per-atom array with 3 columns:

```
compute com all com/chunk mychunk
compute 10 all chunk/spread/atom mychunk c_com[*]
compute 10 all chunk/spread/atom mychunk c_com[1] c_com[2] c_com[3]
```

Here is an example of writing a dump file the with the center-of-mass (COM) for the chunk each atom is in. The commands below can be added to the bench/in.chain script.

```
compute      cmol all chunk/atom molecule
compute      com all com/chunk cmol
compute      comchunk all chunk/spread/atom cmol c_com[*]
dump         1 all custom 50 tmp.dump id mol type x y z c_comchunk[*]
dump_modify  1 sort id
```

The same per-chunk data for each atom could be used to define per-atom forces for the *fix addforce* command. In this example the forces act to pull atoms of an extended polymer chain towards its COM in an attractive manner.

```
compute      prop all property/atom xu yu zu
variable     k equal 0.1
variable     fx atom v_k*(c_comchunk[1]-c_prop[1])
variable     fy atom v_k*(c_comchunk[2]-c_prop[2])
variable     fz atom v_k*(c_comchunk[3]-c_prop[3])
fix          3 all addforce v_fx v_fy v_fz
```

Note that *compute property/atom* is used to generate unwrapped coordinates for use in the per-atom force calculation, so that the effect of periodic boundaries is accounted for properly.

Over time this applied force could shrink each polymer chain's radius of gyration in a polymer mixture simulation. Here is output from the bench/in.chain script. Thermo output is shown for 1000 steps, where the last column is the average radius of gyration over all 320 chains in the 32000 atom system:

```
compute      gyr all gyration/chunk cmol
variable     ave equal ave(c_gyr)
thermo_style custom step etotal press v_ave

      0      22.394765      4.6721833      5.128278
     100     22.445002      4.8166709      5.0348372
```

(continues on next page)

(continued from previous page)

200	22.500128	4.8790392	4.9364875
300	22.534686	4.9183766	4.8590693
400	22.557196	4.9492211	4.7937849
500	22.571017	4.9161853	4.7412008
600	22.573944	5.0229708	4.6931243
700	22.581804	5.0541301	4.6440647
800	22.584683	4.9691734	4.6000016
900	22.59128	5.0247538	4.5611513
1000	22.586832	4.94697	4.5238362

Here is an example for using one set of chunks, defined for molecules, to compute the dipole moment vector for each chunk. E.g. for water molecules. Then spreading those values to each atom in each chunk. Then defining a second set of chunks based on spatial bins. And finally, using the *fix ave/chunk* command to calculate an average dipole moment vector per spatial bin.

```
compute      cmol all chunk/atom molecule
compute      dipole all dipole/chunk cmol
compute      spread all chunk/spread/atom cmol c_dipole[1] c_dipole[2] c_dipole[3]
compute      cspatial all chunk/atom bin/1d z lower 0.1 units reduced
fix          ave all ave/chunk 100 10 1000 cspatial c_spread[*]
```

Note that the *fix ave/chunk* command requires per-atom values as input. That is why the compute chunk/spread/atom command is used to assign per-chunk values to each atom in the chunk. If a molecule straddles bin boundaries, each of its atoms contributes in a weighted manner to the average dipole moment of the spatial bin it is in.

### 18.14.4 Output info

This compute calculates a per-atom vector or array, which can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The output is a per-atom vector if a single input value is specified, otherwise a per-atom array is output. The number of columns in the array is the number of inputs provided. The per-atom values for the vector or each column of the array will be in whatever *units* the corresponding input value is in.

The vector or array values are “intensive”.

### 18.14.5 Restrictions

none

### 18.14.6 Related commands

*compute chunk/atom, fix ave/chunk, compute reduce/chunk*

### 18.14.7 Default

none

## 18.15 compute cluster/atom command

## 18.16 compute fragment/atom command

## 18.17 compute aggregate/atom command

### 18.17.1 Syntax

```
compute ID group-ID cluster/atom cutoff
compute ID group-ID fragment/atom keyword value ...
compute ID group-ID aggregate/atom cutoff
```

- ID, group-ID are documented in *compute* command
- *cluster/atom* or *fragment/atom* or *aggregate/atom* = style name of this compute command
- cutoff = distance within which to label atoms as part of same cluster (distance units)
- zero or more keyword/value pairs may be appended to *fragment/atom*
- keyword = *single*

*single* value = *yes* or *no* to treat single atoms (no bonds) as fragments

### 18.17.2 Examples

```
compute 1 all cluster/atom 3.5
compute 1 all fragment/atom
compute 1 all fragment/atom single no
compute 1 all aggregate/atom 3.5
```

### 18.17.3 Description

Define a computation that assigns each atom a cluster, fragment, or aggregate ID. Only atoms in the compute group are clustered and assigned cluster IDs. Atoms not in the compute group are assigned an ID = 0.

A cluster is defined as a set of atoms, each of which is within the cutoff distance from one or more other atoms in the cluster. If an atom has no neighbors within the cutoff distance, then it is a 1-atom cluster.

A fragment is similarly defined as a set of atoms, each of which has a bond to another atom in the fragment. Bonds can be defined initially via the *data file* or *create\_bonds* commands, or dynamically by fixes which create or break bonds



like *fix bond/react*, *fix bond/create*, *fix bond/swap*, or *fix bond/break*. The cluster ID or fragment ID of every atom in the cluster will be set to the smallest atom ID of any atom in the cluster or fragment, respectively.

For the *fragment/atom* style, the *single* keyword determines whether single atoms (not bonded to another atom) are treated as one-atom fragments or not, based on the *yes* or *no* setting. If the setting is *no* (the default), their fragment IDs are set to 0.

An aggregate is defined by combining the rules for clusters and fragments (i.e., a set of atoms, where each of them is within the cutoff distance from one or more atoms within a fragment that is part of the same cluster). This measure can be used to track molecular assemblies like micelles.

For computes *cluster/atom* and *aggregate/atom* a neighbor list needed to compute cluster IDs is constructed each time the compute is invoked. Thus it can be inefficient to compute/dump this quantity too frequently or to have multiple *cluster/atom* or *aggregate/atom* style computes.

---

**Note:** If you have a bonded system, then the settings of *special\_bonds* command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the *special\_bonds* command, and means those pairwise interactions do not appear in the neighbor list. Because this fix uses the neighbor list, it also means those pairs will not be included when computing the clusters. This does not apply when using long-range coulomb (*coul/long*, *coul/msm*, *coul/wolf* or similar. One way to get around this would be to set *special\_bond* scaling factors to very tiny numbers that are not exactly zero (e.g.,  $1.0 \times 10^{-50}$ ). Another workaround is to write a dump file and use the *rerun* command to compute the clusters for snapshots in the dump file. The rerun script can use a *special\_bonds* command that includes all pairs in the neighbor list.

---

---

**Note:** For the compute *fragment/atom* style, each fragment is identified using the current bond topology. This will not account for bonds broken by the *bond\_style quartic* command because it does not perform a full update of the bond topology data structures within LAMMPS.

---

## 18.17.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The per-atom vector values will be an ID > 0, as explained above.

## 18.17.5 Restrictions

none

## 18.17.6 Related commands

*compute coord/atom*

### 18.17.7 Default

The default for fragment/atom is single=no.

## 18.18 compute cna/atom command

### 18.18.1 Syntax

```
compute ID group-ID cna/atom cutoff
```

- ID, group-ID are documented in *compute* command
- cna/atom = style name of this compute command
- cutoff = cutoff distance for nearest neighbors (distance units)

### 18.18.2 Examples

```
compute 1 all cna/atom 3.08
```

### 18.18.3 Description

Define a computation that calculates the CNA (Common Neighbor Analysis) pattern for each atom in the group. In solid-state systems the CNA pattern is a useful measure of the local crystal structure around an atom. The CNA methodology is described in (*Faken*) and (*Tsuzuki*).

Currently, there are five kinds of CNA patterns LAMMPS recognizes:

- fcc = 1
- hcp = 2
- bcc = 3
- icosahedral = 4
- unknown = 5

The value of the CNA pattern will be 0 for atoms not in the specified compute group. Note that normally a CNA calculation should only be performed on mono-component systems.

The CNA calculation can be sensitive to the specified cutoff value. You should insure the appropriate nearest neighbors of an atom are found within the cutoff distance for the presumed crystal structure (e.g., 12 nearest neighbor for perfect FCC and HCP crystals, 14 nearest neighbors for perfect BCC crystals). These formulas can be used to obtain a good cutoff distance:

$$r_c^{\text{fcc}} = \frac{1}{2} \left( \frac{\sqrt{2}}{2} + 1 \right) a \approx 0.8536a$$

$$r_c^{\text{bcc}} = \frac{1}{2} (\sqrt{2} + 1) a \approx 1.207a$$

$$r_c^{\text{hcp}} = \frac{1}{2} \left( 1 + \sqrt{\frac{4 + 2x^2}{3}} \right) a$$

where  $a$  is the lattice constant for the crystal structure concerned and in the HCP case,  $x = (c/a)/1.633$ , where 1.633 is the ideal  $c/a$  for HCP crystals.

Also note that since the CNA calculation in LAMMPS uses the neighbors of an owned atom to find the nearest neighbors of a ghost atom, the following relation should also be satisfied:

$$r_c + r_s > 2 * \text{cutoff}$$

where  $r_c$  is the cutoff distance of the potential,  $r_s$  is the skin distance as specified by the *neighbor* command, and cutoff is the argument used with the `compute cna/atom` command. LAMMPS will issue a warning if this is not the case.

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (e.g. each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently or to have multiple `compute/dump` commands, each with a *cna/atom* style.

### 18.18.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The per-atom vector values will be a number from 0 to 5, as explained above.

### 18.18.5 Restrictions

none

### 18.18.6 Related commands

*compute centro/atom*

### 18.18.7 Default

none

---

**(Faken)** Faken, Jonsson, *Comput Mater Sci*, 2, 279 (1994).

**(Tsuzuki)** Tsuzuki, Branicio, Rino, *Comput Phys Comm*, 177, 518 (2007).

## 18.19 compute cnp/atom command

### 18.19.1 Syntax

`compute ID group-ID cnp/atom cutoff`

- ID, group-ID are documented in *compute* command
- cnp/atom = style name of this compute command
- cutoff = cutoff distance for nearest neighbors (distance units)

## 18.19.2 Examples

```
compute 1 all cnp/atom 3.08
```

## 18.19.3 Description

Define a computation that calculates the Common Neighborhood Parameter (CNP) for each atom in the group. In solid-state systems the CNP is a useful measure of the local crystal structure around an atom and can be used to characterize whether the atom is part of a perfect lattice, a local defect (e.g., a dislocation or stacking fault), or at a surface.

The value of the CNP parameter will be 0.0 for atoms not in the specified compute group. Note that normally a CNP calculation should only be performed on single component systems.

This parameter is computed using the following formula from (*Tsuzuki*)

$$Q_i = \frac{1}{n_i} \sum_{j=1}^{n_i} \left\| \sum_{k=1}^{n_{ij}} \vec{R}_{ik} + \vec{R}_{jk} \right\|^2$$

where the index  $j$  goes over the  $n_i$  nearest neighbors of atom  $i$ , and the index  $k$  goes over the  $n_{ij}$  common nearest neighbors between atom  $i$  and atom  $j$ .  $\vec{R}_{ik}$  and  $\vec{R}_{jk}$  are the vectors connecting atom  $k$  to atoms  $i$  and  $j$ . The quantity in the double sum is computed for each atom.

The CNP calculation is sensitive to the specified cutoff value. You should ensure that the appropriate nearest neighbors of an atom are found within the cutoff distance for the presumed crystal structure. E.g. 12 nearest neighbor for perfect FCC and HCP crystals, 14 nearest neighbors for perfect BCC crystals. These formulas can be used to obtain a good cutoff distance:

$$r_c^{\text{fcc}} = \frac{1}{2} \left( \frac{\sqrt{2}}{2} + 1 \right) a \approx 0.8536a$$

$$r_c^{\text{bcc}} = \frac{1}{2} (\sqrt{2} + 1) a \approx 1.207a$$

$$r_c^{\text{hcp}} = \frac{1}{2} \left( 1 + \sqrt{\frac{4 + 2x^2}{3}} \right) a$$

where  $a$  is the lattice constant for the crystal structure concerned and in the HCP case,  $x = (c/a)/1.633$ , where 1.633 is the ideal  $c/a$  for HCP crystals.

Also note that since the CNP calculation in LAMMPS uses the neighbors of an owned atom to find the nearest neighbors of a ghost atom, the following relation should also be satisfied:

$$r_c + r_s > 2 * \text{cutoff}$$

where  $r_c$  is the cutoff distance of the potential,  $r_s$  is the skin distance as specified by the *neighbor* command, and cutoff is the argument used with the compute cnp/atom command. LAMMPS will issue a warning if this is not the case.

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (e.g., each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently or to have multiple compute/dump commands, each with a *cnp/atom* style.

### 18.19.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values will be real positive numbers. Some typical CNP values:

```
FCC lattice = 0.0
BCC lattice = 0.0
HCP lattice = 4.4

FCC (111) surface = 13.0
FCC (100) surface = 26.5
FCC dislocation core = 11
```

### 18.19.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 18.19.6 Related commands

*compute cna/atom compute centro/atom*

### 18.19.7 Default

none

---

(**Tsuzuki**) Tsuzuki, Branicio, Rino, Comput Phys Comm, 177, 518 (2007).

## 18.20 compute com command

### 18.20.1 Syntax

```
compute ID group-ID com
```

- ID, group-ID are documented in *compute* command
- com = style name of this compute command

## 18.20.2 Examples

```
compute 1 all com
```

## 18.20.3 Description

Define a computation that calculates the center-of-mass of the group of atoms, including all effects due to atoms passing through periodic boundaries.

A vector of three quantities is calculated by this compute, which are the  $(x, y, z)$  coordinates of the center of mass.

---

**Note:** The coordinates of an atom contribute to the center-of-mass in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the *set image* command.

---

## 18.20.4 Output info

This compute calculates a global vector of length 3, which can be accessed by indices 1–3 by any command that uses global vector values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The vector values are “intensive”. The vector values will be in distance *units*.

## 18.20.5 Restrictions

none

## 18.20.6 Related commands

*compute com/chunk*

## 18.20.7 Default

none

# 18.21 compute com/chunk command

## 18.21.1 Syntax

```
compute ID group-ID com/chunk chunkID
```

- ID, group-ID are documented in *compute* command
- com/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command

## 18.21.2 Examples

```
compute 1 fluid com/chunk molchunk
```

## 18.21.3 Description

Define a computation that calculates the center-of-mass for multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

This compute calculates the  $(x, y, z)$  coordinates of the center of mass for each chunk, which includes all effects due to atoms passing through periodic boundaries.

Note that only atoms in the specified group contribute to the calculation. The *compute chunk/atom* command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

---

**Note:** The coordinates of an atom contribute to the chunk’s center-of-mass in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the *set image* command.

---

The simplest way to output the results of the compute com/chunk calculation to a file is to use the *fix ave/time* command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk all com/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk[*] file tmp.out mode vector
```

## 18.21.4 Output info

This compute calculates a global array where the number of rows = the number of chunks *Nchunk* as calculated by the specified *compute chunk/atom* command. The number of columns is 3 for the  $(x, y, z)$  center-of-mass coordinates of each chunk. These values can be accessed by any command that uses global array values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The array values are “intensive”. The array values will be in distance *units*.

### 18.21.5 Restrictions

none

### 18.21.6 Related commands

*compute com*

### 18.21.7 Default

none

## 18.22 compute contact/atom command

### 18.22.1 Syntax

```
compute ID group-ID contact/atom group2-ID
```

- ID, group-ID are documented in *compute* command
- contact/atom = style name of this compute command
- group2-ID = optional argument to restrict which atoms to consider for contacts (see below)

### 18.22.2 Examples

```
compute 1 all contact/atom  
compute 1 all contact/atom mygroup
```

### 18.22.3 Description

Define a computation that calculates the number of contacts for each atom in a group.

The contact number is defined for finite-size spherical particles as the number of neighbor atoms which overlap the central particle, meaning that their distance of separation is less than or equal to the sum of the radii of the two particles.

The value of the contact number will be 0.0 for atoms not in the specified compute group.

### 18.22.4 Output info

This compute calculates a per-atom vector, whose values can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The per-atom vector values will be a number  $\geq 0.0$ , as explained above.

The optional *group2-ID* argument allows to specify from which group atoms contribute to the coordination number. Default setting is group 'all.'



### 18.22.5 Restrictions

This compute is part of the GRANULAR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This compute requires that atoms store a radius as defined by the *atom\_style sphere* command.

### 18.22.6 Related commands

*compute coord/atom*

### 18.22.7 Default

*group2-ID* = all

none

## 18.23 compute coord/atom command

Accelerator Variants: *coord/atom/kk*

### 18.23.1 Syntax

```
compute ID group-ID coord/atom style args ...
```

- ID, group-ID are documented in *compute* command
- coord/atom = style name of this compute command
- style = *cutoff* or *orientorder*

```
cutoff args = cutoff [group group2-ID] typeN
  cutoff = distance within which to count coordination neighbors (distance units)
  group group2-ID = select group-ID to restrict which atoms to consider for
  ↪coordination number (optional)
  typeN = atom type for Nth coordination count (see asterisk form below)
orientorder args = orientorderID threshold
  orientorderID = ID of an orientorder/atom compute
  threshold = minimum value of the product of two "connected" atoms
```

### 18.23.2 Examples

```
compute 1 all coord/atom cutoff 2.0
compute 1 all coord/atom cutoff 6.0 1 2
compute 1 all coord/atom cutoff 6.0 2*4 5*8 *
compute 1 solute coord/atom cutoff 2.0 group solvent
compute 1 all coord/atom orientorder 2 0.5
```

### 18.23.3 Description

This compute performs calculations between neighboring atoms to determine a coordination value. The specific calculation and the meaning of the resulting value depend on the *cstyle* keyword used.

The *cutoff* *cstyle* calculates one or more traditional coordination numbers for each atom. A coordination number is defined as the number of neighbor atoms with specified atom type(s), and optionally within the specified group, that are within the specified cutoff distance from the central atom. The compute group selects only the central atoms; all neighboring atoms, unless selected by type, type range, or group option, are included in the coordination number tally.

The optional *group* keyword allows to specify from which group atoms contribute to the coordination number. Default setting is group 'all.'

The *typeN* keywords allow specification of which atom types contribute to each coordination number. One coordination number is computed for each of the *typeN* keywords listed. If no *typeN* keywords are listed, a single coordination number is calculated, which includes atoms of all types (same as the "\*" format, see below).

The *typeN* keywords can be specified in one of two ways. An explicit numeric value can be used, as in the second example above. Or a wild-card asterisk can be used to specify a range of atom types. This takes the form "\*" or "\*n" or "m\*" or "m\*n". If *N* is the number of atom types, then an asterisk with no numeric values means all types from 1 to *N*. A leading asterisk means all types from 1 to *n* (inclusive). A trailing asterisk means all types from *m* to *N* (inclusive). A middle asterisk means all types from *m* to *n* (inclusive).

The *orientorder* *cstyle* calculates the number of "connected" neighbor atoms *j* around each central atom *i*. For this *cstyle*, connected is defined by the orientational order parameter calculated by the [compute orientorder/atom](#) command. This *cstyle* thus allows one to apply the ten Wolde's criterion to identify crystal-like atoms in a system, as discussed in [ten Wolde](#).

The ID of the previously specified [compute orientorder/atom](#) command is specified as *orientorderID*. The compute must invoke its *components* option to calculate components of the *Ybar\_lm* vector for each atoms, as described in its documentation. Note that [orientorder/atom](#) compute defines its own criteria for identifying neighboring atoms. If the scalar product ( $Ybar\_lm(i), Ybar\_lm(j)$ ), calculated by the [orientorder/atom](#) compute is larger than the specified *threshold*, then *i* and *j* are connected, and the coordination value of *i* is incremented by one.

For all *cstyle* settings, all coordination values will be 0.0 for atoms not in the specified compute group.

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (i.e., each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently.

---

**Note:** If you have a bonded system, then the settings of [special\\_bonds](#) command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the [special\\_bonds](#) command, and means those pairwise interactions do not appear in the neighbor list. Because this fix uses the neighbor list, it also means those pairs will not be included in the coordination count. One way to get around this, is to write a dump file, and use the [rerun](#) command to compute the coordination for snapshots in the dump file. The rerun script can use a [special\\_bonds](#) command that includes all pairs in the neighbor list.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* [command-line switch](#) when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 18.23.4 Output info

For *cstyle* cutoff, this compute can calculate a per-atom vector or array. If single *type1* keyword is specified (or if none are specified), this compute calculates a per-atom vector. If multiple *typeN* keywords are specified, this compute calculates a per-atom array, with *N* columns.

For *cstyle* orientorder, this compute calculates a per-atom vector.

These values can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The per-atom vector or array values will be a number  $\geq 0.0$ , as explained above.

### 18.23.5 Restrictions

none

### 18.23.6 Related commands

*compute cluster/atom compute orientorder/atom*

### 18.23.7 Default

group = all

---

(tenWolde) P. R. ten Wolde, M. J. Ruiz-Montero, D. Frenkel, J. Chem. Phys. 104, 9932 (1996).

## 18.24 compute damage/atom command

### 18.24.1 Syntax

```
compute ID group-ID damage/atom
```

- ID, group-ID are documented in *compute* command
- damage/atom = style name of this compute command

## 18.24.2 Examples

```
compute 1 all damage/atom
```

## 18.24.3 Description

Define a computation that calculates the per-atom damage for each atom in a group. This is a quantity relevant for *Peridynamics models*. See [this document](#) for an overview of LAMMPS commands for Peridynamics modeling.

The “damage” of a Peridynamics particles is based on the bond breakage between the particle and its neighbors. If all the bonds are broken the particle is considered to be fully damaged.

See the [PDLAMMPS user guide](#) for a formal definition of “damage” and more details about Peridynamics as it is implemented in LAMMPS.

This command can be used with all the Peridynamic pair styles.

The damage value will be 0.0 for atoms not in the specified compute group.

## 18.24.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values are unitless numbers (damage)  $\geq 0.0$ .

## 18.24.5 Restrictions

This compute is part of the PERI package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 18.24.6 Related commands

*compute dilatation/atom, compute plasticity/atom*

## 18.24.7 Default

none

# 18.25 compute dihedral command

## 18.25.1 Syntax

```
compute ID group-ID dihedral
```

- ID, group-ID are documented in *compute* command
- dihedral = style name of this compute command

## 18.25.2 Examples

```
compute 1 all dihedral
```

## 18.25.3 Description

Define a computation that extracts the dihedral energy calculated by each of the dihedral sub-styles used in the *dihedral\_style hybrid* command. These values are made accessible for output or further processing by other commands. The group specified for this command is ignored.

This compute is useful when using *dihedral\_style hybrid* if you want to know the portion of the total energy contributed by one or more of the hybrid sub-styles.

## 18.25.4 Output info

This compute calculates a global vector of length  $N$ , where  $N$  is the number of sub\_styles defined by the *dihedral\_style hybrid* command, which can be accessed by the indices 1 through  $N$ . These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector values are “extensive” and will be in energy *units*.

## 18.25.5 Restrictions

none

## 18.25.6 Related commands

*compute pe*, *compute pair*

## 18.25.7 Default

none

## 18.26 compute dihedral/local command

### 18.26.1 Syntax

```
compute ID group-ID dihedral/local value1 value2 ... keyword args ...
```

- ID, group-ID are documented in *compute* command
- dihedral/local = style name of this compute command
- one or more values may be appended
- value = *phi* or *v\_name*
  - phi* = tabulate dihedral angles
  - v\_name* = equal-style variable with name (see below)

- zero or more keyword/args pairs may be appended
- keyword = *set*  

```
set args = phi name  
  phi = only currently allowed arg  
  name = name of variable to set with phi
```

## 18.26.2 Examples

```
compute 1 all dihedral/local phi  
compute 1 all dihedral/local phi v_cos set phi p
```

## 18.26.3 Description

Define a computation that calculates properties of individual dihedral interactions. The number of datums generated, aggregated across all processors, equals the number of dihedral angles in the system, modified by the group parameter as explained below.

The value *phi* ( $\phi$ ) is the dihedral angle, as defined in the diagram on the *dihedral\_style* doc page.

The value *v\_name* can be used together with the *set* keyword to compute a user-specified function of the dihedral angle  $\phi$ . The *name* specified for the *v\_name* value is the name of an *equal-style variable* which should evaluate a formula based on a variable which will store the angle  $\phi$ . This other variable must be an *internal-style variable* defined in the input script; its initial numeric value can be anything. It must be an internal-style variable, because this command resets its value directly. The *set* keyword is used to identify the name of this other variable associated with  $\phi$ .

Note that the value of  $\phi$  for each angle which stored in the internal variable is in radians, not degrees.

As an example, these commands can be added to the bench/in.rhodo script to compute the  $\cos \phi$  and  $\cos^2 \phi$  of every dihedral angle in the system and output the statistics in various ways:

```
variable p internal 0.0  
variable cos equal cos(v_p)  
variable cossq equal cos(v_p)*cos(v_p)  
  
compute 1 all property/local datum1 datum2 datum3 datum4 dtype  
compute 2 all dihedral/local phi v_cos v_cossq set phi p  
dump 1 all local 100 tmp.dump c_1[*] c_2[*]  
  
compute 3 all reduce ave c_2[*]  
thermo_style custom step temp press c_3[*]  
  
fix 10 all ave/histo 10 10 100 -1 1 20 c_2[2] mode vector file tmp.histo
```

The *dump local* command will output the angle ( $\phi$ ),  $\cos(\phi)$ , and  $\cos^2(\phi)$  for every dihedral in the system. The *thermo\_style* command will print the average of those quantities via the *compute reduce* command with thermo output. And the *fix ave/histo* command will histogram the cosine(angle) values and write them to a file.

---

The local data stored by this command is generated by looping over all the atoms owned on a processor and their dihedrals. A dihedral will only be included if all four atoms in the dihedral are in the specified compute group.

Note that as atoms migrate from processor to processor, there will be no consistent ordering of the entries within the local vector or array from one timestep to the next. The only consistency that is guaranteed is that the ordering on a

particular timestep will be the same for local vectors or arrays generated by other compute commands. For example, dihedral output from the *compute property/local* command can be combined with data from this command and output by the *dump local* command in a consistent way.

Here is an example of how to do this:

```
compute 1 all property/local dtype datom1 datom2 datom3 datom4
compute 2 all dihedral/local phi
dump 1 all local 1000 tmp.dump index c_1[1] c_1[2] c_1[3] c_1[4] c_1[5] c_2[1]
```

## 18.26.4 Output info

This compute calculates a local vector or local array depending on the number of values. The length of the vector or number of rows in the array is the number of dihedrals. If a single value is specified, a local vector is produced. If two or more values are specified, a local array is produced where the number of columns is equal to the number of values. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The output for *phi* will be in degrees.

## 18.26.5 Restrictions

none

## 18.26.6 Related commands

*dump local*, *compute property/local*

## 18.26.7 Default

none

## 18.27 compute dilatation/atom command

### 18.27.1 Syntax

```
compute ID group-ID dilatation/atom
```

- ID, group-ID are documented in compute command
- dilatation/atom = style name of this compute command

## 18.27.2 Examples

```
compute 1 all dilatation/atom
```

## 18.27.3 Description

Define a computation that calculates the per-atom dilatation for each atom in a group. This is a quantity relevant for *Peridynamics models*. See [this document](#) for an overview of LAMMPS commands for Peridynamics modeling.

For small deformation, dilatation of is the measure of the volumetric strain.

The dilatation  $\theta$  for each peridynamic particle  $i$  is calculated as a sum over its neighbors with unbroken bonds, where the contribution of the  $ij$  pair is a function of the change in bond length (versus the initial length in the reference state), the volume fraction of the particles and an influence function. See the [PDLAMMPS user guide](#) for a formal definition of dilatation.

This command can only be used with a subset of the Peridynamic *pair styles*: peri/lps, peri/ves and peri/eps.

The dilatation value will be 0.0 for atoms not in the specified compute group.

## 18.27.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values are unitless numbers ( $\theta \geq 0.0$ ).

## 18.27.5 Restrictions

This compute is part of the PERI package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 18.27.6 Related commands

*compute damage/atom, compute plasticity/atom*

## 18.27.7 Default

none

# 18.28 compute dipole command

## 18.28.1 Syntax

```
compute ID group-ID dipole arg
```

- ID, group-ID are documented in *compute* command
- dipole = style name of this compute command



- `arg = mass` or `geometry` = use COM or geometric center for charged chunk correction (optional)

## 18.28.2 Examples

```
compute 1 fluid dipole
compute dw water dipole geometry
```

## 18.28.3 Description

Define a computation that calculates the dipole vector and total dipole for a group of atoms.

This compute calculates the  $x,y,z$  coordinates of the dipole vector and the total dipole moment for the atoms in the compute group. This includes all effects due to atoms passing through periodic boundaries. For a group with a net charge the resulting dipole is made position independent by subtracting the position vector of the center of mass or geometric center times the net charge from the computed dipole vector. Both per-atom charges and per-atom dipole moments, if present, contribute to the computed dipole.

---

**Note:** The coordinates of an atom contribute to the dipole in “unwrapped” form, by using the image flags associated with each atom. See the `dump custom` command for a discussion of “unwrapped” coordinates. See the Atoms section of the `read_data` command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the `set image` command.

---

## 18.28.4 Output info

This compute calculations a global scalar containing the magnitude of the computed dipole moment and a global vector of length 3 with the dipole vector. See the `Howto output` page for an overview of LAMMPS output options.

The computed values are “intensive”. The array values will be in dipole units (i.e., charge *units* times distance *units*).

## 18.28.5 Restrictions

none

## 18.28.6 Related commands

`compute dipole/chunk`

## 18.28.7 Default

Using the center of mass is the default setting for the net charge correction.

## 18.29 compute dipole/chunk command

### 18.29.1 Syntax

```
compute ID group-ID dipole/chunk chunkID arg
```

- ID, group-ID are documented in *compute* command
- dipole/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command
- arg = *mass* or *geometry* = use COM or geometric center for charged chunk correction (optional)

### 18.29.2 Examples

```
compute 1 fluid dipole/chunk molchunk  
compute dw water dipole/chunk 1 geometry
```

### 18.29.3 Description

Define a computation that calculates the dipole vector and total dipole for multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

This compute calculates the  $(x, y, z)$  coordinates of the dipole vector and the total dipole moment for each chunk, which includes all effects due to atoms passing through periodic boundaries. For chunks with a net charge the resulting dipole is made position independent by subtracting the position vector of the center of mass or geometric center times the net charge from the computed dipole vector. Both per-atom charges and per-atom dipole moments, if present, contribute to the computed dipole.

Note that only atoms in the specified group contribute to the calculation. The *compute chunk/atom* command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

---

**Note:** The coordinates of an atom contribute to the chunk’s dipole in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the *set image* command.

---

The simplest way to output the results of the compute com/chunk calculation to a file is to use the *fix ave/time* command, for example:

```
compute cc1 all chunk/atom molecule  
compute myChunk all dipole/chunk cc1  
fix 1 all ave/time 100 1 100 c_myChunk[*] file tmp.out mode vector
```

## 18.29.4 Output info

This compute calculates a global array where the number of rows = the number of chunks *Nchunk* as calculated by the specified `compute chunk/atom` command. The number of columns is 4 for the  $(x, y, z)$  dipole vector components and the total dipole of each chunk. These values can be accessed by any command that uses global array values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The array values are “intensive”. The array values will be in dipole units (i.e., charge *units* times distance *units*).

## 18.29.5 Restrictions

none

## 18.29.6 Related commands

`compute com/chunk`, `compute dipole`

## 18.29.7 Default

Using the center of mass is the default setting for the net charge correction.

# 18.30 compute displace/atom command

## 18.30.1 Syntax

```
compute ID group-ID displace/atom
```

- ID, group-ID are documented in `compute` command
- displace/atom = style name of this compute command
- zero or more keyword/arg pairs may be appended
- keyword = `refresh`  
     *replace arg* = name of per-atom variable

## 18.30.2 Examples

```
compute 1 all displace/atom
compute 1 all displace/atom refresh myVar
```

### 18.30.3 Description

Define a computation that calculates the current displacement of each atom in the group from its original (reference) coordinates, including all effects due to atoms passing through periodic boundaries.

A vector of four quantities per atom is calculated by this compute. The first three elements of the vector are the  $(dx, dy, dz)$  displacements. The fourth component is the total displacement (i.e.,  $\sqrt{dx^2 + dy^2 + dz^2}$ ).

The displacement of an atom is from its original position at the time the compute command was issued. The value of the displacement will be 0.0 for atoms not in the specified compute group.

---

**Note:** Initial coordinates are stored in “unwrapped” form, by using the image flags associated with each atom. See the `dump custom` command for a discussion of “unwrapped” coordinates. See the Atoms section of the `read_data` command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the `set image` command.

---



---

**Note:** If you want the quantities calculated by this compute to be continuous when running from a `restart file`, then you should use the same ID for this compute, as in the original run. This is so that the fix this compute creates to store per-atom quantities will also have the same ID, and thus be initialized correctly with time = 0 atom coordinates from the restart file.

---

The `refresh` option can be used in conjunction with the “`dump_modify refresh`” command to generate incremental dump files.

The definition and motivation of an incremental dump file is as follows. Instead of outputting all atoms at each snapshot (with some associated values), you may only wish to output the subset of atoms with a value that has changed in some way compared to the value the last time that atom was output. In some scenarios this can result in a dramatically smaller dump file. If desired, by post-processing the sequence of snapshots, the values for all atoms at all timesteps can be inferred.

A concrete example using this compute, is a simulation of atom diffusion in a solid, represented as atoms on a lattice. Diffusive hops are rare. Imagine that when a hop occurs an atom moves more than a distance  $D_{hop}$ . For any snapshot we only want to output atoms that have hopped since the last snapshot. This can be accomplished with something like the following commands:

```
write_dump      all custom tmp.dump id type x y z      # see comment below

variable       Dhop equal 0.6
variable       check atom "c_dsp[4] > v_Dhop"
compute       dsp all displace/atom refresh check
dump          1 all custom 100 tmp.dump id type x y z
dump_modify   1 append yes thresh c_dsp[4] > ${Dhop} &
              refresh c_dsp delay 100
```

The `dump_modify thresh` command will only output atoms that have displaced more than 0.6 Å on each snapshot (assuming metal units). The `dump_modify refresh` option triggers a call to this compute at the end of every dump.

The `refresh` argument for this compute is the ID of an *atom-style variable* which calculates a Boolean value (0 or 1) based on the same criterion used by `dump_modify thresh`. This compute evaluates the atom-style variable. For each atom that returns 1 (true), the original (reference) coordinates of the atom (stored by this compute) are updated.

The effect of these commands is that a particular atom will only be output in the dump file on the snapshot after it makes a diffusive hop. It will not be output again until it makes another hop.

Note that in the first snapshot of a subsequent run, no atoms will be typically be output. That is because the initial displacement for all atoms is 0.0. If an initial dump snapshot is desired, containing the initial reference positions of all atoms, one way to do this is illustrated above. An initial `write_dump` command can be used before the first run. It will contain the positions of all the atoms, Options in the `dump_modify` command above will append new output to that same file and delay the output until a later timestep. The `delay` setting avoids a second time = 0 snapshot which would be empty.

---

### 18.30.4 Output info

This compute calculates a per-atom array with four columns, which can be accessed by indices 1–4 by any command that uses per-atom values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The per-atom array values will be in distance *units*.

This compute supports the `refresh` option as explained above, for use in conjunction with `dump_modify refresh` to generate incremental dump files.

### 18.30.5 Restrictions

none

### 18.30.6 Related commands

*compute msd, dump custom, fix store/state*

### 18.30.7 Default

none

## 18.31 compute dpd command

### 18.31.1 Syntax

```
compute ID group-ID dpd
```

- ID, group-ID are documented in *compute* command
- dpd = style name of this compute command

## 18.31.2 Examples

```
compute 1 all dpd
```

---

## 18.31.3 Description

Define a computation that accumulates the total internal conductive energy ( $U^{\text{cond}}$ ), the total internal mechanical energy ( $U^{\text{mech}}$ ), the total chemical energy ( $U^{\text{chem}}$ ) and the *harmonic* average of the internal temperature ( $\theta_{\text{avg}}$ ) for the entire system of particles. See the `compute dpd/atom` command if you want per-particle internal energies and internal temperatures.

The system internal properties are computed according to the following relations:

$$U^{\text{cond}} = \sum_{i=1}^N u_i^{\text{cond}}$$
$$U^{\text{mech}} = \sum_{i=1}^N u_i^{\text{mech}}$$
$$U^{\text{chem}} = \sum_{i=1}^N u_i^{\text{chem}}$$
$$U = \sum_{i=1}^N (u_i^{\text{cond}} + u_i^{\text{mech}} + u_i^{\text{chem}})$$
$$\theta_{\text{avg}} = \left( \frac{1}{N} \sum_{i=1}^N \frac{1}{\theta_i} \right)^{-1}$$

where  $N$  is the number of particles in the system.

---

## 18.31.4 Output info

This compute calculates a global vector of length 5 ( $U^{\text{cond}}$ ,  $U^{\text{mech}}$ ,  $U^{\text{chem}}$ ,  $\theta_{\text{avg}}$ ,  $N$ ), which can be accessed by indices 1 through 5. See the [Howto output](#) page for an overview of LAMMPS output options.

The vector values will be in energy and temperature *units*.

## 18.31.5 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This command also requires use of the `atom_style dpd` command.

## 18.31.6 Related commands

*compute dpd/atom*, *thermo\_style*

## 18.31.7 Default

none

---

**(Larentzos)** J.P. Larentzos, J.K. Brennan, J.D. Moore, and W.D. Mattson, “LAMMPS Implementation of Constant Energy Dissipative Particle Dynamics (DPD-E)”, ARL-TR-6863, U.S. Army Research Laboratory, Aberdeen Proving Ground, MD (2014).

## 18.32 compute dpd/atom command

### 18.32.1 Syntax

```
compute ID group-ID dpd/atom
```

- ID, group-ID are documented in *compute* command
- dpd/atom = style name of this compute command

### 18.32.2 Examples

```
compute 1 all dpd/atom
```

### 18.32.3 Description

Define a computation that accesses the per-particle internal conductive energy ( $u^{\text{cond}}$ ), internal mechanical energy ( $u^{\text{mech}}$ ), internal chemical energy ( $u^{\text{chem}}$ ) and internal temperatures ( $\theta$ ) for each particle in a group. See the *compute dpd* command if you want the total internal conductive energy, the total internal mechanical energy, the total chemical energy and average internal temperature of the entire system or group of dpd particles.

### 18.32.4 Output info

This compute calculates a per-particle array with four columns ( $u^{\text{cond}}$ ,  $u^{\text{mech}}$ ,  $u^{\text{chem}}$ ,  $\theta$ ), which can be accessed by indices 1–4 by any command that uses per-particle values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The per-particle array values will be in energy ( $u^{\text{cond}}$ ,  $u^{\text{mech}}$ ,  $u^{\text{chem}}$ ) and temperature ( $\theta$ ) *units*.

### 18.32.5 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This command also requires use of the *atom\_style dpd* command.

### 18.32.6 Related commands

*dump custom*, *compute dpd*

### 18.32.7 Default

none

---

**(Larentzos)** J.P. Larentzos, J.K. Brennan, J.D. Moore, and W.D. Mattson, “LAMMPS Implementation of Constant Energy Dissipative Particle Dynamics (DPD-E)”, ARL-TR-6863, U.S. Army Research Laboratory, Aberdeen Proving Ground, MD (2014).

## 18.33 compute edpd/temp/atom command

### 18.33.1 Syntax

```
compute ID group-ID edpd/temp/atom
```

- ID, group-ID are documented in *compute* command
- edpd/temp/atom = style name of this compute command

### 18.33.2 Examples

```
compute 1 all edpd/temp/atom
```

### 18.33.3 Description

Define a computation that calculates the per-atom temperature for each eDPD particle in a group.

The temperature is a local temperature derived from the internal energy of each eDPD particle based on the local equilibrium hypothesis. For more details please see (*Espanol1997*) and (*Li2014*).



### 18.33.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values will be in temperature *units*.

### 18.33.5 Restrictions

This compute is part of the DPD-MESO package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 18.33.6 Related commands

*pair\_style edpd*

### 18.33.7 Default

none

---

(Espanol1997) Espanol, Europhys Lett, 40(6): 631-636 (1997). DOI: 10.1209/epl/i1997-00515-8

(Li2014) Li, Tang, Lei, Caswell, Karniadakis, J Comput Phys, 265: 113-127 (2014). DOI: 10.1016/j.jcp.2014.02.003.

## 18.34 compute efield/atom command

### 18.34.1 Syntax

```
compute ID group-ID efield/atom keyword val
```

- ID, group-ID are documented in [compute](#) command
- efield/atom = style name of this compute command
- zero or more keyword/value pairs may be appended
- keyword = *pair* or *kpace*
  - pair* args = *yes* or *no*
  - kpace* args = *yes* or *no*

### 18.34.2 Examples

```
compute 1 all efield/atom
compute 1 all efield/atom pair yes kspace no
```

Used in input scripts:

```
examples/PACKAGES/dielectric/in.confined
examples/PACKAGES/dielectric/in.nopbc
```

### 18.34.3 Description

Define a computation that calculates the electric field at each atom in a group. The compute should only be enabled with *pair* and *kspace* styles that are provided by the DIELECTRIC package because only these styles compute the per-atom electric field at every time step.

The electric field is a 3-component vector. The value of the electric field components will be 0.0 for atoms not in the specified compute group.

---

The keyword/value option pairs are used in the following ways.

For the *pair* and *kspace* keywords, the real-space and reciprocal-space contributions to the electric field can be turned off and on.

### 18.34.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The per-atom vector values will be in electric field *units*.

### 18.34.5 Restrictions

This compute is part of the DIELECTRIC package. It is only enabled if LAMMPS was built with that package.

### 18.34.6 Related commands

*dump custom*

### 18.34.7 Default

The option defaults are *pair* = yes and *kspace* = yes.

## 18.35 compute entropy/atom command

### 18.35.1 Syntax

```
compute ID group-ID entropy/atom sigma cutoff keyword value ...
```

- ID, group-ID are documented in *compute* command
- entropy/atom = style name of this compute command
- sigma = width of Gaussians used in the  $g(r)$  smoothing
- cutoff = cutoff for the  $g(r)$  calculation
- one or more keyword/value pairs may be appended

keyword = *avg* or *local*  
*avg* args = *neigh cutoff2*  
     *neigh value* = *yes* or *no* = whether to average the pair entropy over neighbors  
     *cutoff2* = cutoff for the averaging over neighbors  
*local* arg = *yes* or *no* = use the local density around each atom to normalize the  $g(r)$

## 18.35.2 Examples

```
compute 1 all entropy/atom 0.25 5.
compute 1 all entropy/atom 0.25 5. avg yes 5.
compute 1 all entropy/atom 0.125 7.3 avg yes 5.1 local yes
```

## 18.35.3 Description

Define a computation that calculates the pair entropy fingerprint for each atom in the group. The fingerprint is useful to distinguish between ordered and disordered environments, for instance liquid and solid-like environments, or glassy and crystalline-like environments. Some applications could be the identification of grain boundaries, a melt-solid interface, or a solid cluster emerging from the melt. The advantage of this parameter over others is that no a priori information about the solid structure is required.

This parameter for atom  $i$  is computed using the following formula from (*Piaggi*) and (*Nettleton*),

$$s_S^i = -2\pi\rho k_B \int_0^{r_m} [g(r) \ln g(r) - g(r) + 1] r^2 dr$$

where  $r$  is a distance,  $g(r)$  is the radial distribution function of atom  $i$ , and  $\rho$  is the density of the system. The  $g(r)$  computed for each atom  $i$  can be noisy and therefore it is smoothed using

$$g_m^i(r) = \frac{1}{4\pi\rho r^2} \sum_j \frac{1}{\sqrt{2\pi\sigma^2}} e^{-(r-r_{ij})^2/(2\sigma^2)}$$

where the sum over  $j$  goes through the neighbors of atom  $i$  and  $\sigma$  is a parameter to control the smoothing.

The input parameters are *sigma* the smoothing parameter  $\sigma$ , and the *cutoff* for the calculation of  $g(r)$ .

If the keyword *avg* has the setting *yes*, then this compute also averages the parameter over the neighbors of atom  $i$  according to

$$\langle s_S^i \rangle = \frac{\sum_j s_S^j + s_S^i}{N + 1},$$

where the sum over  $j$  goes over the neighbors of atom  $i$  and  $N$  is the number of neighbors. This procedure provides a sharper distinction between order and disorder environments. In this case the input parameter *cutoff2* is the cutoff for the averaging over the neighbors and must also be specified.

If the *avg yes* option is used, the effective cutoff of the neighbor list should be *cutoff+cutoff2* and therefore it might be necessary to increase the skin of the neighbor list with:

```
neighbor <skin distance> bin
```

See *neighbor* for details.

If the *local yes* option is used, the  $g(r)$  is normalized by the local density around each atom, that is to say the density around each atom is the number of neighbors within the neighbor list cutoff divided by the corresponding volume. This option can be useful when dealing with inhomogeneous systems such as those that have surfaces.

Here are typical input parameters for fcc aluminum (lattice constant 4.05 Å),

```
compute 1 all entropy/atom 0.25 5.7 avg yes 3.7
```

and for bcc sodium (lattice constant 4.23 Angstroms),

```
compute 1 all entropy/atom 0.25 7.3 avg yes 5.1
```

### 18.35.4 Output info

By default, this compute calculates the pair entropy value for each atom as a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The pair entropy values have units of the Boltzmann constant. They are always negative, and lower values (lower entropy) correspond to more ordered environments.

### 18.35.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 18.35.6 Related commands

*compute cna/atom compute centro/atom*

### 18.35.7 Default

The default values for the optional keywords are avg = no and local = no.

---

**(Piaggi)** Piaggi and Parrinello, J Chem Phys, 147, 114112 (2017).

**(Nettleton)** Nettleton and Green, J Chem Phys, 29, 6 (1958).

## 18.36 compute erotate/asphere command

### 18.36.1 Syntax

```
compute ID group-ID erotate/asphere
```

- ID, group-ID are documented in *compute* command
- erotate/asphere = style name of this compute command

## 18.36.2 Examples

```
compute 1 all erotate/sphere
```

## 18.36.3 Description

Define a computation that calculates the rotational kinetic energy of a group of aspherical particles. The aspherical particles can be ellipsoids, or line segments, or triangles. See the *atom\_style* and *read\_data* commands for descriptions of these options.

For all 3 types of particles, the rotational kinetic energy is computed as  $\frac{1}{2}I\omega^2$ , where  $I$  is the inertia tensor for the aspherical particle and  $\omega$  is its angular velocity, which is computed from its angular momentum if needed.

---

**Note:** For *2d models*, ellipsoidal particles are treated as ellipsoids, not ellipses, meaning their moments of inertia will be the same as in 3d.

---

## 18.36.4 Output info

This compute calculates a global scalar (the KE). This value can be used by any command that uses a global scalar value from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “extensive”. The scalar value will be in energy *units*.

## 18.36.5 Restrictions

This compute requires that ellipsoidal particles atoms store a shape and quaternion orientation and angular momentum as defined by the *atom\_style ellipsoid* command.

This compute requires that line segment particles atoms store a length and orientation and angular velocity as defined by the *atom\_style line* command.

This compute requires that triangular particles atoms store a size and shape and quaternion orientation and angular momentum as defined by the *atom\_style tri* command.

All particles in the group must be of finite size. They cannot be point particles.

## 18.36.6 Related commands

none

*compute erotate/sphere*

### 18.36.7 Default

none

## 18.37 compute erotate/rigid command

### 18.37.1 Syntax

```
compute ID group-ID erotate/rigid fix-ID
```

- ID, group-ID are documented in *compute* command
- erotate/rigid = style name of this compute command
- fix-ID = ID of rigid body fix

### 18.37.2 Examples

```
compute 1 all erotate/rigid myRigid
```

### 18.37.3 Description

Define a computation that calculates the rotational kinetic energy of a collection of rigid bodies, as defined by one of the *fix rigid* command variants.

The rotational energy of each rigid body is computed as  $\frac{1}{2}I\omega_{\text{body}}^2$ , where  $I$  is the inertia tensor for the rigid body and  $\omega_{\text{body}}$  is its angular velocity vector. Both  $I$  and  $\omega_{\text{body}}$  are in the frame of reference of the rigid body (i.e.,  $I$  is diagonal).

The *fix-ID* should be the ID of one of the *fix rigid* commands which defines the rigid bodies. The group specified in the compute command is ignored. The rotational energy of all the rigid bodies defined by the fix rigid command is included in the calculation.

### 18.37.4 Output info

This compute calculates a global scalar (the summed rotational energy of all the rigid bodies). This value can be used by any command that uses a global scalar value from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “extensive”. The scalar value will be in energy *units*.

### 18.37.5 Restrictions

This compute is part of the RIGID package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 18.37.6 Related commands

*compute ke/rigid*

## 18.37.7 Default

none

# 18.38 compute erotate/sphere command

## 18.38.1 Syntax

```
compute ID group-ID erotate/sphere
```

- ID, group-ID are documented in *compute* command
- erotate/sphere = style name of this compute command

## 18.38.2 Examples

```
compute 1 all erotate/sphere
```

## 18.38.3 Description

Define a computation that calculates the rotational kinetic energy of a group of spherical particles.

The rotational energy is computed as  $\frac{1}{2}I\omega^2$ , where  $I$  is the moment of inertia for a sphere and  $\omega$  is the particle's angular velocity.

---

**Note:** For *2d models*, particles are treated as spheres, not disks, meaning their moment of inertia will be the same as in 3d.

---

## 18.38.4 Output info

This compute calculates a global scalar (the KE). This value can be used by any command that uses a global scalar value from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “extensive”. The scalar value will be in energy *units*.

### 18.38.5 Restrictions

This compute requires that atoms store a radius and angular velocity (omega) as defined by the *atom\_style sphere* command.

All particles in the group must be finite-size spheres or point particles. They cannot be aspherical. Point particles will not contribute to the rotational energy.

### 18.38.6 Related commands

*compute erotate/sphere*

### 18.38.7 Default

none

## 18.39 compute erotate/sphere/atom command

### 18.39.1 Syntax

```
compute ID group-ID erotate/sphere/atom
```

- ID, group-ID are documented in *compute* command
- erotate/sphere/atom = style name of this compute command

### 18.39.2 Examples

```
compute 1 all erotate/sphere/atom
```

### 18.39.3 Description

Define a computation that calculates the rotational kinetic energy for each particle in a group.

The rotational energy is computed as  $\frac{1}{2}I\omega^2$ , where  $I$  is the moment of inertia for a sphere and  $\omega$  is the particle's angular velocity.

---

**Note:** For *2d models*, particles are treated as spheres, not disks, meaning their moment of inertia will be the same as in 3d.

---

The value of the rotational kinetic energy will be 0.0 for atoms not in the specified compute group or for point particles with a radius of 0.0.



### 18.39.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The per-atom vector values will be in energy *units*.

### 18.39.5 Restrictions

none

### 18.39.6 Related commands

*dump custom*

### 18.39.7 Default

none

## 18.40 compute event/displace command

### 18.40.1 Syntax

```
compute ID group-ID event/displace threshold
```

- ID, group-ID are documented in *compute* command
- event/displace = style name of this compute command
- threshold = minimum distance any particle must move to trigger an event (distance units)

### 18.40.2 Examples

```
compute 1 all event/displace 0.5
```

### 18.40.3 Description

Define a computation that flags an “event” if any particle in the group has moved a distance greater than the specified threshold distance when compared to a previously stored reference state (i.e., the previous event). This compute is typically used in conjunction with the *prd* and *tad* commands, to detect if a transition to a new minimum energy basin has occurred.

This value calculated by the compute is equal to 0 if no particle has moved far enough, and equal to 1 if one or more particles have moved further than the threshold distance.

**Note:** If the system is undergoing significant center-of-mass motion, due to thermal motion, an external force, or an initial net momentum, then this compute will not be able to distinguish that motion from local atom displacements and may generate “false positives”.

---

### 18.40.4 Output info

This compute calculates a global scalar (the flag). This value can be used by any command that uses a global scalar value from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The scalar value will be a 0 or 1 as explained above.

### 18.40.5 Restrictions

This command can only be used if LAMMPS was built with the REPLICA package. See the *Build package* doc page for more info.

### 18.40.6 Related commands

*prd, tad*

### 18.40.7 Default

none

## 18.41 compute fabric command

### 18.41.1 Syntax

```
compute ID group-ID fabric cutoff attribute ... keyword values ...
```

- ID, group-ID are documented in *compute* command
- fabric = style name of this compute command
- cutoff = *type* or *radius*
  - type* = cutoffs determined based on atom types
  - radius* = cutoffs determined based on atom diameters (atom style sphere)
- one or more attributes may be appended
- attribute = *contact* or *branch* or *force/normal* or *force/tangential*
  - contact* = contact tensor
  - branch* = branch tensor
  - force/normal* = normal force tensor
  - force/tangential* = tangential force tensor
- zero or more keyword/value pairs may be appended
- keyword = *type/include*

```
type/include value = arg1 arg2
arg = separate lists of types (see below)
```

## 18.41.2 Examples

```
compute 1 all fabric type contact force/normal type/include 1,2 3*4
compute 1 all fabric radius force/normal force/tangential
```

## 18.41.3 Description

Define a compute that calculates various fabric tensors for pairwise interaction (*Ouadfel*). Fabric tensors are commonly used to quantify the anisotropy or orientation of granular contacts but can also be used to characterize the direction of pairwise interactions in general systems. The *type* and *radius* settings are used to select whether interactions cutoffs are determined by atom types or by the sum of atomic radii (atom style sphere), respectively. Calling this compute is roughly the cost of a pair style invocation as it involves a loop over the neighbor list. If the normal or tangential force tensors are requested, it will be more expensive than a pair style invocation as it will also recalculate all pair forces.

Four fabric tensors are available: the contact, branch, normal force, or tangential force tensor. The contact tensor is calculated as

$$C_{ab} = \frac{15}{2}(\phi_{ab} - \text{Tr}(\phi)\delta_{ab})$$

where  $a$  and  $b$  are the  $x, y, z$  directions,  $\delta_{ab}$  is the Kronecker delta function, and the tensor  $\phi$  is defined as

$$\phi_{ab} = \sum_{n=1}^{N_p} \frac{r_a r_b}{r^2}$$

where  $n$  loops over the  $N_p$  pair interactions in the simulation,  $r_a$  is the  $a$  component of the radial vector between the two pairwise interacting particles, and  $r$  is the magnitude of the radial vector.

The branch tensor is calculated as

$$B_{ab} = \frac{15}{6\text{Tr}(D)}(D_{ab} - \text{Tr}(D)\delta_{ab})$$

where the tensor  $D$  is defined as

$$D_{ab} = \sum_{n=1}^{N_p} \frac{1}{N_c(r^2 + C_{cd}r_c r_d)} \frac{r_a r_b}{r}$$

where  $N_c$  is the total number of contacts in the system and the subscripts  $c$  and  $d$  indices are summed according to Einstein notation.

The normal force fabric tensor is calculated as

$$F_{ab}^n = \frac{15}{6\text{Tr}(N)}(N_{ab} - \text{Tr}(N)\delta_{ab})$$

where the tensor  $N$  is defined as

$$N_{ab} = \sum_{n=1}^{N_p} \frac{1}{N_c(r^2 + C_{cd}r_c r_d)} \frac{r_a r_b}{r^2} f_n$$

and  $f_n$  is the magnitude of the normal, central-body force between the two atoms.

Finally, the tangential force fabric tensor is only defined for pair styles that apply tangential forces to particles, namely granular pair styles. It is calculated as

$$F_{ab}^t = \frac{15}{9 \text{Tr}(N)} (T_{ab} - \text{Tr}(T)\delta_{ab})$$

where the tensor  $T$  is defined as

$$T_{ab} = \sum_{n=1}^{N_p} \frac{1}{N_c(r^2 + C_{cd}r_c r_d)} \frac{r_a r_b}{r^2} f_t$$

and  $f_t$  is the magnitude of the tangential force between the two atoms.

The *type/include* keyword filters interactions based on the types of the two atoms. Interactions between two atoms are only included in calculations if the atom types are in the two lists. Each list consists of a series of type ranges separated by commas. The range can be specified as a single numeric value, or a wildcard asterisk can be used to specify a range of values. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. For example, if  $M$  is the number of atom types, then an asterisk with no numeric values means all types from 1 to  $M$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $m$  to  $M$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive). Multiple *type/include* keywords may be added.

#### 18.41.4 Output info

This compute calculates a local vector of doubles and a scalar. The vector stores the unique components of the first requested tensor in the order  $xx, yy, zz, xy, xz, yz$  followed by the same components for all subsequent tensors. The length of the vector is therefore six times the number of requested tensors. The scalar output is the number of pairwise interactions included in the calculation of the fabric tensor.

#### 18.41.5 Restrictions

This fix is part of the GRANULAR package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

Currently, compute *fabric* does not support pair styles with many-body interactions. It also does not support models with long-range Coulombic or dispersion forces, i.e. the `kspace_style` command in LAMMPS. It also does not support the following fixes which add rigid-body constraints: *fix shake*, *fix rattle*, *fix rigid*, *fix rigid/small*. It does not support granular pair styles that extend beyond the contact of atomic radii (e.g., JKR and DMT).

#### 18.41.6 Related commands

none

#### 18.41.7 Default

none

---

**(Quadfel)** Quadfel and Rothenburg “Stress-force-fabric relationship for assemblies of ellipsoids”, *Mechanics of Materials* (2001). ([link to paper](#))

## 18.42 compute fep command

### 18.42.1 Syntax

```
compute ID group-ID fep temp attribute args ... keyword value ...
```

- ID, group-ID are documented in the *compute* command
- fep = name of this compute command
- temp = external temperature (as specified for constant-temperature run)
- one or more attributes with args may be appended
- attribute = *pair* or *atom*

```
pair args = pstyle pparam I J v_delta
pstyle = pair style name (e.g., lj/cut)
pparam = parameter to perturb
I,J = type pair(s) to set parameter for
v_delta = variable with perturbation to apply (in the units of the parameter)
atom args = aparam I v_delta
aparam = charge = parameter to perturb
I = type to set parameter for
v_delta = variable with perturbation to apply (in the units of the parameter)
```

- zero or more keyword/value pairs may be appended
- keyword = *tail* or *volume*

```
tail value = no or yes
no = ignore tail correction to pair energies (usually small in fep)
yes = include tail correction to pair energies
volume value = no or yes
no = ignore volume changes (e.g., in NVE or NVT trajectories)
yes = include volume changes (e.g., in NPT trajectories)
```

### 18.42.2 Examples

```
compute 1 all fep 298 pair lj/cut epsilon 1 * v_delta pair lj/cut sigma 1 * v_delta
↪ volume yes
compute 1 all fep 300 atom charge 2 v_delta
```

Example input scripts available: examples/PACKAGES/fep

### 18.42.3 Description

Apply a perturbation to parameters of the interaction potential and recalculate the pair potential energy without changing the atomic coordinates from those of the reference, unperturbed system. This compute can be used to calculate free energy differences using several methods, such as free-energy perturbation (FEP), finite-difference thermodynamic integration (FDTI) or Bennet's acceptance ratio method (BAR).

The potential energy of the system is decomposed in three terms: a background term corresponding to interaction sites whose parameters remain constant, a reference term  $U_0$  corresponding to the initial interactions of the atoms that will

undergo perturbation, and a term  $U_1$  corresponding to the final interactions of these atoms:

$$U(\lambda) = U_{\text{bg}} + U_1(\lambda) + U_0(\lambda)$$

A coupling parameter  $\lambda$  varying from 0 to 1 connects the reference and perturbed systems:

$$\begin{aligned}\lambda = 0 &\Rightarrow U = U_{\text{bg}} + U_0 \\ \lambda = 1 &\Rightarrow U = U_{\text{bg}} + U_1\end{aligned}$$

It is possible but not necessary that the coupling parameter (or a function thereof) appears as a multiplication factor of the potential energy. Therefore, this compute can apply perturbations to interaction parameters that are not directly proportional to the potential energy (e.g.,  $\sigma$  in Lennard-Jones potentials).

This command can be combined with *fix adapt* to perform multistage free-energy perturbation calculations along step-wise alchemical transformations during a simulation run:

$$\Delta_0^1 A = \sum_{i=0}^{n-1} \Delta_{\lambda_i}^{\lambda_{i+1}} A = -k_B T \sum_{i=0}^{n-1} \ln \left\langle \exp \left( -\frac{U(\lambda_{i+1}) - U(\lambda_i)}{k_B T} \right) \right\rangle_{\lambda_i}$$

This compute is suitable for the finite-difference thermodynamic integration (FDTI) method (*Mezei*), which is based on an evaluation of the numerical derivative of the free energy by a perturbation method using a very small  $\delta$ :

$$\Delta_0^1 A = \int_{\lambda=0}^{\lambda=1} \left( \frac{\partial A(\lambda)}{\partial \lambda} \right)_{\lambda} d\lambda \approx \sum_{i=0}^{n-1} w_i \frac{A(\lambda_i + \delta) - A(\lambda_i)}{\delta}$$

where  $w_i$  are weights of a numerical quadrature. The *fix adapt* command can be used to define the stages of  $\lambda$  at which the derivative is calculated and averaged.

The compute *fep* calculates the exponential Boltzmann term and also the potential energy difference  $U_1 - U_0$ . By choosing a very small perturbation  $\delta$  the thermodynamic integration method can be implemented using a numerical evaluation of the derivative of the potential energy with respect to  $\lambda$ :

$$\Delta_0^1 A = \int_{\lambda=0}^{\lambda=1} \left\langle \frac{\partial U(\lambda)}{\partial \lambda} \right\rangle_{\lambda} d\lambda \approx \sum_{i=0}^{n-1} w_i \left\langle \frac{U(\lambda_i + \delta) - U(\lambda_i)}{\delta} \right\rangle_{\lambda_i}$$

Another technique to calculate free energy differences is the acceptance ratio method (*Bennet*), which can be implemented by calculating the potential energy differences with  $\delta = 1.0$  on both the forward and reverse routes:

$$\left\langle \frac{1}{1 + \exp[(U_1 - U_0 - \Delta_0^1 A)/k_B T]} \right\rangle_0 = \left\langle \frac{1}{1 + \exp[(U_0 - U_1 + \Delta_0^1 A)/k_B T]} \right\rangle_1$$

The value of the free energy difference is determined by numerical root finding to establish the equality.

Concerning the choice of how the atomic parameters are perturbed in order to setup an alchemical transformation route, several strategies are available, such as single-topology or double-topology strategies (*Pearlman*). The latter does not require modification of bond lengths, angles or other internal coordinates.

NOTES: This compute command does not take kinetic energy into account, therefore the masses of the particles should not be modified between the reference and perturbed states, or along the alchemical transformation route. This compute command does not change bond lengths or other internal coordinates (*Boresch, Karplus*).

The *pair* attribute enables various parameters of potentials defined by the *pair\_style* and *pair\_coeff* commands to be changed, if the pair style supports it.

The *pstyle* argument is the name of the pair style. For example, *pstyle* could be specified as “lj/cut”. The *pparam* argument is the name of the parameter to change. This is a list of pair styles and parameters that can be used with this compute. See the doc pages for individual pair styles and their energy formulas for the meaning of these parameters:

<i>born</i>	a,b,c	type pairs
<i>buck, buck/coul/cut, buck/coul/long, buck/coul/msm</i>	a,c	type pairs
<i>buck/mdf</i>	a,c	type pairs
<i>coul/cut</i>	scale	type pairs
<i>coul/cut/soft</i>	lambda	type pairs
<i>coul/long, coul/msm</i>	scale	type pairs
<i>coul/long/soft</i>	scale, lambda	type pairs
<i>eam</i>	scale	type pairs
<i>gauss</i>	a	type pairs
<i>lennard/mdf</i>	a,b	type pairs
<i>lj/class2</i>	epsilon,sigma	type pairs
<i>lj/class2/coul/cut, lj/class2/coul/long</i>	epsilon,sigma	type pairs
<i>lj/cut</i>	epsilon,sigma	type pairs
<i>lj/cut/soft</i>	epsilon,sigma,lambda	type pairs
<i>lj/cut/coul/cut, lj/cut/coul/long, lj/cut/coul/msm</i>	epsilon,sigma	type pairs
<i>lj/cut/coul/cut/soft, lj/cut/coul/long/soft</i>	epsilon,sigma,lambda	type pairs
<i>lj/cut/tip4p/cut, lj/cut/tip4p/long</i>	epsilon,sigma	type pairs
<i>lj/cut/tip4p/long/soft</i>	epsilon,sigma,lambda	type pairs
<i>lj/expand</i>	epsilon,sigma,delta	type pairs
<i>lj/mdf</i>	epsilon,sigma	type pairs
<i>lj/sf/dipole/sf</i>	epsilon,sigma,scale	type pairs
<i>mie/cut</i>	epsilon,sigma,gamR,gamA	type pairs
<i>morse, morse/smooth/linear</i>	d0,r0,alpha	type pairs
<i>morse/soft</i>	d0,r0,alpha,lambda	type pairs
<i>nm/cut</i>	e0,r0,nn,mm	type pairs
<i>nm/cut/coul/cut, nm/cut/coul/long</i>	e0,r0,nn,mm	type pairs
<i>ufm</i>	epsilon,sigma,scale	type pairs
<i>soft</i>	a	type pairs

Note that it is easy to add new potentials and their parameters to this list. All it typically takes is adding an `extract()` method to the `pair_*.cpp` file associated with the potential.

Similar to the `pair_coeff` command, I and J can be specified in one of two ways. Explicit numeric values can be used for each, as in the first example above.  $I \leq J$  is required. LAMMPS sets the coefficients for the symmetric J,I interaction to the same values. A wild-card asterisk can be used in place of or in conjunction with the I,J arguments to set the coefficients for multiple pairs of atom types. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the number of atom types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $m$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive). Note that only type pairs with  $I \leq J$  are considered; if asterisks imply type pairs where  $J < I$ , they are ignored.

If `pair_style hybrid` or `hybrid/overlay` is being used, then the `pstyle` will be a sub-style name. You must specify I,J arguments that correspond to type pair values defined (via the `pair_coeff` command) for that sub-style.

The `v_name` argument for keyword `pair` is the name of an *equal-style variable* which will be evaluated each time this compute is invoked. It should be specified as `v_name`, where name is the variable name.

The `atom` attribute enables atom properties to be changed. The `aparam` argument is the name of the parameter to change. This is the current list of atom parameters that can be used with this compute:

- `charge` = charge on particle

The `v_name` argument for keyword `pair` is the name of an *equal-style variable* which will be evaluated each time this compute is invoked. It should be specified as `v_name`, where name is the variable name.

The *tail* keyword controls the calculation of the tail correction to “van der Waals” pair energies beyond the cutoff, if this has been activated via the *pair\_modify* command. If the perturbation is small, the tail contribution to the energy difference between the reference and perturbed systems should be negligible.

If the keyword *volume = yes*, then the Boltzmann term is multiplied by the volume so that correct ensemble averaging can be performed over trajectories during which the volume fluctuates or changes (*Allen and Tildesley*):

$$\Delta_0^1 A = -k_B T \sum_{i=0}^{n-1} \ln \frac{\langle V \exp\left(-\frac{U(\lambda_{i+1}) - U(\lambda_i)}{k_B T}\right) \rangle_{\lambda_i}}{\langle V \rangle_{\lambda_i}}$$

---

### 18.42.4 Output info

This compute calculates a global vector of length 3 which contains the energy difference ( $U_1 - U_0$ ) as `c_ID[1]`, the Boltzmann factor  $\exp(-(U_1 - U_0)/k_B T)$ , or  $V \exp(-(U_1 - U_0)/k_B T)$ , as `c_ID[2]` and the volume of the simulation box  $V$  as `c_ID[3]`.  $U_1$  is the pair potential energy obtained with the perturbed parameters and  $U_0$  is the pair potential energy obtained with the unperturbed parameters. The energies include `kspc` terms if these are used in the simulation.

These output results can be used by any command that uses a global scalar or vector from a compute as input. See the *Howto output* page for an overview of LAMMPS output options. For example, the computed values can be averaged using *fix ave/time*.

The values calculated by this compute are “extensive”.

### 18.42.5 Restrictions

This compute is distributed as the FEP package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 18.42.6 Related commands

*fix adapt/fep, fix ave/time, pair\_style .../soft*

### 18.42.7 Default

The option defaults are *tail = no, volume = no*.

---

**(Pearlman)** Pearlman, J Chem Phys, 98, 1487 (1994)

**(Mezei)** Mezei, J Chem Phys, 86, 7084 (1987)

**(Bennet)** Bennet, J Comput Phys, 22, 245 (1976)

**(BoreschKarplus)** Boresch and Karplus, J Phys Chem A, 103, 103 (1999)

**(AllenTildesley)** Allen and Tildesley, Computer Simulation of Liquids, Oxford University Press (1987)



## 18.43 compute fep/ta command

### 18.43.1 Syntax

```
compute ID group-ID fep/ta temp plane scale_factor keyword value ...
```

- ID, group-ID are documented in the *compute* command
- fep/ta = name of this compute command
- temp = external temperature (as specified for constant-temperature run)
- plane = *xy* or *xz* or *yz*
- scale\_factor = multiplicative factor for change in plane area
- zero or more keyword/value pairs may be appended
- keyword = *tail*  
*tail* value = *no* or *yes*  
*no* = ignore tail correction to pair energies (usually small in fep)  
*yes* = include tail correction to pair energies

### 18.43.2 Examples

```
compute 1 all fep/ta 298 xy 1.0005
```

### 18.43.3 Description

Define a computation that calculates the change in the free energy due to a test-area (TA) perturbation (*Gloor*). The test-area approach can be used to determine the interfacial tension of the system in a single simulation:

$$\gamma = \lim_{\Delta\mathcal{A} \rightarrow 0} \left( \frac{\Delta A_{0 \rightarrow 1}}{\Delta\mathcal{A}} \right)_{N,V,T} = -\frac{k_B T}{\Delta\mathcal{A}} \ln \left\langle \exp \left( \frac{-(U_1 - U_0)}{k_B T} \right) \right\rangle_0$$

During the perturbation, both axes of *plane* are scaled by multiplying  $\sqrt{\text{scale\_factor}}$ , while the other axis divided by *scale\_factor* such that the overall volume of the system is maintained.

The *tail* keyword controls the calculation of the tail correction to “van der Waals” pair energies beyond the cutoff, if this has been activated via the *pair\_modify* command. If the perturbation is small, the tail contribution to the energy difference between the reference and perturbed systems should be negligible.

### 18.43.4 Output info

This compute calculates a global vector of length 3 which contains the energy difference ( $U_1 - U_0$ ) as c\_ID[1], the Boltzmann factor  $\exp(-(U_1 - U_0)/k_B T)$ , as c\_ID[2] and the change in the *plane* area  $\Delta\mathcal{A}$  as c\_ID[3].  $U_1$  is the potential energy of the perturbed state and  $U_0$  is the potential energy of the reference state. The energies include kspace terms if these are used in the simulation.

These output results can be used by any command that uses a global scalar or vector from a compute as input. See the *Howto output* page for an overview of LAMMPS output options. For example, the computed values can be averaged using *fix ave/time*.

### 18.43.5 Restrictions

Constraints, like fix shake, may lead to incorrect values for energy difference.

This compute is distributed as the FEP package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 18.43.6 Related commands

*compute fep*

### 18.43.7 Default

The option defaults are *tail = no*.

---

(Gloor) Gloor, J Chem Phys, 123, 134703 (2005)

## 18.44 compute global/atom command

### 18.44.1 Syntax

```
compute ID group-ID style index input1 input2 ...
```

- ID, group-ID are documented in *compute* command
- global/atom = style name of this compute command
- index = c\_ID, c\_ID[N], f\_ID, f\_ID[N], v\_name

```
c_ID = per-atom vector calculated by a compute with ID
c_ID[I] = Ith column of per-atom array calculated by a compute with ID
f_ID = per-atom vector calculated by a fix with ID
f_ID[I] = Ith column of per-atom array calculated by a fix with ID
v_name = per-atom vector calculated by an atom-style variable with name
```

- one or more inputs can be listed
- input = c\_ID, c\_ID[N], f\_ID, f\_ID[N], v\_name

```
c_ID = global vector calculated by a compute with ID
c_ID[I] = Ith column of global array calculated by a compute with ID, I can include ↵
↵wildcard (see below)
f_ID = global vector calculated by a fix with ID
f_ID[I] = Ith column of global array calculated by a fix with ID, I can include ↵
↵wildcard (see below)
v_name = global vector calculated by a vector-style variable with name
```

## 18.44.2 Examples

```
compute 1 all global/atom c_chunk c_com[1] c_com[2] c_com[3]
compute 1 all global/atom c_chunk c_com[*]
```

## 18.44.3 Description

Define a calculation that assigns global values to each atom from vectors or arrays of global values. The specified *index* parameter is used to determine which global value is assigned to each atom.

The *index* parameter must reference a per-atom vector or array from a *compute* or *fix* or the evaluation of an atom-style *variable*. Each *input* value must reference a global vector or array from a *compute* or *fix* or the evaluation of an vector-style *variable*. Details are given below.

The *index* value for an atom is used as an index  $I$  (from 1 to  $N$ , where  $N$  is the number of atoms) into the vector associated with each of the input values. The  $I$ th value from the input vector becomes one output value for that atom. If the atom is not in the specified group, or the index  $I < 1$  or  $I > M$ , where  $M$  is the actual length of the input vector, then an output value of 0.0 is assigned to the atom.

An example of how this command is useful, is in the context of “chunks” which are static or dynamic subsets of atoms. The *compute chunk/atom* command assigns unique chunk IDs to each atom. Its output can be used as the *index* parameter for this command. Various other computes with “chunk” in their style name, such as *compute com/chunk* or *compute msd/chunk*, calculate properties for each chunk. The output of these commands are global vectors or arrays, with one or more values per chunk, and can be used as input values for this command. This command will then assign the global chunk value to each atom in the chunk, producing a per-atom vector or per-atom array as output. The per-atom values can then be output to a dump file or used by any command that uses per-atom values from a compute as input, as discussed on the *Howto output* doc page.

As a concrete example, these commands will calculate the displacement of each atom from the center-of-mass of the molecule it is in, and dump those values to a dump file. In this case, each molecule is a chunk.

```
compute cc1 all chunk/atom molecule
compute myChunk all com/chunk cc1
compute prop all property/atom xu yu zu
compute glob all global/atom c_cc1 c_myChunk[*]
variable dx atom c_prop[1]-c_glob[1]
variable dy atom c_prop[2]-c_glob[2]
variable dz atom c_prop[3]-c_glob[3]
variable dist atom sqrt(v_dx*v_dx+v_dy*v_dy+v_dz*v_dz)
dump 1 all custom 100 tmp.dump id xu yu zu c_glob[1] c_glob[2] c_glob[3] &
      v_dx v_dy v_dz v_dist
dump_modify 1 sort id
```

You can add these commands to the bench/in.chain script to see how they work.

Note that for input values from a compute or fix, the bracketed index  $I$  can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the size of the vector (for *mode* = scalar) or the number of columns in the array (for *mode* = vector), then an asterisk with no numeric values means all indices from 1 to  $N$ . A leading asterisk means all indices from 1 to  $n$  (inclusive). A trailing asterisk means all indices from  $m$  to  $N$  (inclusive). A middle asterisk means all indices from  $m$  to  $n$  (inclusive).

Using a wildcard is the same as if the individual columns of the array had been listed one by one. For example, the following two compute global/atom commands are equivalent, since the *compute com/chunk* command creates a global array with three columns:

```
compute cc1 all chunk/atom molecule
compute com all com/chunk cc1
compute 1 all global/atom c_cc1 c_com[1] c_com[2] c_com[3]
compute 1 all global/atom c_cc1 c_com[*]
```

---

This section explains the *index* parameter. Note that it must reference per-atom values, as contrasted with the *input* values, which must reference global values.

Note that all of these options generate floating point values. When they are used as an index into the specified input vectors, they are simply rounded down to convert the value to integer indices. The final values should range from 1 to  $N$  (inclusive), since they are used to access values from  $N$ -length vectors.

If *index* begins with “c\_”, a compute ID must follow which has been previously defined in the input script. The compute must generate per-atom quantities. See the individual *compute* doc page for details. If no bracketed integer is appended, the per-atom vector calculated by the compute is used. If a bracketed integer is appended, the  $I$ th column of the per-atom array calculated by the compute is used. Users can also write code for their own compute styles and *add them to LAMMPS*. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

If *index* begins with “f\_”, a fix ID must follow which has been previously defined in the input script. The Fix must generate per-atom quantities. See the individual *fix* page for details. Note that some fixes only produce their values on certain timesteps, which must be compatible with when compute global/atom references the values, else an error results. If no bracketed integer is appended, the per-atom vector calculated by the fix is used. If a bracketed integer is appended, the  $I$ th column of the per-atom array calculated by the fix is used. Users can also write code for their own fix style and *add them to LAMMPS*. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

If *index* begins with “v\_”, a variable name must follow which has been previously defined in the input script. It must be an *atom-style variable*. Atom-style variables can reference thermodynamic keywords and various per-atom attributes, or invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of generating per-atom quantities to use as *index*.

---

This section explains the kinds of *input* values that can be used. Note that inputs reference global values, as contrasted with the *index* parameter which must reference per-atom values.

If a value begins with “c\_”, a compute ID must follow which has been previously defined in the input script. The compute must generate a global vector or array. See the individual *compute* doc page for details. If no bracketed integer is appended, the vector calculated by the compute is used. If a bracketed integer is appended, the  $I$ th column of the array calculated by the compute is used. Users can also write code for their own compute styles and *add them to LAMMPS*. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

If a value begins with “f\_”, a fix ID must follow which has been previously defined in the input script. The fix must generate a global vector or array. See the individual *fix* doc page for details. Note that some fixes only produce their values on certain timesteps, which must be compatible with when compute global/atom references the values, else an error results. If no bracketed integer is appended, the vector calculated by the fix is used. If a bracketed integer is appended, the  $I$ th column of the array calculated by the fix is used. Users can also write code for their own fix style and *add them to LAMMPS*. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

If a value begins with “v\_”, a variable name must follow which has been previously defined in the input script. It must be a *vector-style variable*. Vector-style variables can reference thermodynamic keywords and various other attributes of atoms, or invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of generating a vector of global quantities which the *index* parameter will reference for assignment of global values to atoms.

### 18.44.4 Output info

If a single input is specified this compute produces a per-atom vector. If multiple inputs are specified, this compute produces a per-atom array values, where the number of columns is equal to the number of inputs specified. These values can be used by any command that uses per-atom vector or array values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The per-atom vector or array values will be in whatever units the corresponding input values are in.

### 18.44.5 Restrictions

none

### 18.44.6 Related commands

*compute, fix, variable, compute chunk/atom, compute reduce*

### 18.44.7 Default

none

## 18.45 compute group/group command

### 18.45.1 Syntax

```
compute ID group-ID group/group group2-ID keyword value ...
```

- ID, group-ID are documented in *compute* command
- group/group = style name of this compute command
- group2-ID = group ID of second (or same) group
- zero or more keyword/value pairs may be appended
- keyword = *pair* or *kpace* or *boundary* or *molecule*

*pair* value = *yes* or *no*

*kpace* value = *yes* or *no*

*boundary* value = *yes* or *no*

*molecule* value = *off* or *inter* or *intra*

## 18.45.2 Examples

```
compute 1 lower group/group upper
compute 1 lower group/group upper kspace yes
compute mine fluid group/group wall
```

## 18.45.3 Description

Define a computation that calculates the total energy and force interaction between two groups of atoms: the compute group and the specified group2. The two groups can be the same.

If the *pair* keyword is set to *yes*, which is the default, then the the interaction energy will include a pair component which is defined as the pairwise energy between all pairs of atoms where one atom in the pair is in the first group and the other is in the second group. Likewise, the interaction force calculated by this compute will include the force on the compute group atoms due to pairwise interactions with atoms in the specified group2.

---

**Note:** The energies computed by the *pair* keyword do not include tail corrections, even if they are enabled via the *pair\_modify* command.

---

If the *molecule* keyword is set to *inter* or *intra* than an additional check is made based on the molecule IDs of the two atoms in each pair before including their pairwise interaction energy and force. For the *inter* setting, the two atoms must be in different molecules. For the *intra* setting, the two atoms must be in the same molecule.

If the *kspace* keyword is set to *yes*, which is not the default, and if a *kspace\_style* is defined, then the interaction energy will include a Kspace component which is the long-range Coulombic energy between all the atoms in the first group and all the atoms in the second group. Likewise, the interaction force calculated by this compute will include the force on the compute group atoms due to long-range Coulombic interactions with atoms in the specified group2.

Normally the long-range Coulombic energy converges only when the net charge of the unit cell is zero. However, one can assume the net charge of the system is neutralized by a uniform background plasma, and a correction to the system energy can be applied to reduce artifacts. For more information see (*Bogusz*). If the *boundary* keyword is set to *yes*, which is the default, and *kspace* contributions are included, then this energy correction term will be added to the total group-group energy. This correction term does not affect the force calculation and will be zero if one or both of the groups are charge neutral. This energy correction term is the same as that included in the regular Ewald and PPPM routines.

---

**Note:** The *molecule* setting only affects the group/group contributions calculated by the *pair* keyword. It does not affect the group/group contributions calculated by the *kspace* keyword.

---

This compute does not calculate any bond or angle or dihedral or improper interactions between atoms in the two groups.

---

The pairwise contributions to the group-group interactions are calculated by looping over a neighbor list. The Kspace contribution to the group-group interactions require essentially the same amount of work (FFTs, Ewald summation) as computing long-range forces for the entire system. Thus it can be costly to invoke this compute too frequently.

---

**Note:** If you have a bonded system, then the settings of *special\_bonds* command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the *special\_bonds* command, and means those pairwise interactions do not appear in the neighbor list. Because this compute uses a neighbor list, it

also means those pairs will not be included in the `group/group` interaction. This does not apply when using long-range Coulomb interactions (*coul/long*, *coul/msm*, *coul/wolf* or similar). One way to get around this would be to set *special\_bond* scaling factors to very tiny numbers that are not exactly zero (e.g.,  $1.0 \times 10^{-50}$ ). Another workaround would be to write a dump file and use the *rerun* command to compute the `group/group` interactions for snapshots in the dump file. The rerun script can use a *special\_bonds* command that includes all pairs in the neighbor list.

---

If you desire a breakdown of the interactions into a pairwise and `Kspace` component, simply invoke the `compute` twice with the appropriate *yes/no* settings for the *pair* and *kpace* keywords. This is no more costly than using a single `compute` with both keywords set to *yes*. The individual contributions can be summed in a *variable* if desired.

This [document](#) describes how the long-range `group-group` calculations are performed.

---

### 18.45.4 Output info

This `compute` calculates a global scalar (the energy) and a global vector of length 3 (force), which can be accessed by indices 1–3. These values can be used by any command that uses global scalar or vector values from a `compute` as input. See the [Howto output](#) page for an overview of LAMMPS output options.

Both the scalar and vector values calculated by this `compute` are “extensive”. The scalar value will be in energy *units*. The vector values will be in force *units*.

### 18.45.5 Restrictions

Not all pair styles can be evaluated in a pairwise mode as required by this `compute`. For example, three-body and other many-body potentials, such as *Tersoff* and *Stillinger-Weber* cannot be used. *EAM* potentials will re-use previously computed embedding term contributions, so the computed pairwise forces and energies are based on the whole system and not valid if particles have been moved since.

Not all *Kspace styles* support the calculation of `group/group` interactions. The regular *ewald* and *pppm* styles do.

### 18.45.6 Related commands

none

### 18.45.7 Default

The option defaults are `pair = yes`, `kpace = no`, `boundary = yes`, `molecule = off`.

---

Bogusz et al, J Chem Phys, 108, 7070 (1998)

## 18.46 compute gyration command

### 18.46.1 Syntax

```
compute ID group-ID gyration
```

- ID, group-ID are documented in *compute* command
- gyration = style name of this compute command

### 18.46.2 Examples

```
compute 1 molecule gyration
```

### 18.46.3 Description

Define a computation that calculates the radius of gyration  $R_g$  of the group of atoms, including all effects due to atoms passing through periodic boundaries.

$R_g$  is a measure of the size of the group of atoms, and is computed as the square root of the  $R_g^2$  value in this formula

$$R_g^2 = \frac{1}{M} \sum_i m_i (r_i - r_{\text{cm}})^2$$

where  $M$  is the total mass of the group,  $r_{\text{cm}}$  is the center-of-mass position of the group, and the sum is over all atoms in the group.

A  $R_g^2$  tensor, stored as a 6-element vector, is also calculated by this compute. The formula for the components of the tensor is the same as the above formula, except that  $(r_i - r_{\text{cm}})^2$  is replaced by  $(r_{i,x} - r_{\text{cm},x}) \cdot (r_{i,y} - r_{\text{cm},y})$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx, yy, zz, xy, xz, yz$ . Note that unlike the scalar  $R_g$ , each of the six values of the tensor is effectively a “squared” value, since the cross-terms may be negative and taking a square root would be invalid.

---

**Note:** The coordinates of an atom contribute to  $R_g$  in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the *set image* command.

---

### 18.46.4 Output info

This compute calculates a global scalar ( $R_g$ ) and a global vector of length 6 ( $R_g^2$  tensor), which can be accessed by indices 1–6. These values can be used by any command that uses a global scalar value or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar and vector values calculated by this compute are “intensive”. The scalar and vector values will be in distance and distance<sup>2</sup> *units*, respectively.



## 18.46.5 Restrictions

none

## 18.46.6 Related commands

*compute gyration/chunk*, *compute gyration/shape*

## 18.46.7 Default

none

# 18.47 compute gyration/chunk command

## 18.47.1 Syntax

```
compute ID group-ID gyration/chunk chunkID keyword value ...
```

- ID, group-ID are documented in *compute* command
- gyration/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command
- zero or more keyword/value pairs may be appended
- keyword = *tensor*  
*tensor* value = none

## 18.47.2 Examples

```
compute 1 molecule gyration/chunk molchunk
compute 2 molecule gyration/chunk molchunk tensor
```

## 18.47.3 Description

Define a computation that calculates the radius of gyration  $R_g$  for multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

This compute calculates the radius of gyration  $R_g$  for each chunk, which includes all effects due to atoms passing through periodic boundaries.

$R_g$  is a measure of the size of a chunk, and is computed by the formula

$$R_g^2 = \frac{1}{M} \sum_i m_i (r_i - r_{cm})^2$$

where  $M$  is the total mass of the chunk,  $r_{\text{cm}}$  is the center-of-mass position of the chunk, and the sum is over all atoms in the chunk.

Note that only atoms in the specified group contribute to the calculation. The `compute chunk/atom` command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

If the `tensor` keyword is specified, then the scalar  $R_g$  value is not calculated, but an  $R_g$  tensor is instead calculated for each chunk. The formula for the components of the tensor is the same as the above formula, except that  $(r_i - r_{\text{cm}})^2$  is replaced by  $(r_{i,x} - r_{\text{cm},x}) \cdot (r_{i,y} - r_{\text{cm},y})$  for the  $xy$  component, and so on. The six components of the tensor are ordered  $xx, yy, zz, xy, xz, yz$ .

---

**Note:** The coordinates of an atom contribute to  $R_g$  in “unwrapped” form, by using the image flags associated with each atom. See the `dump custom` command for a discussion of “unwrapped” coordinates. See the Atoms section of the `read_data` command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the `set image` command.

---

The simplest way to output the results of the compute gyration/chunk calculation to a file is to use the `fix ave/time` command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk all gyration/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk file tmp.out mode vector
```

## 18.47.4 Output info

This compute calculates a global vector if the `tensor` keyword is not specified and a global array if it is. The length of the vector or number of rows in the array = the number of chunks  $N_{\text{chunk}}$  as calculated by the specified `compute chunk/atom` command. If the `tensor` keyword is specified, the global array has six columns. The vector or array can be accessed by any command that uses global values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

All the vector or array values calculated by this compute are “intensive”. The vector or array values will be in distance *units*, since they are the square root of values represented by the formula above.

## 18.47.5 Restrictions

none

## 18.47.6 Related commands

none

`compute gyration`

### 18.47.7 Default

none

## 18.48 compute gyration/shape command

### 18.48.1 Syntax

```
compute ID group-ID gyration/shape compute-ID
```

- ID, group-ID are documented in *compute* command
- gyration/shape = style name of this compute command
- compute-ID = ID of *compute gyration* command

### 18.48.2 Examples

```
compute 1 molecule gyration/shape pe
```

### 18.48.3 Description

Define a computation that calculates the eigenvalues of the gyration tensor of a group of atoms and three shape parameters. The computation includes all effects due to atoms passing through periodic boundaries.

The three computed shape parameters are the asphericity,  $b$ , the acylindricity,  $c$ , and the relative shape anisotropy,  $k$ , viz.,

$$b = l_z - \frac{1}{2}(l_y + l_x)$$

$$c = l_y - l_x$$

$$k = \frac{3}{2} \frac{l_x^2 + l_y^2 + l_z^2}{(l_x + l_y + l_z)^2} - \frac{1}{2}$$

where  $l_x \leq l_y \leq l_z$  are the three eigenvalues of the gyration tensor. A general description of these parameters is provided in (*Mattice*) while an application to polymer systems can be found in (*Theodorou*). The asphericity is always non-negative and zero only when the three principal moments are equal. This zero condition is met when the distribution of particles is spherically symmetric (hence the name asphericity) but also whenever the particle distribution is symmetric with respect to the three coordinate axes (e.g., when the particles are distributed uniformly on a cube, tetrahedron or other Platonic solid). The acylindricity is always non-negative and zero only when the two principal moments are equal. This zero condition is met when the distribution of particles is cylindrically symmetric (hence the name, acylindricity), but also whenever the particle distribution is symmetric with respect to the two coordinate axes (e.g., when the particles are distributed uniformly on a regular prism). The relative shape anisotropy is bounded between zero (if all points are spherically symmetric) and one (if all points lie on a line).

---

**Note:** The coordinates of an atom contribute to the gyration tensor in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the *set image* command.

---

### 18.48.4 Output info

This compute calculates a global vector of length 6, which can be accessed by indices 1–6. The first three values are the eigenvalues of the gyration tensor followed by the asphericity, the acylindricity and the relative shape anisotropy. The computed values can be used by any command that uses global vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector values calculated by this compute are “intensive”. The first five vector values will be in distance2 *units* while the sixth one is dimensionless.

### 18.48.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 18.48.6 Related commands

*compute gyration*

### 18.48.7 Default

none

---

(**Mattice**) Mattice, Suter, Conformational Theory of Large Molecules, Wiley, New York, 1994.

(**Theodorou**) Theodorou, Suter, Macromolecules, 18, 1206 (1985).

## 18.49 compute gyration/shape/chunk command

### 18.49.1 Syntax

```
compute ID group-ID gyration/shape/chunk compute-ID
```

- ID, group-ID are documented in *compute* command
- gyration/shape/chunk = style name of this compute command
- compute-ID = ID of *compute gyration/chunk* command

### 18.49.2 Examples

```
compute 1 molecule gyration/shape/chunk pe
```

### 18.49.3 Description

Define a computation that calculates the eigenvalues of the gyration tensor and three shape parameters of multiple chunks of atoms. The computation includes all effects due to atoms passing through periodic boundaries.

The three computed shape parameters are the asphericity,  $b$ , the acylindricity,  $c$ , and the relative shape anisotropy,  $k$ , viz.,

$$b = l_z - \frac{1}{2}(l_y + l_x)$$

$$c = l_y - l_x$$

$$k = \frac{3}{2} \frac{l_x^2 + l_y^2 + l_z^2}{(l_x + l_y + l_z)^2} - \frac{1}{2}$$

where  $l_x \leq l_y \leq l_z$  are the three eigenvalues of the gyration tensor. A general description of these parameters is provided in (*Mattice*) while an application to polymer systems can be found in (*Theodorou*). The asphericity is always non-negative and zero only when the three principal moments are equal. This zero condition is met when the distribution of particles is spherically symmetric (hence the name asphericity) but also whenever the particle distribution is symmetric with respect to the three coordinate axes (e.g., when the particles are distributed uniformly on a cube, tetrahedron, or other Platonic solid). The acylindricity is always non-negative and zero only when the two principal moments are equal. This zero condition is met when the distribution of particles is cylindrically symmetric (hence the name, acylindricity), but also whenever the particle distribution is symmetric with respect to the two coordinate axes (e.g., when the particles are distributed uniformly on a regular prism). The relative shape anisotropy is bounded between 0 (if all points are spherically symmetric) and 1 (if all points lie on a line).

The tensor keyword must be specified in the compute gyration/chunk command.

---

**Note:** The coordinates of an atom contribute to the gyration tensor in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the *set image* command.

---

### 18.49.4 Output info

This compute calculates a global array with six columns, which can be accessed by indices 1–6. The first three columns are the eigenvalues of the gyration tensor followed by the asphericity, the acylindricity and the relative shape anisotropy. The computed values can be used by any command that uses global array values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The array calculated by this compute is “intensive”. The first five columns will be in distance<sup>2</sup> *units* while the sixth one is dimensionless.

### 18.49.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 18.49.6 Related commands

*compute gyration/chunk compute gyration/shape*

## 18.49.7 Default

none

---

(**Mattice**) Mattice, Suter, Conformational Theory of Large Molecules, Wiley, New York, 1994.

(**Theodorou**) Theodorou, Suter, Macromolecules, 18, 1206 (1985).

## 18.50 compute heat/flux command

### 18.50.1 Syntax

```
compute ID group-ID heat/flux ke-ID pe-ID stress-ID
```

- ID, group-ID are documented in *compute* command
- heat/flux = style name of this compute command
- ke-ID = ID of a compute that calculates per-atom kinetic energy
- pe-ID = ID of a compute that calculates per-atom potential energy
- stress-ID = ID of a compute that calculates per-atom stress

### 18.50.2 Examples

```
compute myFlux all heat/flux myKE myPE myStress
```

### 18.50.3 Description

Define a computation that calculates the heat flux vector based on contributions from atoms in the specified group. This can be used by itself to measure the heat flux through a set of atoms (e.g., a region between two thermostatted reservoirs held at different temperatures), or to calculate a thermal conductivity using the equilibrium Green-Kubo formalism.

For other non-equilibrium ways to compute a thermal conductivity, see the *Howto kappa* doc page. These include use of the *fix thermal/conductivity* command for the Muller-Plathe method. Or the *fix heat* command which can add or subtract heat from groups of atoms.

The compute takes three arguments which are IDs of other *computes*. One calculates per-atom kinetic energy (*ke-ID*), one calculates per-atom potential energy (*pe-ID*), and the third calculates per-atom stress (*stress-ID*).

---

**Note:** These other computes should provide values for all the atoms in the group this compute specifies. That means the other computes could use the same group as this compute, or they can just use group “all” (or any group whose atoms are superset of the atoms in this compute’s group). LAMMPS does not check for this.

---

In case of two-body interactions, the heat flux  $\mathbf{J}$  is defined as

$$\begin{aligned}\mathbf{J} &= \frac{1}{V} \left[ \sum_i e_i \mathbf{v}_i - \sum_i \mathbf{S}_i \mathbf{v}_i \right] \\ &= \frac{1}{V} \left[ \sum_i e_i \mathbf{v}_i + \sum_{i<j} (\mathbf{F}_{ij} \cdot \mathbf{v}_j) \mathbf{r}_{ij} \right] \\ &= \frac{1}{V} \left[ \sum_i e_i \mathbf{v}_i + \frac{1}{2} \sum_{i<j} (\mathbf{F}_{ij} \cdot (\mathbf{v}_i + \mathbf{v}_j)) \mathbf{r}_{ij} \right]\end{aligned}$$

$e_i$  in the first term of the equation is the per-atom energy (potential and kinetic). This is calculated by the computes *ke-ID* and *pe-ID*.  $\mathbf{S}_i$  in the second term is the per-atom stress tensor calculated by the compute *stress-ID*. See *compute stress/atom* and *compute centroid/stress/atom* for possible definitions of atomic stress  $\mathbf{S}_i$  in the case of bonded and many-body interactions. The tensor multiplies  $\mathbf{v}_i$  by a  $3 \times 3$  matrix to yield a vector. Note that as discussed below, the  $1/V$  scaling factor in the equation for  $\mathbf{J}$  is **not** included in the calculation performed by these computes; you need to add it for a volume appropriate to the atoms included in the calculation.

**Note:** The *compute pe/atom* and *compute stress/atom* commands have options for which terms to include in their calculation (pair, bond, etc). The heat flux calculation will thus include exactly the same terms. Normally you should use *compute stress/atom virial* or *compute centroid/stress/atom virial* so as not to include a kinetic energy term in the heat flux.

**Warning:** The compute *heat/flux* has been reported to produce unphysical values for angle, dihedral, improper and constraint force contributions when used with *compute stress/atom*, as discussed in (*Surblys2019*), (*Boone*) and (*Surblys2021*). You are strongly advised to use *compute centroid/stress/atom*, which has been implemented specifically for such cases.

**Warning:** Due to an implementation detail, the *y* and *z* components of heat flux from *fix rigid* contribution when computed via *compute stress/atom* are highly unphysical and should not be used.

The Green–Kubo formulas relate the ensemble average of the auto-correlation of the heat flux  $\mathbf{J}$  to the thermal conductivity  $\kappa$ :

$$\kappa = \frac{V}{k_B T^2} \int_0^\infty \langle J_x(0) J_x(t) \rangle dt = \frac{V}{3k_B T^2} \int_0^\infty \langle \mathbf{J}(0) \cdot \mathbf{J}(t) \rangle dt$$

The heat flux can be output every so many timesteps (e.g., via the *thermo\_style custom* command). Then as a post-processing operation, an auto-correlation can be performed, its integral estimated, and the Green–Kubo formula above evaluated.

The *fix ave/correlate* command can calculate the auto-correlation. The trap() function in the *variable* command can calculate the integral.

An example LAMMPS input script for solid argon is appended below. The result should be an average conductivity  $\approx 0.29 \text{ W/m} \cdot \text{K}$ .

## 18.50.4 Output info

This compute calculates a global vector of length 6. The first three components are the  $x$ ,  $y$ , and  $z$  components of the full heat flux vector (i.e.,  $J_x$ ,  $J_y$ , and  $J_z$ ). The next three components are the  $x$ ,  $y$ , and  $z$  components of just the convective portion of the flux (i.e., the first term in the equation for  $\mathbf{J}$ ). Each component can be accessed by indices 1–6. These values can be used by any command that uses global vector values from a compute as input. See the *Howto output* documentation for an overview of LAMMPS output options.

The vector values calculated by this compute are “extensive”, meaning they scale with the number of atoms in the simulation. They can be divided by the appropriate volume to get a flux, which would then be an “intensive” value, meaning independent of the number of atoms in the simulation. Note that if the compute group is “all”, then the appropriate volume to divide by is the simulation box volume. However, if a group with a subset of atoms is used, it should be the volume containing those atoms.

The vector values will be in energy\*velocity *units*. Once divided by a volume the units will be that of flux, namely energy/area/time *units*

## 18.50.5 Restrictions

none

## 18.50.6 Related commands

*fix thermal/conductivity, fix ave/correlate, variable*

## 18.50.7 Default

none

---

## Example Input File

```
# Sample LAMMPS input script for thermal conductivity of solid Ar

units      real
variable  T equal 70
variable  V equal vol
variable  dt equal 4.0
variable  p equal 200      # correlation length
variable  s equal 10      # sample interval
variable  d equal $p*$s   # dump interval

# convert from LAMMPS real units to SI

variable  kB equal 1.3806504e-23  # [J/K] Boltzmann
variable  kCal2J equal 4186.0/6.02214e23
variable  A2m equal 1.0e-10
variable  fs2s equal 1.0e-15
variable  convert equal ${kCal2J}*${kCal2J}/${fs2s}/${A2m}
```

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```

# setup problem

dimension      3
boundary       p p p
lattice        fcc 5.376 orient x 1 0 0 orient y 0 1 0 orient z 0 0 1
region         box block 0 4 0 4 0 4
create_box     1 box
create_atoms   1 box
mass           1 39.948
pair_style     lj/cut 13.0
pair_coeff     * * 0.2381 3.405
timestep       ${dt}
thermo         $d

# equilibration and thermalization

velocity       all create $T 102486 mom yes rot yes dist gaussian
fix            NVT all nvt temp $T $T 10 drag 0.2
run            8000

# thermal conductivity calculation, switch to NVE if desired

#unfix        NVT
#fix          NVE all nve

reset_timestep 0
compute       myKE all ke/atom
compute       myPE all pe/atom
compute       myStress all stress/atom NULL virial
compute       flux all heat/flux myKE myPE myStress
variable      Jx equal c_flux[1]/vol
variable      Jy equal c_flux[2]/vol
variable      Jz equal c_flux[3]/vol
fix           JJ all ave/correlate $s $p $d &
              c_flux[1] c_flux[2] c_flux[3] type auto file J0Jt.dat ave running
variable      scale equal ${convert}/${kB}/$T/$T/$V*$s*${dt}
variable      k11 equal trap(f_JJ[3])*${scale}
variable      k22 equal trap(f_JJ[4])*${scale}
variable      k33 equal trap(f_JJ[5])*${scale}
thermo_style  custom step temp v_Jx v_Jy v_Jz v_k11 v_k22 v_k33
run           100000
variable      k equal (v_k11+v_k22+v_k33)/3.0
variable      ndens equal count(all)/vol
print         "average conductivity: $k[W/mK] @ $T K, ${ndens} /A\^3"

```

(Surblys2019) Surblys, Matsubara, Kikugawa, Ohara, Phys Rev E, 99, 051301(R) (2019).

(Boone) Boone, Babaei, Wilmer, J Chem Theory Comput, 15, 5579–5587 (2019).

(Surblys2021) Surblys, Matsubara, Kikugawa, Ohara, J Appl Phys 130, 215104 (2021).

## 18.51 compute hexorder/atom command

### 18.51.1 Syntax

```
compute ID group-ID hexorder/atom keyword values ...
```

- ID, group-ID are documented in *compute* command
- hexorder/atom = style name of this compute command
- one or more keyword/value pairs may be appended

keyword = *degree* or *nnn* or *cutoff*  
*cutoff* value = distance cutoff  
*nnn* value = number of nearest neighbors  
*degree* value = degree *n* of order parameter

### 18.51.2 Examples

```
compute 1 all hexorder/atom
compute 1 all hexorder/atom degree 4 nnn 4 cutoff 1.2
```

### 18.51.3 Description

Define a computation that calculates  $q_n$  the bond-orientational order parameter for each atom in a group. The hexatic ( $n = 6$ ) order parameter was introduced by *Nelson and Halperin* as a way to detect hexagonal symmetry in two-dimensional systems. For each atom,  $q_n$  is a complex number (stored as two real numbers) defined as follows:

$$q_n = \frac{1}{nnn} \sum_{j=1}^{nnn} e^{ni\theta(r_{ij})}$$

where the sum is over the *nnn* nearest neighbors of the central atom. The angle  $\theta$  is formed by the bond vector  $r_{ij}$  and the  $x$  axis.  $\theta$  is calculated only using the  $x$  and  $y$  components, whereas the distance from the central atom is calculated using all three  $x$ ,  $y$ , and  $z$  components of the bond vector. Neighbor atoms not in the group are included in the order parameter of atoms in the group.

The optional keyword *cutoff* defines the distance cutoff used when searching for neighbors. The default value, also the maximum allowable value, is the cutoff specified by the pair style.

The optional keyword *nnn* defines the number of nearest neighbors used to calculate  $q_n$ . The default value is 6. If the value is NULL, then all neighbors up to the distance cutoff are used.

The optional keyword *degree* sets the degree  $n$  of the order parameter. The default value is 6. For a perfect hexagonal lattice with  $nnn = 6$ ,  $q_6 = e^{6i\phi}$  for all atoms, where the constant  $0 < \phi < \frac{\pi}{3}$  depends only on the orientation of the lattice relative to the  $x$  axis. In an isotropic liquid, local neighborhoods may still exhibit weak hexagonal symmetry, but because the orientational correlation decays quickly with distance, the value of  $\phi$  will be different for different atoms, and so when  $q_6$  is averaged over all the atoms in the system,  $|\langle q_6 \rangle| \ll 1$ .

The value of  $q_n$  is set to zero for atoms not in the specified compute group, as well as for atoms that have less than *nnn* neighbors within the distance cutoff.

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (i.e. each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently.

---

**Note:** If you have a bonded system, then the settings of *special\_bonds* command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the *special\_bonds* command, and means those pairwise interactions do not appear in the neighbor list. Because this fix uses the neighbor list, it also means those pairs will not be included in the order parameter. This difficulty can be circumvented by writing a dump file, and using the *rerun* command to compute the order parameter for snapshots in the dump file. The rerun script can use a *special\_bonds* command that includes all pairs in the neighbor list.

---

### 18.51.4 Output info

This compute calculates a per-atom array with 2 columns, giving the real and imaginary parts  $q_n$ , a complex number restricted to the unit disk of the complex plane (i.e.,  $\Re(q_n)^2 + \Im(q_n)^2 \leq 1$ ).

These values can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

### 18.51.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 18.51.6 Related commands

*compute orientorder/atom*, *compute coord/atom*, *compute centro/atom*

### 18.51.7 Default

The option defaults are *cutoff* = pair style cutoff, *nnn* = 6, *degree* = 6

---

(Nelson) Nelson, Halperin, Phys Rev B, 19, 2457 (1979).

## 18.52 compute hma command

### 18.52.1 Syntax

```
compute ID group-ID hma temp-ID keyword ...
```

- ID, group-ID are documented in *compute* command
- hma = style name of this compute command
- temp-ID = ID of fix that specifies the set temperature during canonical simulation
- one or more keywords or keyword/argument pairs must be appended
- keyword = *anharmonic* or *u* or *p* or *cv*

*anharmonic* = compute will return anharmonic property values  
*u* = compute will return potential energy  
*p* value = Pharm = compute will return pressure  
     Pharm = difference between the harmonic pressure and lattice pressure  
           as described below  
*cv* = compute will return the heat capacity

## 18.52.2 Examples

```

compute 2 all hma 1 u
compute 2 all hma 1 anharmonic u p 0.9
compute 2 all hma 1 u cv
  
```

## 18.52.3 Description

Define a computation that calculates the properties of a solid (potential energy, pressure or heat capacity), using the harmonically-mapped averaging (HMA) method. This command yields much higher precision than the equivalent compute commands (*compute pe*, *compute pressure*, etc.) commands during a canonical simulation of an atomic crystal. Specifically, near melting HMA can yield averages of a given precision an order of magnitude faster than conventional methods, and this only improves as the temperatures is lowered. This is particularly important for evaluating the free energy by thermodynamic integration, where the low-temperature contributions are the greatest source of statistical uncertainty. Moreover, HMA has other advantages, including smaller potential-truncation effects, finite-size effects, smaller timestep inaccuracy, faster equilibration and shorter decorrelation time.

HMA should not be used if atoms are expected to diffuse. It is also restricted to simulations in the NVT ensemble. While this compute may be used with any potential in LAMMPS, it will provide inaccurate results for potentials that do not go to 0 at the truncation distance; *pair\_style lj/smooth/linear* and Ewald summation should work fine, while *pair\_style lj/cut* will perform poorly unless the potential is shifted (via *pair\_modify* shift) or the cutoff is large. Furthermore, computation of the heat capacity with this compute is restricted to those that implement the *single\_hessian* method in Pair. Implementing *single\_hessian* in additional pair styles is simple. Please contact Andrew Schultz (ajs42 at buffalo.edu) and David Kofke (kofke at buffalo.edu) if your desired pair style does not have this method. This is the list of pair styles that currently implement *single\_hessian*:

- *pair\_style lj/smooth/linear*

In this method, the analytically known harmonic behavior of a crystal is removed from the traditional ensemble averages, which leads to an accurate and precise measurement of the anharmonic contributions without contamination by noise produced by the already-known harmonic behavior. A detailed description of this method can be found in (Moustafa). The potential energy is computed by the formula:

$$\langle U \rangle_{\text{HMA}} = \frac{d}{2}(N-1)k_B T + \left\langle U + \frac{1}{2} \vec{F} \cdot \Delta \vec{r} \right\rangle$$

where  $N$  is the number of atoms in the system,  $k_B$  is Boltzmann's constant,  $T$  is the temperature,  $d$  is the dimensionality of the system (2 or 3 for 2d/3d),  $\vec{F} \cdot \Delta \vec{r}$  is the sum of dot products of the atomic force vectors and displacement (from lattice sites) vectors, and  $U$  is the sum of pair, bond, angle, dihedral, improper, kspace (long-range), and fix energies.

The pressure is computed by the formula:

$$\langle P \rangle_{\text{HMA}} = \Delta \hat{P} + \left\langle P_{\text{vir}} + \frac{\beta \Delta \hat{P} - \rho}{d(N-1)} \vec{F} \cdot \Delta \vec{r} \right\rangle$$

where  $\rho$  is the number density of the system,  $\Delta \hat{P}$  is the difference between the harmonic and lattice pressure,  $P_{\text{vir}}$  is the virial pressure computed as the sum of pair, bond, angle, dihedral, improper, kspace (long-range), and fix contributions

to the force on each atom, and  $k_B = 1/k_B T$ . Although the method will work for any value of  $\Delta\hat{P}$  specified (use pressure *units*), the precision of the resultant pressure is sensitive to  $\Delta\hat{P}$ ; the precision tends to be best when  $\Delta\hat{P}$  is the actual the difference between the lattice pressure and harmonic pressure.

$$\langle C_V \rangle_{\text{HMA}} = \frac{d}{2}(N-1)k_B + \frac{1}{k_B T^2} \left( \langle U_{\text{HMA}}^2 \rangle - \langle U_{\text{HMA}} \rangle^2 \right) + \frac{1}{4T} \langle \vec{F} \cdot \Delta\vec{r} + \Delta r \cdot \Phi \cdot \Delta\vec{r} \rangle$$

where  $\Phi$  is the Hessian matrix. The compute hma command computes the full expression for  $C_V$  except for the  $\langle U_{\text{HMA}} \rangle^2$  in the variance term, which can be obtained by passing the *u* keyword; you must add this extra contribution to the  $C_V$  value reported by this compute. The variance term can cause significant round-off error when computing  $C_V$ . To address this, the *anharmonic* keyword can be passed and/or the output format can be specified with more digits.

```
thermo_modify format float '%22.15e'
```

The *anharmonic* keyword will instruct the compute to return anharmonic properties rather than the full properties, which include lattice, harmonic and anharmonic contributions. When using this keyword, the compute must be first active (it must be included via a *thermo\_style custom* command) while the atoms are still at their lattice sites (before equilibration).

The temp-ID specified with compute hma command should be same as the fix-ID of the Nose–Hoover (*fix nvt*) or Berendsen (*fix temp/berendsen*) thermostat used for the simulation. While using this command, the Langevin thermostat (*fix langevin*) should be avoided as its extra forces interfere with the HMA implementation.

**Note:** Compute hma command should be used right after the energy minimization, when the atoms are at their lattice sites. The simulation should not be started before this command has been used in the input script.

The following example illustrates the placement of this command in the input script:

```
min_style cg
minimize 1e-35 1e-15 50000 500000
compute 1 all hma thermostaid u
fix thermostaid all nvt temp 600.0 600.0 100.0
```

**Note:** Compute hma should be used when the atoms of the solid do not diffuse. Diffusion will reduce the precision in the potential energy computation.

**Note:** The *fix\_modify energy yes* command must also be specified if a fix is to contribute potential energy to this command.

An example input script that uses this compute is included in examples/PACKAGES/hma/ along with corresponding LAMMPS output showing that the HMA properties fluctuate less than the corresponding conventional properties.

### 18.52.4 Output info

This compute calculates a global vector that includes the  $n$  properties requested as arguments to the command (the potential energy, pressure and/or heat capacity). The elements of the vector can be accessed by indices 1– $n$  by any command that uses global vector values as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector values calculated by this compute are “extensive”. The scalar value will be in energy *units*.

### 18.52.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is enabled only if LAMMPS was built with that package. See the *Build package* page for more info.

Usage restricted to canonical (NVT) ensemble simulation only.

### 18.52.6 Related commands

*compute pe*, *compute pressure*

*dynamical matrix* provides a finite difference formulation of the Hessian provided by Pair’s `single_hessian`, which is used by this compute.

### 18.52.7 Default

none

---

(**Moustafa**) Sabry G. Moustafa, Andrew J. Schultz, and David A. Kofke, *Very fast averaging of thermal properties of crystals by molecular simulation*, Phys. Rev. E [92], 043303 (2015)

## 18.53 compute improper command

### 18.53.1 Syntax

```
compute ID group-ID improper
```

- ID, group-ID are documented in *compute* command
- improper = style name of this compute command

### 18.53.2 Examples

```
compute 1 all improper
```

### 18.53.3 Description

Define a computation that extracts the improper energy calculated by each of the improper sub-styles used in the *improper\_style hybrid* command. These values are made accessible for output or further processing by other commands. The group specified for this command is ignored.

This compute is useful when using *improper\_style hybrid* if you want to know the portion of the total energy contributed by one or more of the hybrid sub-styles.

### 18.53.4 Output info

This compute calculates a global vector of length  $N$ , where  $N$  is the number of sub\_styles defined by the *improper\_style hybrid* command. These styles can be accessed by the indices 1 through  $N$ . These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector values are “extensive” and will be in energy *units*.

### 18.53.5 Restrictions

none

### 18.53.6 Related commands

*compute pe*, *compute pair*

### 18.53.7 Default

none

## 18.54 compute improper/local command

### 18.54.1 Syntax

```
compute ID group-ID improper/local value1 value2 ...
```

- ID, group-ID are documented in *compute* command
- improper/local = style name of this compute command
- one or more values may be appended
- value = *chi*  
*chi* = tabulate improper angles

## 18.54.2 Examples

```
compute 1 all improper/local chi
```

## 18.54.3 Description

Define a computation that calculates properties of individual improper interactions. The number of datums generated, aggregated across all processors, equals the number of impropers in the system, modified by the group parameter as explained below.

The value *chi* is the improper angle, as defined in the doc pages for the individual improper styles listed on *improper\_style* doc page.

The local data stored by this command is generated by looping over all the atoms owned on a processor and their impropers. An improper will only be included if all four atoms in the improper are in the specified compute group.

Note that as atoms migrate from processor to processor, there will be no consistent ordering of the entries within the local vector or array from one timestep to the next. The only consistency that is guaranteed is that the ordering on a particular timestep will be the same for local vectors or arrays generated by other compute commands. For example, improper output from the *compute property/local* command can be combined with data from this command and output by the *dump local* command in a consistent way.

Here is an example of how to do this:

```
compute 1 all property/local itype iatom1 iatom2 iatom3 iatom4
compute 2 all improper/local chi
dump 1 all local 1000 tmp.dump index c_1[1] c_1[2] c_1[3] c_1[4] c_1[5] c_2[1]
```

## 18.54.4 Output info

This compute calculates a local vector or local array depending on the number of keywords. The length of the vector or number of rows in the array is the number of impropers. If a single keyword is specified, a local vector is produced. If two or more keywords are specified, a local array is produced where the number of columns = the number of keywords. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The output for *chi* will be in degrees.

## 18.54.5 Restrictions

none

## 18.54.6 Related commands

*dump local*, *compute property/local*



## 18.54.7 Default

none

## 18.55 compute inertia/chunk command

### 18.55.1 Syntax

```
compute ID group-ID inertia/chunk chunkID
```

- ID, group-ID are documented in *compute* command
- inertia/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command

### 18.55.2 Examples

```
compute 1 fluid inertia/chunk molchunk
```

### 18.55.3 Description

Define a computation that calculates the inertia tensor for multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

This compute calculates the six components of the symmetric inertia tensor for each chunk, ordered  $I_{xx}, I_{yy}, I_{zz}, I_{xy}, I_{yz}, I_{xz}$ . The calculation includes all effects due to atoms passing through periodic boundaries.

Note that only atoms in the specified group contribute to the calculation. The *compute chunk/atom* command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

---

**Note:** The coordinates of an atom contribute to the chunk’s inertia tensor in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the *set image* command.

---

The simplest way to output the results of the compute inertia/chunk calculation to a file is to use the *fix ave/time* command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk all inertia/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk[*] file tmp.out mode vector
```

### 18.55.4 Output info

This compute calculates a global array where the number of rows = the number of chunks *Nchunk* as calculated by the specified *compute chunk/atom* command. The number of columns is 6, one for each of the 6 components of the inertia tensor for each chunk, ordered as listed above. These values can be accessed by any command that uses global array values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The array values are “intensive”. The array values will be in mass\*distance<sup>2</sup> *units*.

### 18.55.5 Restrictions

none

### 18.55.6 Related commands

*variable inertia()* function

### 18.55.7 Default

none

## 18.56 compute ke command

### 18.56.1 Syntax

```
compute ID group-ID ke
```

- ID, group-ID are documented in *compute* command
- ke = style name of this compute command

### 18.56.2 Examples

```
compute 1 all ke
```

### 18.56.3 Description

Define a computation that calculates the translational kinetic energy of a group of particles.

The kinetic energy of each particle is computed as  $\frac{1}{2}mv^2$ , where *m* and *v* are the mass and velocity of the particle, respectively.

There is a subtle difference between the quantity calculated by this compute and the kinetic energy calculated by the *ke* or *etotal* keyword used in thermodynamic output, as specified by the *thermo\_style* command. For this compute, kinetic energy is “translational” kinetic energy, calculated by the simple formula above. For thermodynamic output, the *ke* keyword infers kinetic energy from the temperature of the system with  $\frac{1}{2}k_B T$  of energy for each degree of freedom. For the default temperature computation via the *compute temp* command, these are the same. However, different computes that calculate temperature can subtract out different non-thermal components of velocity and/or include different degrees of freedom (translational, rotational, etc.).

### 18.56.4 Output info

This compute calculates a global scalar (the summed KE). This value can be used by any command that uses a global scalar value from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “extensive”. The scalar value will be in energy *units*.

### 18.56.5 Restrictions

none

### 18.56.6 Related commands

*compute erotate/sphere*

### 18.56.7 Default

none

## 18.57 compute ke/atom command

### 18.57.1 Syntax

```
compute ID group-ID ke/atom
```

- ID, group-ID are documented in *compute* command
- ke/atom = style name of this compute command

### 18.57.2 Examples

```
compute 1 all ke/atom
```

### 18.57.3 Description

Define a computation that calculates the per-atom translational kinetic energy for each atom in a group.

The kinetic energy is simply  $\frac{1}{2}mv^2$ , where  $m$  is the mass and  $v$  is the velocity of each atom.

The value of the kinetic energy will be 0.0 for atoms not in the specified compute group.

### 18.57.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values will be in energy *units*.

### 18.57.5 Restrictions

none

### 18.57.6 Related commands

*dump custom*

### 18.57.7 Default

none

## 18.58 compute ke/atom/eff command

### 18.58.1 Syntax

```
compute ID group-ID ke/atom/eff
```

- ID, group-ID are documented in *compute* command
- ke/atom/eff = style name of this compute command

### 18.58.2 Examples

```
compute 1 all ke/atom/eff
```

### 18.58.3 Description

Define a computation that calculates the per-atom translational (nuclei and electrons) and radial kinetic energy (electron only) in a group. The particles are assumed to be nuclei and electrons modeled with the *electronic force field*.

The kinetic energy for each nucleus is computed as  $\frac{1}{2}mv^2$ , where  $m$  corresponds to the corresponding nuclear mass, and the kinetic energy for each electron is computed as  $\frac{1}{2}(m_e v^2 + \frac{3}{4}m_e s^2)$ , where  $m_e$  and  $v$  correspond to the mass and translational velocity of each electron, and  $s$  to its radial velocity, respectively.

There is a subtle difference between the quantity calculated by this compute and the kinetic energy calculated by the *ke* or *etotal* keyword used in thermodynamic output, as specified by the *thermo\_style* command. For this compute, kinetic energy is “translational” plus electronic “radial” kinetic energy, calculated by the simple formula above. For thermodynamic output, the *ke* keyword infers kinetic energy from the temperature of the system with  $\frac{1}{2}k_B T$  of energy for each (nuclear-only) degree of freedom in eFF.

---

**Note:** The temperature in eFF should be monitored via the `compute temp/eff` command, which can be printed with thermodynamic output by using the `thermo_modify` command, as shown in the following example:

---

```
compute      effTemp all temp/eff
thermo_style custom step etotal pe ke temp press
thermo_modify temp effTemp
```

The value of the kinetic energy will be 0.0 for atoms (nuclei or electrons) not in the specified compute group.

## 18.58.4 Output info

This compute calculates a scalar quantity for each atom, which can be accessed by any command that uses per-atom computes as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values will be in energy *units*.

## 18.58.5 Restrictions

This compute is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 18.58.6 Related commands

*dump custom*

## 18.58.7 Default

none

# 18.59 compute ke/eff command

## 18.59.1 Syntax

```
compute ID group-ID ke/eff
```

- ID, group-ID are documented in [compute](#) command
- ke/eff = style name of this compute command

## 18.59.2 Examples

```
compute 1 all ke/eff
```

## 18.59.3 Description

Define a computation that calculates the kinetic energy of motion of a group of eFF particles (nuclei and electrons), as modeled with the *electronic force field*.

The kinetic energy for each nucleus is computed as  $\frac{1}{2}mv^2$  and the kinetic energy for each electron is computed as  $\frac{1}{2}(m_e v^2 + \frac{3}{4}m_e s^2)$ , where  $m$  corresponds to the nuclear mass,  $m_e$  to the electron mass,  $v$  to the translational velocity of each particle, and  $s$  to the radial velocity of the electron, respectively.

There is a subtle difference between the quantity calculated by this compute and the kinetic energy calculated by the *ke* or *etotal* keyword used in thermodynamic output, as specified by the *thermo\_style* command. For this compute, kinetic energy is “translational” and “radial” (only for electrons) kinetic energy, calculated by the simple formula above. For thermodynamic output, the *ke* keyword infers kinetic energy from the temperature of the system with  $\frac{1}{2}k_B T$  of energy for each degree of freedom. For the eFF temperature computation via the *compute temp\_eff* command, these are the same. But different computes that calculate temperature can subtract out different non-thermal components of velocity and/or include other degrees of freedom.

**Warning:** The temperature in eFF models should be monitored via the *compute temp/eff* command, which can be printed with thermodynamic output by using the *thermo\_modify* command, as shown in the following example:

```
compute      effTemp all temp/eff
thermo_style custom step etotal pe ke temp press
thermo_modify temp effTemp
```

See *compute temp/eff*.

## 18.59.4 Output info

This compute calculates a global scalar (the KE). This value can be used by any command that uses a global scalar value from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “extensive”. The scalar value will be in energy *units*.

## 18.59.5 Restrictions

This compute is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 18.59.6 Related commands

none

## 18.59.7 Default

none

# 18.60 compute ke/rigid command

## 18.60.1 Syntax

```
compute ID group-ID ke/rigid fix-ID
```

- ID, group-ID are documented in *compute* command
- ke = style name of this compute command
- fix-ID = ID of rigid body fix

## 18.60.2 Examples

```
compute 1 all ke/rigid myRigid
```

## 18.60.3 Description

Define a computation that calculates the translational kinetic energy of a collection of rigid bodies, as defined by one of the *fix rigid* command variants.

The kinetic energy of each rigid body is computed as  $\frac{1}{2}MV_{\text{cm}}^2$ , where  $M$  is the total mass of the rigid body, and  $V_{\text{cm}}$  is its center-of-mass velocity.

The *fix-ID* should be the ID of one of the *fix rigid* commands which defines the rigid bodies. The group specified in the compute command is ignored. The kinetic energy of all the rigid bodies defined by the fix rigid command is included in the calculation.

## 18.60.4 Output info

This compute calculates a global scalar (the summed KE of all the rigid bodies). This value can be used by any command that uses a global scalar value from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “extensive”. The scalar value will be in energy *units*.

### 18.60.5 Restrictions

This compute is part of the RIGID package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 18.60.6 Related commands

*compute erotate/rigid*

### 18.60.7 Default

none

## 18.61 compute mesont command

### 18.61.1 Syntax

```
compute ID group-ID mesont style
```

- ID, group-ID are documented in [compute](#) command
- mesont = style name of the compute command
- style = *estretch* or *ebend* or *etube*

### 18.61.2 Examples

```
compute 1 all mesont estretch
```

### 18.61.3 Description

These computes define computations for the stretching (*estretch*), bending (*ebend*), and intertube (*etube*) per-node (atom) and total energies. The evaluated value is selected by the style parameter passed to the compute (*estretch*, *ebend*, or *etube*).

### 18.61.4 Output info

These computes calculate per-node (per-atom) vectors, which can be accessed by any command that uses per-atom values from a compute as input, and global scalars. See the [Howto output](#) page for an overview of LAMMPS output options.

The computed values are provided in energy *units*.



### 18.61.5 Restrictions

These computes are part of the MESONT package. They are only enabled if LAMMPS is built with that package. See the *Build package* doc page for more info. In addition, *mesont pair\_style* must be used.

### 18.61.6 Related commands

*dump custom*

### 18.61.7 Default

none

## 18.62 compute mliap command

### 18.62.1 Syntax

```
compute ID group-ID mliap ... keyword values ...
```

- ID, group-ID are documented in *compute* command
- mliap = style name of this compute command
- two or more keyword/value pairs must be appended
- keyword = *model* or *descriptor* or *gradgradflag*
  - model* values = style
    - style = *linear* or *quadratic* or *mliappy*
  - descriptor* values = style filename
    - style = *sna*
    - filename = name of file containing descriptor definitions
  - gradgradflag* value = 0/1
    - toggle gradgrad method for force gradient

### 18.62.2 Examples

```
compute mliap model linear descriptor sna Ta06A.mliap.descriptor
```

### 18.62.3 Description

Compute style *mliap* provides a general interface to the gradient of machine-learning interatomic potentials with respect to model parameters. It is used primarily for calculating the gradient of energy, force, and stress components with respect to model parameters, which is useful when training *mliap pair\_style* models to match target data. It provides separate definitions of the interatomic potential functional form (*model*) and the geometric quantities that characterize the atomic positions (*descriptor*). By defining *model* and *descriptor* separately, it is possible to use many different models with a given descriptor, or many different descriptors with a given model. Currently, the compute supports just two models, *linear* and *quadratic*, and one descriptor, *sna*, the SNAP descriptor used by *pair\_style sna*, including

the linear, quadratic, and chem variants. Work is currently underway to extend the interface to handle neural network energy models, and it is also straightforward to add new descriptor styles.

The compute *mliap* command must be followed by two keywords *model* and *descriptor* in either order.

The *model* keyword is followed by the model style (*linear*, *quadratic* or *mliappy*). The *mliappy* model is only available if LAMMPS is built with the *mliappy* Python module. There are [specific installation instructions](#) for that module.

The *descriptor* keyword is followed by a descriptor style, and additional arguments. The compute currently supports two descriptor styles *sna* and *so3*, but it is straightforward to add additional descriptor styles. The SNAP descriptor style *sna* is the same as that used by *pair\_style sna*, including the linear, quadratic, and chem variants. A single additional argument specifies the descriptor filename containing the parameters and setting used by the SNAP descriptor. The descriptor filename usually ends in the *.mliap.descriptor* extension. The format of this file is identical to the descriptor file in the *pair\_style mliap*, and is described in detail there.

---

**Note:** The number of LAMMPS atom types (and the value of *nelems* in the model) must match the value of *nelems* in the descriptor file.

---

Compute *mliap* calculates a global array containing gradient information. The number of columns in the array is  $nelems \times nparams + 1$ . The first row of the array contain the derivative of potential energy with respect to. to each parameter and each element. The last six rows of the array contain the corresponding derivatives of the virial stress tensor, listed in Voigt notation: *pxx*, *pyy*, *pzz*, *pyz*, *pxz*, and *pxy*. In between the energy and stress rows are the  $3N$  rows containing the derivatives of the force components. See section below on output for a detailed description of how rows and columns are ordered.

The element in the last column of each row contains the potential energy, force, or stress, according to the row. These quantities correspond to the user-specified reference potential that must be subtracted from the target data when training a model. The potential energy calculation uses the built in compute *thermo\_pe*. The stress calculation uses a compute called *mliap\_press* that is automatically created behind the scenes, according to the following command:

```
compute mliap_press all pressure NULL virial
```

See section below on output for a detailed explanation of the data layout in the global array.

The optional keyword *gradgradflag* controls how the force gradient is calculated. A value of 1 requires that the model provide the matrix of double gradients of energy with respect to both parameters and descriptors. For the linear and quadratic models this matrix is sparse and so is easily calculated and stored. For other models, this matrix may be prohibitively expensive to calculate and store. A value of 0 requires that the descriptor provide the derivative of the descriptors with respect to the position of every neighbor atom. This is not optimal for linear and quadratic models, but may be a better choice for more complex models.

Atoms not in the group do not contribute to this compute. Neighbor atoms not in the group do not contribute to this compute. The neighbor list needed to compute this quantity is constructed each time the calculation is performed (i.e., each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently.

---

**Note:** If the user-specified reference potentials includes bonded and non-bonded pairwise interactions, then the settings of *special\_bonds* command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the *special\_bonds* command, and means those pairwise interactions do not appear in the neighbor list. Because this fix uses the neighbor list, it also means those pairs will not be included in the calculation. The *rerun* command is not an option here, since the reference potential is required for the last column of the global array. A work-around is to prevent pairwise interactions from being removed by explicitly adding a *tiny* positive value for every pairwise interaction that would otherwise be set to zero in the *special\_bonds* command.

---

## 18.62.4 Output info

Compute *mliap* evaluates a global array. The columns are arranged into *nelems* blocks, listed in order of element *I*. Each block contains one column for each of the *nparams* model parameters. A final column contains the corresponding energy, force component on an atom, or virial stress component. The rows of the array appear in the following order:

- 1 row: Derivatives of potential energy with respect to each parameter of each element.
- $3N$  rows: Derivatives of force components; the *x*, *y*, and *z* components of the force on atom *i* appear in consecutive rows. The atoms are sorted based on atom ID.
- 6 rows: Derivatives of the virial stress tensor with respect to each parameter of each element. The ordering of the rows follows Voigt notation: *pxx*, *pyy*, *pzz*, *pyz*, *pxz*, *pxy*.

These values can be accessed by any command that uses a global array from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options. To see how this command can be used within a Python workflow to train machine-learning interatomic potentials, see the examples in [FitSNAP](#).

## 18.62.5 Restrictions

This compute is part of the ML-IAP package. It is only enabled if LAMMPS was built with that package. In addition, building LAMMPS with the ML-IAP package requires building LAMMPS with the ML-SNAP package. The *mliappy* model also requires building LAMMPS with the PYTHON package. See the [Build package](#) page for more info.

## 18.62.6 Related commands

*pair\_style mliap*

## 18.62.7 Default

The keyword defaults are `gradgradflag = 1`

# 18.63 compute momentum command

## 18.63.1 Syntax

```
compute ID group-ID momentum
```

- ID, group-ID are documented in [compute](#) command
- momentum = style name of this compute command

## 18.63.2 Examples

```
compute 1 all momentum
```

## 18.63.3 Description

Define a computation that calculates the translational momentum  $p$  of a group of particles. It is computed as the sum  $\vec{p} = \sum_i m_i \cdot \vec{v}_i$  over all particles in the compute group, where  $m$  and  $v$  are the mass and velocity vector of the particle, respectively.

## 18.63.4 Output info

This compute calculates a global vector (the summed momentum) of length 3. This value can be used by any command that uses a global vector value from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The vector value calculated by this compute is “extensive”. The vector value will be in mass\*velocity *units*.

## 18.63.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 18.63.6 Related commands

## 18.63.7 Default

none

# 18.64 compute msd command

## 18.64.1 Syntax

```
compute ID group-ID msd keyword values ...
```

- ID, group-ID are documented in *compute* command
- msd = style name of this compute command
- zero or more keyword/value pairs may be appended
- keyword = *com* or *average*
  - com* value = *yes* or *no*
  - average* value = *yes* or *no*

## 18.64.2 Examples

```
compute 1 all msd
compute 1 upper msd com yes average yes
```

## 18.64.3 Description

Define a computation that calculates the mean-squared displacement (MSD) of the group of atoms, including all effects due to atoms passing through periodic boundaries. For computation of the non-Gaussian parameter of mean-squared displacement, see the *compute msd/nongauss* command.

A vector of four quantities is calculated by this compute. The first three elements of the vector are the squared  $dx$ ,  $dy$ , and  $dz$  displacements, summed and averaged over atoms in the group. The fourth element is the total squared displacement (i.e.,  $dx^2 + dy^2 + dz^2$ ), summed and averaged over atoms in the group.

The slope of the mean-squared displacement (MSD) versus time is proportional to the diffusion coefficient of the diffusing atoms.

The displacement of an atom is from its reference position. This is normally the original position at the time the compute command was issued, unless the *average* keyword is set to *yes*. The value of the displacement will be 0.0 for atoms not in the specified compute group.

If the *com* option is set to *yes* then the effect of any drift in the center-of-mass of the group of atoms is subtracted out before the displacement of each atom is calculated.

If the *average* option is set to *yes* then the reference position of an atom is based on the average position of that atom, corrected for center-of-mass motion if requested. The average position is a running average over all previous calls to the compute, including the current call. So on the first call it is current position, on the second call it is the arithmetic average of the current position and the position on the first call, and so on. Note that when using this option, the precise value of the mean square displacement will depend on the number of times the compute is called. So, for example, changing the frequency of thermo output may change the computed displacement. Also, the precise values will be changed if a single simulation is broken up into two parts, using either multiple run commands or a restart file. It only makes sense to use this option if the atoms are not diffusing, so that their average positions relative to the center of mass of the system are stationary. The most common case is crystalline solids undergoing thermal motion.

---

**Note:** Initial coordinates are stored in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g. to 0) before invoking this compute by using the *set image* command.

---



---

**Note:** If you want the quantities calculated by this compute to be continuous when running from a *restart file*, then you should use the same ID for this compute, as in the original run. This is so that the fix this compute creates to store per-atom quantities will also have the same ID, and thus be initialized correctly with atom reference positions from the restart file. When *average* is set to *yes*, then the atom reference positions are restored correctly, but not the number of samples used obtain them. As a result, the reference positions from the restart file are combined with subsequent positions as if they were from a single sample, instead of many, which will change the values of *msd* somewhat.

---

### 18.64.4 Output info

This compute calculates a global vector of length 4, which can be accessed by indices 1–4 by any command that uses global vector values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The vector values are “intensive”. The vector values will be in distance<sup>2</sup> *units*.

### 18.64.5 Restrictions

Compute *msd* cannot be used with a dynamic group.

### 18.64.6 Related commands

*compute msd/nongauss*, *compute displace\_atom*, *fix store/state*, *compute msd/chunk*

### 18.64.7 Default

The option default are `com = no`, `average = no`.

## 18.65 compute msd/chunk command

### 18.65.1 Syntax

```
compute ID group-ID msd/chunk chunkID
```

- ID, group-ID are documented in *compute* command
- msd/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command

### 18.65.2 Examples

```
compute 1 all msd/chunk molchunk
```

### 18.65.3 Description

Define a computation that calculates the mean-squared displacement (MSD) for multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

Four quantities are calculated by this compute for each chunk. The first 3 quantities are the squared *dx*, *dy*, and *dz* displacements of the center-of-mass. The fourth component is the total squared displacement (i.e.,  $dx^2 + dy^2 + dz^2$ ) of the center-of-mass. These calculations include all effects due to atoms passing through periodic boundaries.

Note that only atoms in the specified group contribute to the calculation. The `compute chunk/atom` command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

The slope of the mean-squared displacement (MSD) versus time is proportional to the diffusion coefficient of the diffusing chunks.

The displacement of the center-of-mass of the chunk is from its original center-of-mass position, calculated on the timestep this compute command was first invoked.

---

**Note:** The number of chunks *Nchunk* calculated by the `compute chunk/atom` command must remain constant each time this compute is invoked, so that the displacement for each chunk from its original position can be computed consistently. If *Nchunk* does not remain constant, an error will be generated. If needed, you can enforce a constant *Nchunk* by using the `nchunk once` or `ids once` options when specifying the `compute chunk/atom` command.

---



---

**Note:** This compute stores the original position (of the center-of-mass) of each chunk. When a displacement is calculated on a later timestep, it is assumed that the same atoms are assigned to the same chunk ID. However LAMMPS has no simple way to insure this is the case, though you can use the `ids once` option when specifying the `compute chunk/atom` command. Note that if this is not the case, the MSD calculation does not have a sensible meaning.

---



---

**Note:** The initial coordinates of the atoms in each chunk are stored in “unwrapped” form, by using the image flags associated with each atom. See the `dump custom` command for a discussion of “unwrapped” coordinates. See the Atoms section of the `read_data` command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the `set image` command.

---



---

**Note:** If you want the quantities calculated by this compute to be continuous when running from a *restart file*, then you should use the same ID for this compute, as in the original run. This is so that the fix this compute creates to store per-chunk quantities will also have the same ID, and thus be initialized correctly with chunk reference positions from the restart file.

---

The simplest way to output the results of the compute `msd/chunk` calculation to a file is to use the `fix ave/time` command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk all msd/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk[*] file tmp.out mode vector
```

## 18.65.4 Output info

This compute calculates a global array where the number of rows = the number of chunks *Nchunk* as calculated by the specified `compute chunk/atom` command. The number of columns = 4 for *dx*, *dy*, *dz*, and the total displacement. These values can be accessed by any command that uses global array values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The array values are “intensive”. The array values will be in distance<sup>2</sup> *units*.

## 18.65.5 Restrictions

none

## 18.65.6 Related commands

*compute msd*

## 18.65.7 Default

none

# 18.66 compute msd/nongauss command

## 18.66.1 Syntax

```
compute ID group-ID msd/nongauss keyword values ...
```

- ID, group-ID are documented in *compute* command
- msd/nongauss = style name of this compute command
- zero or more keyword/value pairs may be appended
- keyword = *com*  
*com* value = *yes* or *no*

## 18.66.2 Examples

```
compute 1 all msd/nongauss  
compute 1 upper msd/nongauss com yes
```

## 18.66.3 Description

Define a computation that calculates the mean-squared displacement (MSD) and non-Gaussian parameter (NGP) of the group of atoms, including all effects due to atoms passing through periodic boundaries.

A vector of three quantities is calculated by this compute. The first element of the vector is the total squared displacement,  $dr^2 = dx^2 + dy^2 + dz^2$ , of the atoms, and the second is the fourth power of these displacements,  $dr^4 = (dx^2 + dy^2 + dz^2)^2$ , summed and averaged over atoms in the group. The third component is the non-Gaussian diffusion parameter NGP,

$$\text{NGP}(t) = \frac{3 \langle (r(t) - r(0))^4 \rangle}{5 \langle (r(t) - r(0))^2 \rangle^2} - 1.$$

The NGP is a commonly used quantity in studies of dynamical heterogeneity. Its minimum theoretical value ( $-0.4$ ) occurs when all atoms have the same displacement magnitude.  $\text{NGP} = 0$  for Brownian diffusion, while  $\text{NGP} > 0$  when some mobile atoms move faster than others.



If the *com* option is set to *yes* then the effect of any drift in the center-of-mass of the group of atoms is subtracted out before the displacement of each atom is calculated.

See the *compute msd* page for further important NOTES, which also apply to this compute.

## 18.66.4 Output info

This compute calculates a global vector of length 3, which can be accessed by indices 1–3 by any command that uses global vector values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The vector values are “intensive”. The first vector value will be in distance<sup>2</sup> *units*, the second is in distance<sup>4</sup> units, and the third is dimensionless.

## 18.66.5 Restrictions

Compute *msd/nongauss* cannot be used with a dynamic group.

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 18.66.6 Related commands

*compute msd*

## 18.66.7 Default

The option default is *com = no*.

# 18.67 compute nbond/atom command

## 18.67.1 Syntax

```
compute ID group-ID nbond/atom
```

- ID, group-ID are documented in *compute* command
- nbond/atom = style name of this compute command

## 18.67.2 Examples

```
compute 1 all nbond/atom
```

### 18.67.3 Description

Define a computation that computes the number of bonds each atom is part of. Bonds which are broken are not counted in the tally. See the *Howto broken bonds* page for more information. The number of bonds will be zero for atoms not in the specified compute group. This compute does not depend on Newton bond settings.

### 18.67.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

### 18.67.5 Restrictions

This fix can only be used if LAMMPS was built with the BPM package. See the *Build package* doc page for more info.

### 18.67.6 Related commands

### 18.67.7 Default

none

## 18.68 compute omega/chunk command

### 18.68.1 Syntax

```
compute ID group-ID omega/chunk chunkID
```

- ID, group-ID are documented in *compute* command
- omega/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command

### 18.68.2 Examples

```
compute 1 fluid omega/chunk molchunk
```

### 18.68.3 Description

Define a computation that calculates the angular velocity (omega) of multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

This compute calculates the three components of the angular velocity vector for each chunk via the formula  $\vec{L} = I \cdot \vec{\omega}$ , where  $\vec{L}$  is the angular momentum vector of the chunk,  $I$  is its moment of inertia tensor, and  $\omega$  is the angular velocity of the chunk. The calculation includes all effects due to atoms passing through periodic boundaries.

Note that only atoms in the specified group contribute to the calculation. The `compute chunk/atom` command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

---

**Note:** The coordinates of an atom contribute to the chunk’s angular velocity in “unwrapped” form, by using the image flags associated with each atom. See the `dump custom` command for a discussion of “unwrapped” coordinates. See the Atoms section of the `read_data` command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the `set image` command.

---

The simplest way to output the results of the compute omega/chunk calculation to a file is to use the `fix ave/time` command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk all omega/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk[*] file tmp.out mode vector
```

## 18.68.4 Output info

This compute calculates a global array where the number of rows is the number of chunks  $N_{chunk}$  as calculated by the specified `compute chunk/atom` command. The number of columns is 3 for the three ( $x$ ,  $y$ ,  $z$ ) components of the angular velocity for each chunk. These values can be accessed by any command that uses global array values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The array values are “intensive”. The array values will be in velocity/distance *units*.

## 18.68.5 Restrictions

none

## 18.68.6 Related commands

*variable omega() function*

## 18.68.7 Default

none

## 18.69 compute orientorder/atom command

Accelerator Variants: *orientorder/atom/kk*

### 18.69.1 Syntax

```
compute ID group-ID orientorder/atom keyword values ...
```

- ID, group-ID are documented in *compute* command
- orientorder/atom = style name of this compute command
- one or more keyword/value pairs may be appended

keyword = *cutoff* or *nnn* or *degrees* or *wl* or *wl/hat* or *components* or *chunksize*  
*cutoff* value = distance cutoff  
*nnn* value = number of nearest neighbors  
*degrees* values = nlvalues, 11, 12, ...  
*wl* value = *yes* or *no*  
*wl/hat* value = *yes* or *no*  
*components* value = ldegree  
*chunksize* value = number of atoms in each pass

### 18.69.2 Examples

```
compute 1 all orientorder/atom
compute 1 all orientorder/atom degrees 5 4 6 8 10 12 nnn NULL cutoff 1.5
compute 1 all orientorder/atom wl/hat yes
compute 1 all orientorder/atom components 6
```

### 18.69.3 Description

Define a computation that calculates a set of bond-orientational order parameters  $Q_\ell$  for each atom in a group. These order parameters were introduced by *Steinhardt et al.* as a way to characterize the local orientational order in atomic structures. For each atom,  $Q_\ell$  is a real number defined as follows:

$$\bar{Y}_{\ell m} = \frac{1}{nnn} \sum_{j=1}^{nnn} Y_{\ell m}(\theta(\mathbf{r}_{ij}), \phi(\mathbf{r}_{ij}))$$

$$Q_\ell = \sqrt{\frac{4\pi}{2\ell + 1} \sum_{m=-\ell}^{m=\ell} \bar{Y}_{\ell m} \bar{Y}_{\ell m}^*}$$

The first equation defines the local order parameters as averages of the spherical harmonics  $Y_{\ell m}$  for each neighbor. These are complex number components of the 3D analog of the 2D order parameter  $q_n$ , which is implemented as LAMMPS compute *hexorder/atom*. The summation is over the *nnn* nearest neighbors of the central atom. The angles  $\theta$  and  $\phi$  are the standard spherical polar angles defining the direction of the bond vector  $r_{ij}$ . The phase and sign of  $Y_{\ell m}$  follow the standard conventions, so that  $\text{sign}(Y_{\ell\ell}(0,0)) = (-1)^\ell$ . The second equation defines  $Q_\ell$ , which is a rotationally invariant non-negative amplitude obtained by summing over all the components of degree  $\ell$ .

The optional keyword *cutoff* defines the distance cutoff used when searching for neighbors. The default value, also the maximum allowable value, is the cutoff specified by the pair style.

The optional keyword *nnn* defines the number of nearest neighbors used to calculate  $Q_\ell$ . The default value is 12. If the value is NULL, then all neighbors up to the specified distance cutoff are used.

The optional keyword *degrees* defines the list of order parameters to be computed. The first argument *nvalues* is the number of order parameters. This is followed by that number of non-negative integers giving the degree of each order parameter. Because  $Q_2$  and all odd-degree order parameters are zero for atoms in cubic crystals (see *Steinhardt*), the default order parameters are  $Q_4, Q_6, Q_8, Q_{10}$ , and  $Q_{12}$ . For the FCC crystal with  $nnn = 12$ ,

$$Q_4 = \sqrt{\frac{7}{192}} \approx 0.19094$$

The numerical values of all order parameters up to  $Q_{12}$  for a range of commonly encountered high-symmetry structures are given in Table I of *Mickel et al.*, and these can be reproduced with this compute.

The optional keyword *wl* will output the third-order invariants  $W_\ell$  (see Eq. 1.4 in *Steinhardt*) for the same degrees as for the  $Q_\ell$  parameters. For the FCC crystal with  $nnn = 12$ ,

$$W_4 = -\sqrt{\frac{14}{143}} \left( \frac{49}{4096} \right) \pi^{-3/2} \approx -0.0006722136$$

The optional keyword *wlthat* will output the normalized third-order invariants  $\hat{W}_\ell$  (see Eq. 2.2 in *Steinhardt*) for the same degrees as for the  $Q_\ell$  parameters. For the FCC crystal with  $nnn = 12$ ,

$$\hat{W}_4 = -\frac{7}{3} \sqrt{\frac{2}{429}} \approx -0.159317$$

The numerical values of  $\hat{W}_\ell$  for a range of commonly encountered high-symmetry structures are given in Table I of *Steinhardt*, and these can be reproduced with this keyword.

The optional keyword *components* will output the components of the *normalized* complex vector  $\hat{Y}_{\ell m} = \bar{Y}_{\ell m} / |\bar{Y}_{\ell m}|$  of degree *ldegree*, which must be included in the list of order parameters to be computed. This option can be used in conjunction with *compute coord\_atom* to calculate the ten Wolde's criterion to identify crystal-like particles, as discussed in *ten Wolde*.

The optional keyword *chunksize* is only applicable when using the the KOKKOS package and is ignored otherwise. This keyword controls the number of atoms in each pass used to compute the bond-orientational order parameters and is used to avoid running out of memory. For example if there are 32768 atoms in the simulation and the *chunksize* is set to 16384, the parameter calculation will be broken up into two passes.

The value of  $Q_\ell$  is set to zero for atoms not in the specified compute group, as well as for atoms that have less than *nnn* neighbors within the distance cutoff, unless *nnn* is NULL.

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (i.e., each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently.

---

**Note:** If you have a bonded system, then the settings of *special\_bonds* command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the *special\_bonds* command, and means those pairwise interactions do not appear in the neighbor list. Because this fix uses the neighbor list, it also means those pairs will not be included in the order parameter. This difficulty can be circumvented by writing a dump file, and using the *rerun* command to compute the order parameter for snapshots in the dump file. The rerun script can use a *special\_bonds* command that includes all pairs in the neighbor list.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 18.69.4 Output info

This compute calculates a per-atom array with *nvalues* columns, giving the  $Q_\ell$  values for each atom, which are real numbers in the range  $0 \leq Q_\ell \leq 1$ .

If the keyword *wl* is set to yes, then the  $W_\ell$  values for each atom will be added to the output array, which are real numbers.

If the keyword *wl/hat* is set to yes, then the  $\hat{W}_\ell$  values for each atom will be added to the output array, which are real numbers.

If the keyword *components* is set, then the real and imaginary parts of each component of *normalized*  $\hat{Y}_{\ell m}$  will be added to the output array in the following order:  $\Re(\hat{Y}_{-m})$ ,  $\Im(\hat{Y}_{-m})$ ,  $\Re(\hat{Y}_{-m+1})$ ,  $\Im(\hat{Y}_{-m+1})$ , ...,  $\Re(\hat{Y}_m)$ ,  $\Im(\hat{Y}_m)$ .

In summary, the per-atom array will contain *nvalues* columns, followed by an additional *nvalues* columns if *wl* is set to yes, followed by an additional *nvalues* columns if *wl/hat* is set to yes, followed by an additional  $2*(2* \textit{ldegree}+1)$  columns if the *components* keyword is set.

These values can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options.

## 18.69.5 Restrictions

none

## 18.69.6 Related commands

*compute coord/atom*, *compute centro/atom*, *compute hexorder/atom*

## 18.69.7 Default

The option defaults are *cutoff* = pair style cutoff, *nnn* = 12, *degrees* = 5 4 6 8 10 12 (i.e.,  $Q_4$ ,  $Q_6$ ,  $Q_8$ ,  $Q_{10}$ , and  $Q_{12}$ ), *wl* = no, *wl/hat* = no, *components* off, and *chunksize* = 16384

---

**(Steinhardt)** P. Steinhardt, D. Nelson, and M. Ronchetti, Phys. Rev. B 28, 784 (1983).

**(Mickel)** W. Mickel, S. C. Kapfer, G. E. Schroeder-Turkand, K. Mecke, J. Chem. Phys. 138, 044501 (2013).

**(tenWolde)** P. R. ten Wolde, M. J. Ruiz-Montero, D. Frenkel, J. Chem. Phys. 104, 9932 (1996).

## 18.70 compute pair command

### 18.70.1 Syntax

```
compute ID group-ID pair pstyle [nstyle] [evaluate]
```

- ID, group-ID are documented in *compute* command
- pair = style name of this compute command
- pstyle = style name of a pair style that calculates additional values
- nsub = *n*-instance of a sub-style, if a pair style is used multiple times in a hybrid style
- *evaluate* = *epair* or *evdwl* or *ecoul* or blank (optional)

### 18.70.2 Examples

```
compute 1 all pair gauss
compute 1 all pair lj/cut/coul/cut ecoul
compute 1 all pair tersoff 2 epair
compute 1 all pair reaxff
```

### 18.70.3 Description

Define a computation that extracts additional values calculated by a pair style, and makes them accessible for output or further processing by other commands.

---

**Note:** The group specified for this command is **ignored**.

---

The specified *pstyle* must be a pair style used in your simulation either by itself or as a sub-style in a *pair\_style hybrid* or *hybrid/overlay* command. If the sub-style is used more than once, an additional number *nsub* has to be specified in order to choose which instance of the sub-style will be used by the compute. Not specifying the number in this case will cause the compute to fail.

The *evaluate* setting is optional. All pair styles tally a potential energy *epair* which may be broken into two parts: *evdwl* and *ecoul* such that  $epair = evdwl + ecoul$ . If the pair style calculates Coulombic interactions, their energy will be tallied in *ecoul*. Everything else (whether it is a Lennard-Jones style van der Waals interaction or not) is tallied in *evdwl*. If *evaluate* is blank or specified as *epair*, then *epair* is stored as a global scalar by this compute. This is useful when using *pair\_style hybrid* if you want to know the portion of the total energy contributed by one sub-style. If *evaluate* is specified as *evdwl* or *ecoul*, then just that portion of the energy is stored as a global scalar.

---

**Note:** The energy returned by the *evdwl* keyword does not include tail corrections, even if they are enabled via the *pair\_modify* command.

---

Some pair styles tally additional quantities, e.g. a breakdown of potential energy into 14 components is tallied by the *pair\_style reaxff* command. These values (1 or more) are stored as a global vector by this compute. See the page for *individual pair styles* for info on these values.

## 18.70.4 Output info

This compute calculates a global scalar which is *epair* or *evdwl* or *ecoul*. If the pair style supports it, it also calculates a global vector of length  $\geq 1$ , as determined by the pair style. These values can be used by any command that uses global scalar or vector values from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options.

The scalar and vector values calculated by this compute are “extensive”.

The scalar value will be in energy *units*. The vector values will typically also be in energy *units*, but see the page for the pair style for details.

## 18.70.5 Restrictions

none

## 18.70.6 Related commands

*compute pe, compute bond, fix pair*

## 18.70.7 Default

The keyword defaults are *evaluate = epair*, *nsub = 0*.

# 18.71 compute pair/local command

## 18.71.1 Syntax

```
compute ID group-ID pair/local value1 value2 ... keyword args ...
```

- ID, group-ID are documented in *compute* command
- pair/local = style name of this compute command
- one or more values may be appended
- value = *dist* or *dx* or *dy* or *dz* or *eng* or *force* or *fx* or *fy* or *fz* or *p1* or *p2* or ...
  - dist* = pairwise distance
  - dx,dy,dz* = components of pairwise distance
  - eng* = pairwise energy
  - force* = pairwise force
  - fx,fy,fz* = components of pairwise force
  - p1, p2, ...* = pair style specific quantities for allowed N values
- zero or more keyword/arg pairs may be appended
- keyword = *cutoff*
  - cutoff* arg = *type* or *radius*



## 18.71.2 Examples

```
compute 1 all pair/local eng
compute 1 all pair/local dist eng force
compute 1 all pair/local dist eng fx fy fz
compute 1 all pair/local dist fx fy fz p1 p2 p3
```

## 18.71.3 Description

Define a computation that calculates properties of individual pairwise interactions. The number of datums generated, aggregated across all processors, equals the number of pairwise interactions in the system.

The local data stored by this command is generated by looping over the pairwise neighbor list. Info about an individual pairwise interaction will only be included if both atoms in the pair are in the specified compute group, and if the current pairwise distance is less than the force cutoff distance for that interaction, as defined by the *pair\_style* and *pair\_coeff* commands.

The value *dist* is the distance between the pair of atoms. The values *dx*, *dy*, and *dz* are the  $(x, y, z)$  components of the *distance* between the pair of atoms. This value is always the distance from the atom of lower to the one with the higher id.

The value *eng* is the interaction energy for the pair of atoms.

The value *force* is the force acting between the pair of atoms, which is positive for a repulsive force and negative for an attractive force. The values *fx*, *fy*, and *fz* are the  $(x, y, z)$  components of *force* on atom I.

A pair style may define additional pairwise quantities which can be accessed as *p1* to *pN*, where *N* is defined by the pair style. Most pair styles do not define any additional quantities, so  $N = 0$ . An example of ones that do are the *granular pair styles* which calculate the tangential force between two particles and return its components and magnitude acting on atom *I* for  $N \in \{1, 2, 3, 4\}$ . See individual pair styles for details.

When using *pN* with pair style *hybrid*, the output will be the *N*th quantity from the sub-style that computes the pairwise interaction (based on atom types). If that sub-style does not define a *pN*, the output will be 0.0. The maximum allowed *N* is the maximum number of quantities provided by any sub-style.

When using *pN* with pair style *hybrid/overlay* the quantities from all sub-styles that provide them are concatenated together into one long list. For example, if there are 3 sub-styles and 2 of them have additional output (with 3 and 4 quantities, respectively), then 7 values (*p1* up to *p7*) are defined. The values *p1* to *p3* refer to quantities defined by the first of the two sub-styles. Values *p4* to *p7* refer to quantities from the second of the two sub-styles. If the referenced *pN* is not computed for the specific pairwise interaction (based on atom types), then the output will be 0.0.

The value *dist*, *dx*, *dy* and *dz* will be in distance *units*. The value *eng* will be in energy *units*. The values *force*, *fx*, *fy*, and *fz* will be in force *units*. The values *pN* will be in whatever units the pair style defines.

The optional *cutoff* keyword determines how the force cutoff distance for an interaction is determined. For the default setting of *type*, the pairwise cutoff defined by the *pair\_style* command for the types of the two atoms is used. For the *radius* setting, the sum of the radii of the two particles is used as a cutoff. For example, this is appropriate for granular particles which only interact when they are overlapping, as computed by *granular pair styles*. Note that if a granular model defines atom types such that all particles of a specific type are monodisperse (same diameter), then the two settings are effectively identical.

Note that as atoms migrate from processor to processor, there will be no consistent ordering of the entries within the local vector or array from one timestep to the next. The only consistency that is guaranteed is that the ordering on a particular timestep will be the same for local vectors or arrays generated by other compute commands. For example, pair output from the *compute property/local* command can be combined with data from this command and output by the *dump local* command in a consistent way.

Here is an example of how to do this:

```
compute 1 all property/local patom1 patom2
compute 2 all pair/local dist eng force
dump 1 all local 1000 tmp.dump index c_1[1] c_1[2] c_2[1] c_2[2] c_2[3]
```

**Note:** For pairs, if two atoms I,J are involved in 1–2, 1–3, and 1–4 interactions within the molecular topology, their pairwise interaction may be turned off, and thus they may not appear in the neighbor list, and will not be part of the local data created by this command. More specifically, this will be true of I,J pairs with a weighting factor of 0.0; pairs with a non-zero weighting factor are included. The weighting factors for 1–2, 1–3, and 1–4 pairwise interactions are set by the *special\_bonds* command. An exception is if long-range Coulombics are being computed via the *kspace\_style* command, then atom pairs with weighting factors of zero are still included in the neighbor list, so that a portion of the long-range interaction contribution can be computed in the pair style. Hence in that case, those atom pairs will be part of the local data created by this command.

---

### 18.71.4 Output info

This compute calculates a local vector or local array depending on the number of keywords. The length of the vector or number of rows in the array is the number of pairs. If a single keyword is specified, a local vector is produced. If two or more keywords are specified, a local array is produced where the number of columns = the number of keywords. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The output for *dist* will be in distance *units*. The output for *eng* will be in energy *units*. The output for *force*, *fx*, *fy*, and *fz* will be in force *units*. The output for *pN* will be in whatever units the pair style defines.

### 18.71.5 Restrictions

none

### 18.71.6 Related commands

*dump local*, *compute property/local*

### 18.71.7 Default

The keyword default is `cutoff = type`.

## 18.72 compute pe command

### 18.72.1 Syntax

```
compute ID group-ID pe keyword ...
```

- ID, group-ID are documented in *compute* command
- pe = style name of this compute command
- zero or more keywords may be appended

- keyword = *pair* or *bond* or *angle* or *dihedral* or *improper* or *k-space* or *fix*

## 18.72.2 Examples

```
compute 1 all pe
compute molPE all pe bond angle dihedral improper
```

## 18.72.3 Description

Define a computation that calculates the potential energy of the entire system of atoms. The specified group must be “all”. See the *compute pe/atom* command if you want per-atom energies. These per-atom values could be summed for a group of atoms via the *compute reduce* command.

The energy is calculated by the various pair, bond, etc. potentials defined for the simulation. If no extra keywords are listed, then the potential energy is the sum of pair, bond, angle, dihedral, improper, *k*-space (long-range), and fix energy (i.e., it is as though all the keywords were listed). If any extra keywords are listed, then only those components are summed to compute the potential energy.

The *k*-space contribution requires 1 extra FFT each timestep the energy is calculated, if using the PPPM solver via the *k-space\_style pppm* command. Thus it can increase the cost of the PPPM calculation if it is needed on a large fraction of the simulation timesteps.

Various fixes can contribute to the total potential energy of the system if the *fix* contribution is included. See the doc pages for *individual fixes* for details of which ones compute a potential energy.

---

**Note:** The *fix\_modify energy yes* command must also be specified if a fix is to contribute potential energy to this command.

---

A compute of this style with the ID of “thermo\_pe” is created when LAMMPS starts up, as if this command were in the input script:

```
compute thermo_pe all pe
```

See the “thermo\_style” command for more details.

---

## 18.72.4 Output info

This compute calculates a global scalar (the potential energy). This value can be used by any command that uses a global scalar value from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “extensive”. The scalar value will be in energy *units*.

## 18.72.5 Restrictions

none

## 18.72.6 Related commands

*compute pe/atom*

## 18.72.7 Default

none

# 18.73 compute pe/atom command

## 18.73.1 Syntax

```
compute ID group-ID pe/atom keyword ...
```

- ID, group-ID are documented in *compute* command
- pe/atom = style name of this compute command
- zero or more keywords may be appended
- keyword = *pair* or *bond* or *angle* or *dihedral* or *improper* or *k-space* or *fix*

## 18.73.2 Examples

```
compute 1 all pe/atom  
compute 1 all pe/atom pair  
compute 1 all pe/atom pair bond
```

## 18.73.3 Description

Define a computation that computes the per-atom potential energy for each atom in a group. See the *compute pe* command if you want the potential energy of the entire system.

The per-atom energy is calculated by the various pair, bond, etc potentials defined for the simulation. If no extra keywords are listed, then the potential energy is the sum of pair, bond, angle, dihedral, improper, *k*-space (long-range), and fix energy (i.e., it is as though all the keywords were listed). If any extra keywords are listed, then only those components are summed to compute the potential energy.

Note that the energy of each atom is due to its interaction with all other atoms in the simulation, not just with other atoms in the group.

For an energy contribution produced by a small set of atoms (e.g., 4 atoms in a dihedral or 3 atoms in a Tersoff 3-body interaction), that energy is assigned in equal portions to each atom in the set (e.g., 1/4 of the dihedral energy to each of the four atoms).

The *dihedral\_style charmm* style calculates pairwise interactions between 1–4 atoms. The energy contribution of these terms is included in the pair energy, not the dihedral energy.

The KSpace contribution is calculated using the method in (Heyes) for the Ewald method and a related method for PPPM, as specified by the `kpace_style pppm` command. For PPPM, the calculation requires 1 extra FFT each timestep that per-atom energy is calculated. This document describes how the long-range per-atom energy calculation is performed.

Various fixes can contribute to the per-atom potential energy of the system if the `fix` contribution is included. See the doc pages for *individual fixes* for details of which ones compute a per-atom potential energy.

---

**Note:** The `fix_modify energy yes` command must also be specified if a fix is to contribute per-atom potential energy to this command.

---

As an example of per-atom potential energy compared to total potential energy, these lines in an input script should yield the same result in the last 2 columns of thermo output:

```
compute      peratom all pe/atom
compute      pe all reduce sum c_peratom
thermo_style  custom step temp etotal press pe c_pe
```

---

**Note:** The per-atom energy does not include any Lennard-Jones tail corrections to the energy added by the `pair_modify tail yes` command, since those are contributions to the global system energy.

---

## 18.73.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The per-atom vector values will be in energy *units*.

## 18.73.5 Restrictions

## 18.73.6 Related commands

*compute pe, compute stress/atom*

## 18.73.7 Default

none

---

(Heyes) Heyes, Phys Rev B 49, 755 (1994),

## 18.74 compute plasticity/atom command

### 18.74.1 Syntax

```
compute ID group-ID plasticity/atom
```

- ID, group-ID are documented in compute command
- plasticity/atom = style name of this compute command

### 18.74.2 Examples

```
compute 1 all plasticity/atom
```

### 18.74.3 Description

Define a computation that calculates the per-atom plasticity for each atom in a group. This is a quantity relevant for *Peridynamics models*. See [this document](#) for an overview of LAMMPS commands for Peridynamics modeling.

The plasticity for a Peridynamic particle is the so-called consistency parameter ( $\lambda$ ). For elastic deformation,  $\lambda = 0$ , otherwise  $\lambda > 0$  for plastic deformation. For details, see ([Mitchell](#)) and the PDF doc included in the LAMMPS distribution in `doc/PDF/PDLammps_EPS.pdf`.

This command can be invoked for one of the Peridynamic *pair styles*: `peri/eps`.

The plasticity value will be 0.0 for atoms not in the specified compute group.

### 18.74.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values are unitless numbers  $\lambda \geq 0.0$ .

### 18.74.5 Restrictions

This compute is part of the PERI package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 18.74.6 Related commands

*compute damage/atom, compute dilatation/atom*

## 18.74.7 Default

none

(**Mitchell**) Mitchell, “A non-local, ordinary-state-based viscoelasticity model for peridynamics”, Sandia National Lab Report, 8064:1-28 (2011).

## 18.75 compute pressure command

### 18.75.1 Syntax

```
compute ID group-ID pressure temp-ID keyword ...
```

- ID, group-ID are documented in *compute* command
- pressure = style name of this compute command
- temp-ID = ID of compute that calculates temperature, can be NULL if not needed
- zero or more keywords may be appended
- keyword = *ke* or *pair* or *bond* or *angle* or *dihedral* or *improper* or *kpace* or *fix* or *virial* or *pair/hybrid*

### 18.75.2 Examples

```
compute 1 all pressure thermo_temp
compute 1 all pressure NULL pair bond
compute 1 all pressure NULL pair/hybrid lj/cut
```

### 18.75.3 Description

Define a computation that calculates the pressure of the entire system of atoms. The specified group must be “all”. See the *compute stress/atom* command if you want per-atom pressure (stress). These per-atom values could be summed for a group of atoms via the *compute reduce* command.

The pressure is computed by the formula

$$P = \frac{Nk_B T}{V} + \frac{1}{Vd} \sum_{i=1}^{N'} \vec{r}_i \cdot \vec{f}_i$$

where  $N$  is the number of atoms in the system (see discussion of DOF below),  $k_B$  is the Boltzmann constant,  $T$  is the temperature,  $d$  is the dimensionality of the system (2 for 2d, 3 for 3d), and  $V$  is the system volume (or area in 2d). The second term is the virial, equal to  $-dU/dV$ , computed for all pairwise as well as 2-body, 3-body, 4-body, many-body, and long-range interactions, where  $\vec{r}_i$  and  $\vec{f}_i$  are the position and force vector of atom  $i$ , and the dot indicates the dot product (scalar product). This is computed in parallel for each sub-domain and then summed over all parallel processes. Thus  $N'$  necessarily includes atoms from neighboring sub-domains (so-called ghost atoms) and the position and force vectors of ghost atoms are thus included in the summation. Only when running in serial and without periodic boundary conditions is  $N' = N$  the number of atoms in the system. *Fixes* that impose constraints (e.g., the *fix shake* command) may also contribute to the virial term.

A symmetric pressure tensor, stored as a 6-element vector, is also calculated by this compute. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ . The equation for the  $(I, J)$  components (where  $I$  and  $J$  are  $x$ ,  $y$ , or  $z$ ) is similar to the above formula, except that the first term uses components of the kinetic energy tensor and the second term uses components of the virial tensor:

$$P_{IJ} = \frac{1}{V} \sum_{k=1}^N m_k v_{k_I} v_{k_J} + \frac{1}{V} \sum_{k=1}^{N'} r_{k_I} f_{k_J}.$$

If no extra keywords are listed, the entire equations above are calculated. This includes a kinetic energy (temperature) term and the virial as the sum of pair, bond, angle, dihedral, improper, kspace (long-range), and fix contributions to the force on each atom. If any extra keywords are listed, then only those components are summed to compute temperature or ke and/or the virial. The *virial* keyword means include all terms except the kinetic energy *ke*.

The *pair/hybrid* keyword means to only include contribution from a sub-style in a *hybrid* or *hybrid/overlay* pair style.

Details of how LAMMPS computes the virial efficiently for the entire system, including for many-body potentials and accounting for the effects of periodic boundary conditions are discussed in (*Thompson*).

The temperature and kinetic energy tensor is not calculated by this compute, but rather by the temperature compute specified with the command. If the kinetic energy is not included in the pressure, then the temperature compute is not used and can be specified as NULL. Normally the temperature compute used by compute pressure should calculate the temperature of all atoms for consistency with the virial term, but any compute style that calculates temperature can be used (e.g., one that excludes frozen atoms or other degrees of freedom).

Note that if desired the specified temperature compute can be one that subtracts off a bias to calculate a temperature using only the thermal velocity of the atoms (e.g., by subtracting a background streaming velocity). See the doc pages for individual *compute commands* to determine which ones include a bias.

Also note that the  $N$  in the first formula above is really degrees-of-freedom divided by  $d =$  dimensionality, where the DOF value is calculated by the temperature compute. See the various *compute temperature* styles for details.

A compute of this style with the ID of thermo\_press is created when LAMMPS starts up, as if this command were in the input script:

```
compute thermo_press all pressure thermo_temp
```

where thermo\_temp is the ID of a similarly defined compute of style “temp”. See the *thermo\_style* command for more details.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---



## 18.75.4 Output info

This compute calculates a global scalar (the pressure) and a global vector of length 6 (pressure tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The ordering of values in the symmetric pressure tensor is as follows:  $p_{xx}$ ,  $p_{yy}$ ,  $p_{zz}$ ,  $p_{xy}$ ,  $p_{xz}$ ,  $p_{yz}$ .

The scalar and vector values calculated by this compute are “intensive”. The scalar and vector values will be in pressure *units*.

## 18.75.5 Restrictions

none

## 18.75.6 Related commands

*compute temp*, *compute stress/atom*, *thermo\_style*, *fix numdiff/virial*,

## 18.75.7 Default

none

---

(Thompson) Thompson, Plimpton, Mattson, J Chem Phys, 131, 154107 (2009).

# 18.76 compute pressure/uef command

## 18.76.1 Syntax

```
compute ID group-ID pressure/uef temp-ID keyword ...
```

- ID, group-ID are documented in *compute* command
- pressure/uef = style name of this compute command
- temp-ID = ID of compute that calculates temperature, can be NULL if not needed
- zero or more keywords may be appended
- keyword = *ke* or *pair* or *bond* or *angle* or *dihedral* or *improper* or *kspace* or *fix* or *virial*

## 18.76.2 Examples

```
compute 1 all pressure/uef my_temp_uef
compute 2 all pressure/uef my_temp_uef virial
```

### 18.76.3 Description

This command is used to compute the pressure tensor in the reference frame of the applied flow field when *fix nvt/uef* or *fix npt/uef* is used. It is not necessary to use this command to compute the scalar value of the pressure. A *compute pressure* may be used for that purpose.

The keywords and output information are documented in *compute\_pressure*.

### 18.76.4 Restrictions

This fix is part of the UEF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This command can only be used when *fix nvt/uef* or *fix npt/uef* is active.

The kinetic contribution to the pressure tensor will be accurate only when the compute specified by *temp-ID* is a *compute temp/uef*.

### 18.76.5 Related commands

*compute pressure*, *fix nvt/uef*, *compute temp/uef*

### 18.76.6 Default

none

## 18.77 compute property/atom command

### 18.77.1 Syntax

```
compute ID group-ID property/atom input1 input2 ...
```

- ID, group-ID are documented in *compute* command
- property/atom = style name of this compute command
- input = one or more atom attributes

```
possible attributes = id, mol, proc, type, mass,  
                    x, y, z, xs, ys, zs, xu, yu, zu, ix, iy, iz,  
                    vx, vy, vz, fx, fy, fz,  
                    q, mux, muy, muz, mu,  
                    spx, spy, spz, sp, fmx, fmy, fmz,  
                    nbonds,  
                    radius, diameter, omegax, omegay, omegaz,  
                    angmomx, angmomy, angmomz,  
                    shapex, shapey, shapez,  
                    quatw, quati, quatj, quatk, tqx, tqy, tqz,  
                    end1x, end1y, end1z, end2x, end2y, end2z,  
                    corner1x, corner1y, corner1z,  
                    corner2x, corner2y, corner2z,
```

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```

corner3x, corner3y, corner3z,
i_name, d_name, i2_name[I], d2_name[I],
vfrac, s0, espin, eradius, ervel, erforce,
rho, drho, e, de, cv, buckling,

```

```

id = atom ID
mol = molecule ID
proc = ID of processor that owns atom
type = atom type
mass = atom mass
x,y,z = unscaled atom coordinates
xs,ys,zs = scaled atom coordinates
xu,yu,zu = unwrapped atom coordinates
ix,iy,iz = box image that the atom is in
vx,vy,vz = atom velocities
fx,fy,fz = forces on atoms
q = atom charge
mux,muy,muz = orientation of dipole moment of atom
mu = magnitude of dipole moment of atom
spx, spy, spz = direction of the atomic magnetic spin
sp = magnitude of atomic magnetic spin moment
fmx, fmy, fmz = magnetic force
nbonds = number of bonds assigned to an atom
radius,diameter = radius,diameter of spherical particle
omegax,omegay,omegaz = angular velocity of spherical particle
angmomx,angmomy,angmomz = angular momentum of aspherical particle
shapex,shapey,shapex = 3 diameters of aspherical particle
quatw,quati,quatj,quatk = quaternion components for aspherical or body particles
tqx,tqy,tqz = torque on finite-size particles
end12x, end12y, end12z = end points of line segment
corner123x, corner123y, corner123z = corner points of triangle
i_name = custom integer vector with name
d_name = custom floating point vector with name
i2_name[I] = Ith column of custom integer array with name
d2_name[I] = Ith column of custom floating-point array with name

```

```

PERI package per-atom properties:
vfrac = volume fraction
s0 = max stretch of any bond a particle is part of

```

```

EFF and AWPMD package per-atom properties:
espin = electron spin
eradius = electron radius
eravel = electron radial velocity
erforce = electron radial force

```

```

SPH package per-atom properties:
rho = density of SPH particles
drho = change in density
e = energy
de = change in thermal energy
cv = heat capacity

```

MESONT package per-atom properties:  
 buckling = buckling flag used in mesoscopic simulation of nanotubes

## 18.77.2 Examples

```
compute 1 all property/atom xs vx fx mux
compute 2 all property/atom type
compute 1 all property/atom ix iy iz
compute 3 all property/atom sp spx spy spz
compute 1 all property/atom i_myFlag d_Sxyz[1] d_Sxyz[3]
```

## 18.77.3 Description

Define a computation that simply stores atom attributes for each atom in the group. This is useful so that the values can be used by other *output commands* that take computes as inputs. See for example, the *compute reduce*, *fix ave/atom*, *fix ave/histo*, *fix ave/chunk*, and *atom-style variable* commands.

The list of possible attributes is essentially the same as that used by the *dump custom* command, which describes their meaning, with some additional quantities that are only defined for certain *atom styles*. The goal of this augmented list gives an input script access to any per-atom quantity stored by LAMMPS.

The values are stored in a per-atom vector or array as discussed below. Zeroes are stored for atoms not in the specified group or for quantities that are not defined for a particular particle in the group (e.g., *shapex* if the particle is not an ellipsoid).

Attributes *i\_name*, *d\_name*, *i2\_name*, *d2\_name* refer to custom per-atom integer and floating-point vectors or arrays that have been added via the *fix property/atom* command. When that command is used specific names are given to each attribute which are the “name” portion of these attributes. For arrays *i2\_name* and *d2\_name*, the column of the array must also be included following the name in brackets (e.g., *d2\_xyz[2]* or *i2\_mySpin[3]*).

The additional quantities only accessible via this command, and not directly via the *dump custom* command, are as follows.

*Nbonds* is available for all molecular atom styles and refers to the number of explicit bonds assigned to an atom. Note that if the *newton bond* command is set to *on*, which is the default, then every bond in the system is assigned to only one of the two atoms in the bond. Thus a bond between atoms *I* and *J* may be tallied for either atom *I* or atom *J*. If *newton bond off* is set, it will be tallied with both atom *I* and atom *J*.

The quantities *shapex*, *shapey*, and *shapex* are defined for ellipsoidal particles and define the 3d shape of each particle.

The quantities *quatw*, *quati*, *quatj*, and *quatk* are defined for ellipsoidal particles and body particles and store the 4-vector quaternion representing the orientation of each particle. See the *set* command for an explanation of the quaternion vector.

*End1x*, *end1y*, *end1z*, *end2x*, *end2y*, *end2z*, are defined for line segment particles and define the end points of each line segment.

*Corner1x*, *corner1y*, *corner1z*, *corner2x*, *corner2y*, *corner2z*, *corner3x*, *corner3y*, *corner3z*, are defined for triangular particles and define the corner points of each triangle.

In addition, the various per-atom quantities listed above for specific packages are only accessible by this command.

Changed in version 15Sep2022: The *espin* property was previously called *spin*.

## 18.77.4 Output info

This compute calculates a per-atom vector or per-atom array depending on the number of input values. If a single input is specified, a per-atom vector is produced. If two or more inputs are specified, a per-atom array is produced where the number of columns = the number of inputs. The vector or array can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector or array values will be in whatever *units* the corresponding attribute is in (e.g., velocity units for *vx*, charge units for *q*).

For the spin quantities, *sp* is in the units of the Bohr magneton; *spx*, *spy*, and *spz* are unitless quantities; and *fm<sub>x</sub>*, *fm<sub>y</sub>*, and *fm<sub>z</sub>* are given in rad/THz.

## 18.77.5 Restrictions

none

## 18.77.6 Related commands

*dump custom*, *compute reduce*, *fix ave/atom*, *fix ave/chunk*, *fix property/atom*

## 18.77.7 Default

none

# 18.78 compute property/chunk command

## 18.78.1 Syntax

```
compute ID group-ID property/chunk chunkID input1 input2 ...
```

- ID, group-ID are documented in *compute* command
- property/chunk = style name of this compute command
- input = one or more attributes

attributes = count, id, coord1, coord2, coord3

count = # of atoms in chunk

id = original chunk IDs before compression by *compute chunk/atom*

coord123 = coordinates for spatial bins calculated by *compute chunk/atom*

## 18.78.2 Examples

```
compute 1 all property/chunk count
compute 1 all property/chunk ID coord1
```

## 18.78.3 Description

Define a computation that stores the specified attributes of chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

This compute calculates and stores the specified attributes of chunks as global data so they can be accessed by other *output commands* and used in conjunction with other commands that generate per-chunk data, such as *compute com/chunk* or *compute msd/chunk*.

Note that only atoms in the specified group contribute to the calculation of the *count* attribute. The *compute chunk/atom* command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

The *count* attribute is the number of atoms in the chunk.

The *id* attribute stores the original chunk ID for each chunk. It can only be used if the *compress* keyword was set to *yes* for the *compute chunk/atom* command referenced by chunkID. This means that the original chunk IDs (e.g., molecule IDs) will have been compressed to remove chunk IDs with no atoms assigned to them. Thus a compressed chunk ID of 3 may correspond to an original chunk ID (molecule ID in this case) of 415. The *id* attribute will then be 415 for the third chunk.

The *coordN* attributes can only be used if a *binning* style was used in the *compute chunk/atom* command referenced by chunkID. For *bin/1d*, *bin/2d*, and *bin/3d* styles the attribute is the center point of the bin in the corresponding dimension. Style *bin/1d* only defines a *coord1* attribute. Style *bin/2d* adds a *coord2* attribute. Style *bin/3d* adds a *coord3* attribute.

Note that if the value of the *units* keyword used in the *compute chunk/atom command* is *box* or *lattice*, the *coordN* attributes will be in distance *units*. If the value of the *units* keyword is *reduced*, the *coordN* attributes will be in unitless reduced units (0–1).

The simplest way to output the results of the compute property/chunk calculation to a file is to use the *fix ave/time* command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk1 all property/chunk cc1 count
compute myChunk2 all com/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk1 c_myChunk2[*] file tmp.out mode vector
```

## 18.78.4 Output info

This compute calculates a global vector or global array depending on the number of input values. The length of the vector or number of rows in the array is the number of chunks.

This compute calculates a global vector or global array where the number of rows = the number of chunks *Nchunk* as calculated by the specified *compute chunk/atom* command. If a single input is specified, a global vector is produced. If two or more inputs are specified, a global array is produced where the number of columns = the number of inputs. The vector or array can be accessed by any command that uses global values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector or array values are “intensive”. The values will be unitless or in the units discussed above.

## 18.78.5 Restrictions

none

## 18.78.6 Related commands

*fix ave/chunk*

## 18.78.7 Default

none

# 18.79 compute property/local command

## 18.79.1 Syntax

```
compute ID group-ID property/local attribute1 attribute2 ... keyword args ...
```

- ID, group-ID are documented in *compute* command
- property/local = style name of this compute command
- one or more attributes of the same type (neighbor, pair, bond, angle, dihedral, or improper) may be appended

```
possible attributes = natom1, natom2, ntype1, ntype2,
                    patom1, patom2, ptype1, ptype2,
                    batom1, batom2, btype,
                    aatom1, aatom2, aatom3, atype,
                    datom1, datom2, datom3, datom4, dtype,
                    iatom1, iatom2, iatom3, iatom4, itype
```

– Neighbor attributes

```
natom1, natom2 = store IDs of 2 atoms in each pair (within neighbor cutoff)
ntype1, ntype2 = store types of 2 atoms in each pair (within neighbor cutoff)
```

– Pair attributes

```
patom1, patom2 = store IDs of 2 atoms in each pair (within force cutoff)
ptype1, ptype2 = store types of 2 atoms in each pair (within force cutoff)
```

– Bond attributes

```
batom1, batom2 = store IDs of 2 atoms in each bond
btype = store bond type of each bond
```

– Angle attributes

```
aatom1, aatom2, aatom3 = store IDs of 3 atoms in each angle
atype = store angle type of each angle
```

– Dihedral attributes

```
datom1, datom2, datom3, datom4 = store IDs of 4 atoms in each dihedral
dtype = store dihedral type of each dihedral
```

– Improper attributes

```
iatom1, iatom2, iatom3, iatom4 = store IDs of 4 atoms in each improper
itype = store improper type of each improper
```

- zero or more keyword/arg pairs may be appended
- keyword = *cutoff*  
*cutoff* arg = *type* or *radius*

## 18.79.2 Examples

```
compute 1 all property/local btype batom1 batom2
compute 1 all property/local atype aatom2
```

## 18.79.3 Description

Define a computation that stores the specified attributes as local data so it can be accessed by other *output commands*. If the input attributes refer to bond information, then the number of datums generated, aggregated across all processors, equals the number of bonds in the system. Ditto for pairs, angles, etc.

If multiple attributes are specified then they must all generate the same amount of information, so that the resulting local array has the same number of rows for each column. This means that only bond attributes can be specified together, or angle attributes, etc. Bond and angle attributes cannot be mixed in the same compute property/local command.

If the inputs are pair attributes, the local data is generated by looping over the pairwise neighbor list. Info about an individual pairwise interaction will only be included if both atoms in the pair are in the specified compute group. For *natom1* and *natom2*, all atom pairs in the neighbor list are considered (out to the neighbor cutoff = force cutoff + *neighbor skin*). For *patom1* and *patom2*, the distance between the atoms must be less than the force cutoff distance for that pair to be included, as defined by the *pair\_style* and *pair\_coeff* commands.

The optional *cutoff* keyword determines how the force cutoff distance for an interaction is determined for the *patom1* and *patom2* attributes. For the default setting of *type*, the pairwise cutoff defined by the *pair\_style* command for the types of the two atoms is used. For the *radius* setting, the sum of the radii of the two particles is used as a cutoff. For example, this is appropriate for granular particles which only interact when they are overlapping, as computed



by *granular pair styles*. Note that if a granular model defines atom types such that all particles of a specific type are monodisperse (same diameter), then the two settings are effectively identical.

If the inputs are bond, angle, etc attributes, the local data is generated by looping over all the atoms owned on a processor and extracting bond, angle, etc info. For bonds, info about an individual bond will only be included if both atoms in the bond are in the specified compute group. Likewise for angles, dihedrals, etc.

For bonds and angles, a bonds/angles that have been broken by setting their bond/angle type to 0 will not be included. Bonds/angles that have been turned off (see the *fix shake* or *delete\_bonds* commands) by setting their bond/angle type negative are written into the file. This is consistent with the *compute bond/local* and *compute angle/local* commands

Note that as atoms migrate from processor to processor, there will be no consistent ordering of the entries within the local vector or array from one timestep to the next. The only consistency that is guaranteed is that the ordering on a particular timestep will be the same for local vectors or arrays generated by other compute commands. For example, output from the *compute bond/local* command can be combined with bond atom indices from this command and output by the *dump local* command in a consistent way.

The *natom1* and *natom2* or *patom1* and *patom2* attributes refer to the atom IDs of the 2 atoms in each pairwise interaction computed by the *pair\_style* command. The *ntype1* and *ntype2* or *ptype1* and *ptype2* attributes refer to the atom types of the 2 atoms in each pairwise interaction.

---

**Note:** For pairs, if two atoms  $I, J$  are involved in 1–2, 1–3, 1–4 interactions within the molecular topology, their pairwise interaction may be turned off, and thus they may not appear in the neighbor list, and will not be part of the local data created by this command. More specifically, this may be true of  $I, J$  pairs with a weighting factor of 0.0; pairs with a non-zero weighting factor are included. The weighting factors for 1–2, 1–3, and 1–4 pairwise interactions are set by the *special\_bonds* command.

---

The *batom1* and *batom2* attributes refer to the atom IDs of the 2 atoms in each *bond*. The *btype* attribute refers to the type of the bond, from 1 to  $Nbtypes = \#$  of bond types. The number of bond types is defined in the data file read by the *read\_data* command.

The attributes that start with “a”, “d”, and “i” refer to similar values for *angles*, *dihedrals*, and *impropers*.

## 18.79.4 Output info

This compute calculates a local vector or local array depending on the number of input values. The length of the vector or number of rows in the array is the number of bonds, angles, etc. If a single input is specified, a local vector is produced. If two or more inputs are specified, a local array is produced where the number of columns = the number of inputs. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector or array values will be integers that correspond to the specified attribute.

## 18.79.5 Restrictions

none

## 18.79.6 Related commands

*dump local*, *compute reduce*

## 18.79.7 Default

The keyword default is cutoff = type.

# 18.80 compute ptm/atom command

## 18.80.1 Syntax

```
compute ID group-ID ptm/atom structures threshold group2-ID
```

- ID, group-ID are documented in *compute* command
- ptm/atom = style name of this compute command
- structures = *default* or *all* or any hyphen-separated combination of *fcc*, *hcp*, *bcc*, *ico*, *sc*, *dcub*, *dhex*, or *graphene*  
= structure types to search for
- threshold = lattice distortion threshold (RMSD)
- group2-ID determines which group is used for neighbor selection (optional, default “all”)

## 18.80.2 Examples

```
compute 1 all ptm/atom default 0.1 all
compute 1 all ptm/atom fcc-hcp-dcub-dhex 0.15 all
compute 1 all ptm/atom all 0
```

## 18.80.3 Description

Define a computation that determines the local lattice structure around an atom using the PTM (Polyhedral Template Matching) method. The PTM method is described in (*Larsen*).

Currently, there are seven lattice structures PTM recognizes:

- fcc = 1
- hcp = 2
- bcc = 3
- ico (icosahedral) = 4
- sc (simple cubic) = 5
- dcub (diamond cubic) = 6
- dhex (diamond hexagonal) = 7
- graphene = 8

The value of the PTM structure will be 0 for unknown types and  $-1$  for atoms not in the specified compute group. The choice of structures to search for can be specified using the “structures” argument, which is a hyphen-separated list of structure keywords. Two convenient pre-set options are provided:

- default: fcc-hcp-bcc-ico
- all: fcc-hcp-bcc-ico-sc-dcub-dhex-graphene

The ‘default’ setting detects the same structures as the Common Neighbor Analysis method. The ‘all’ setting searches for all structure types. A performance penalty is incurred for the diamond and graphene structures, so it is not recommended to use this option if it is known that the simulation does not contain these structures.

PTM identifies structures using two steps. First, a graph isomorphism test is used to identify potential structure matches. Next, the deviation is computed between the local structure (in the simulation) and a template of the ideal lattice structure. The deviation is calculated as:

$$\text{RMSD}(\mathbf{u}, \mathbf{v}) = \min_{s, \mathbf{Q}} \sqrt{\frac{1}{N} \sum_{i=1}^N \|s[\vec{u}_i - \bar{\mathbf{u}}] - \mathbf{Q} \cdot \vec{v}_i\|^2}$$

Here,  $\vec{u}$  and  $\vec{v}$  contain the coordinates of the local and ideal structures respectively,  $s$  is a scale factor, and  $\mathbf{Q}$  is a rotation. The best match is identified by the lowest RMSD value, using the optimal scaling, rotation, and correspondence between the points.

The *threshold* keyword sets an upper limit on the maximum permitted deviation before a local structure is identified as disordered. Typical values are in the range 0.1–0.15, but larger values may be desirable at higher temperatures. A value of 0 is equivalent to infinity and can be used if no threshold is desired.

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (e.g., each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently or to have multiple compute/dump commands, each with a *ptm/atom* style. By default the compute processes **all** neighbors unless the optional *group2-ID* argument is given, then only members of that group are considered as neighbors.

## 18.80.4 Output info

This compute calculates a per-atom array, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

Results are stored in the per-atom array in the following order:

- type
- rmsd
- interatomic distance
- qw
- qx
- qy
- qz

The type is a number from  $-1$  to 8. The rmsd is a positive real number. The interatomic distance is computed from the scale factor in the RMSD equation. The  $(qw, qx, qy, qz)$  parameters represent the orientation of the local structure in quaternion form. The reference coordinates for each template (from which the orientation is determined) can be found in the *ptm\_constants.h* file in the PTM source directory. For atoms that are not within the compute group-ID, all values are set to zero.

## 18.80.5 Restrictions

This fix is part of the PTM package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 18.80.6 Related commands

*compute centro/atom compute cna/atom*

## 18.80.7 Default

none

---

(Larsen) Larsen, Schmidt, Schiotz, Modelling Simul Mater Sci Eng, 24, 055007 (2016).

## 18.81 compute rdf command

### 18.81.1 Syntax

```
compute ID group-ID rdf Nbin itype1 jtype1 itype2 jtype2 ... keyword/value ...
```

- ID, group-ID are documented in *compute* command
- rdf = style name of this compute command
- Nbin = number of RDF bins
- itypeN = central atom type for Nth RDF histogram (see asterisk form below)
- jtypeN = distribution atom type for Nth RDF histogram (see asterisk form below)
- zero or more keyword/value pairs may be appended
- keyword = *cutoff*

*cutoff* value = Rcut

Rcut = cutoff distance for RDF computation (distance units)

### 18.81.2 Examples

```
compute 1 all rdf 100
compute 1 all rdf 100 1 1
compute 1 all rdf 100 * 3 cutoff 5.0
compute 1 fluid rdf 500 1 1 1 2 2 1 2 2
compute 1 fluid rdf 500 1*3 2 5 *10 cutoff 3.5
```

### 18.81.3 Description

Define a computation that calculates the radial distribution function (RDF), also called  $g(r)$ , and the coordination number for a group of particles. Both are calculated in histogram form by binning pairwise distances into  $N_{bin}$  bins from 0.0 to the maximum force cutoff defined by the *pair\_style* command or the cutoff distance  $R_{cut}$  specified via the *cutoff* keyword. The bins are of uniform size in radial distance. Thus a single bin encompasses a thin shell of distances in 3d and a thin ring of distances in 2d.

---

**Note:** If you have a bonded system, then the settings of *special\_bonds* command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the *special\_bonds* command, and means those pairwise interactions do not appear in the neighbor list. Because this fix uses a neighbor list, it also means those pairs will not be included in the RDF. This does not apply when using long-range coulomb interactions (*coul/long*, *coul/msm*, *coul/wolf* or similar). One way to get around this would be to set special\_bond scaling factors to very tiny numbers that are not exactly zero (e.g.,  $1.0 \times 10^{-50}$ ). Another workaround is to write a dump file, and use the *rerun* command to compute the RDF for snapshots in the dump file. The rerun script can use a *special\_bonds* command that includes all pairs in the neighbor list.

---

By default the RDF is computed out to the maximum force cutoff defined by the *pair\_style* command. If the *cutoff* keyword is used, then the RDF is computed accurately out to the  $R_{cut} > 0.0$  distance specified.

---

**Note:** Normally, you should only use the *cutoff* keyword if no pair style is defined (e.g., the *rerun* command is being used to post-process a dump file of snapshots) or if you really want the RDF for distances beyond the *pair\_style* force cutoff and cannot easily post-process a dump file to calculate it. This is because using the *cutoff* keyword incurs extra computation and possibly communication, which may slow down your simulation. If you specify  $R_{cut} \leq$  force cutoff, you will force an additional neighbor list to be built at every timestep this command is invoked (or every reneighboring timestep, whichever is less frequent), which is inefficient. LAMMPS will warn you if this is the case. If you specify a  $R_{cut} >$  force cutoff, you must insure ghost atom information out to  $R_{cut} + skin$  is communicated, via the *comm\_modify cutoff* command, else the RDF computation cannot be performed, and LAMMPS will give an error message. The *skin* value is what is specified with the *neighbor* command. In this case, you are forcing a large neighbor list to be built just for the RDF computation, and extra communication to be performed every timestep.

---

The *itypeN* and *jtypeN* arguments are optional. These arguments must come in pairs. If no pairs are listed, then a single histogram is computed for  $g(r)$  between all atom types. If one or more pairs are listed, then a separate histogram is generated for each *itype,jtype* pair.

The *itypeN* and *jtypeN* settings can be specified in one of two ways. An explicit numeric value can be used, as in the fourth example above. Or a wild-card asterisk can be used to specify a range of atom types. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the number of atom types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $m$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive).

If both *itypeN* and *jtypeN* are single values, as in the fourth example above, this means that a  $g(r)$  is computed where atoms of type *itypeN* are the central atom, and atoms of type *jtypeN* are the distribution atom. If either *itypeN* and *jtypeN* represent a range of values via the wild-card asterisk, as in the fifth example above, this means that a  $g(r)$  is computed where atoms of any of the range of types represented by *itypeN* are the central atom, and atoms of any of the range of types represented by *jtypeN* are the distribution atom.

Pairwise distances are generated by looping over a pairwise neighbor list, just as they would be in a *pair\_style* computation. The distance between two atoms  $I$  and  $J$  is included in a specific histogram if the following criteria are met:

- atoms  $I$  and  $J$  are both in the specified compute group
- the distance between atoms  $I$  and  $J$  is less than the maximum force cutoff

- the type of the  $I$  atom matches  $itypeN$  (one or a range of types)
- the type of the  $J$  atom matches  $jtypeN$  (one or a range of types)

It is OK if a particular pairwise distance is included in more than one individual histogram, due to the way the  $itypeN$  and  $jtypeN$  arguments are specified.

The  $g(r)$  value for a bin is calculated from the histogram count by scaling it by the idealized number of how many counts there would be if atoms of type  $jtypeN$  were uniformly distributed. Thus it involves the count of  $itypeN$  atoms, the count of  $jtypeN$  atoms, the volume of the entire simulation box, and the volume of the bin's thin shell in 3d (or the area of the bin's thin ring in 2d).

A coordination number  $coord(r)$  is also calculated, which is the number of atoms of type  $jtypeN$  within the current bin or closer, averaged over atoms of type  $itypeN$ . This is calculated as the area- or volume-weighted sum of  $g(r)$  values over all bins up to and including the current bin, multiplied by the global average volume density of atoms of type  $jtypeN$ .

The simplest way to output the results of the compute rdf calculation to a file is to use the *fix ave/time* command, for example:

```
compute myRDF all rdf 50
fix 1 all ave/time 100 1 100 c_myRDF[*] file tmp.rdf mode vector
```

## 18.81.4 Output info

This compute calculates a global array in which the number of rows is  $Nbins$  and the number of columns is  $1 + 2N_{pairs}$ , where  $N_{pairs}$  is the number of  $I, J$  pairings specified. The first column has the bin coordinate (center of the bin), and each successive set of two columns has the  $g(r)$  and  $coord(r)$  values for a specific set of  $itypeN$  versus  $jtypeN$  interactions, as described above. These values can be used by any command that uses a global values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The array values calculated by this compute are all “intensive”.

The first column of array values will be in distance *units*. The  $g(r)$  columns of array values are normalized numbers  $\geq 0.0$ . The coordination number columns of array values are also numbers  $\geq 0.0$ .

## 18.81.5 Restrictions

The RDF is not computed for distances longer than the force cutoff, since processors (in parallel) do not know about atom coordinates for atoms further away than that distance. If you want an RDF for larger distances, you can use the *rerun* command to post-process a dump file and set the cutoff for the potential to be longer in the rerun script. Note that in the rerun context, the force cutoff is arbitrary, since you are not running dynamics and thus are not changing your model. The definition of  $g(r)$  used by LAMMPS is only appropriate for characterizing atoms that are uniformly distributed throughout the simulation cell. In such cases, the coordination number is still correct and meaningful. As an example, if a large simulation cell contains only one atom of type  $itypeN$  and one of  $jtypeN$ , then  $g(r)$  will register an arbitrarily large spike at whatever distance they happen to be at, and zero everywhere else. The function  $coord(r)$  will show a step change from zero to one at the location of the spike in  $g(r)$ .

---

**Note:** compute rdf can handle dynamic groups and systems where atoms are added or removed, but this causes that certain normalization parameters need to be re-computed in every step and include collective communication operations. This will reduce performance and limit parallel efficiency and scaling. For systems, where only the type of atoms changes (e.g., when using *fix atom/swap*), you need to explicitly request the dynamic normalization updates via *compute\_modify dynamic yes*

---

## 18.81.6 Related commands

*fix ave/time, compute\_modify, compute adf*

## 18.81.7 Default

The keyword defaults are cutoff = 0.0 (use the pairwise force cutoff).

## 18.82 compute reduce command

## 18.83 compute reduce/region command

### 18.83.1 Syntax

```
compute ID group-ID style arg mode input1 input2 ... keyword args ...
```

- ID, group-ID are documented in *compute* command
- style = *reduce* or *reduce/region*
  - reduce* arg = none
  - reduce/region* arg = region-ID
  - region-ID = ID of region to use for choosing atoms
- mode = *sum* or *min* or *max* or *ave* or *sumsq* or *avesq* or *sumabs* or *aveabs*
- one or more inputs can be listed
- input = *x* or *y* or *z* or *vx* or *vy* or *vz* or *fx* or *fy* or *fz* or *c\_ID* or *c\_ID[N]* or *f\_ID* or *f\_ID[N]* or *v\_name*
  - x, y, z, vx, vy, vz, fx, fy, fz* = atom attribute (position, velocity, force component)
  - c\_ID* = per-atom or local vector calculated by a compute with ID
  - c\_ID[I]* = Ith column of per-atom or local array calculated by a compute with ID, I can include wildcard (see below)
  - f\_ID* = per-atom or local vector calculated by a fix with ID
  - f\_ID[I]* = Ith column of per-atom or local array calculated by a fix with ID, I can include wildcard (see below)
  - v\_name* = per-atom vector calculated by an atom-style variable with name
- zero or more keyword/args pairs may be appended
- keyword = *replace*
  - replace* args = *vec1 vec2*
  - vec1* = reduced value from this input vector will be replaced
  - vec2* = replace it with *vec1[N]* where N is index of max/min value from *vec2*





appended, the vector calculated by the compute is used. If a bracketed integer is appended, the  $I$ th column of the array calculated by the compute is used. Users can also write code for their own compute styles and *add them to LAMMPS*. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

If a value begins with “f\_”, a fix ID must follow which has been previously defined in the input script. Fixes can generate per-atom or local quantities. See the individual *fix* page for details. Note that some fixes only produce their values on certain timesteps, which must be compatible with when compute reduce references the values, else an error results. If no bracketed integer is appended, the vector calculated by the fix is used. If a bracketed integer is appended, the  $I$ th column of the array calculated by the fix is used. Users can also write code for their own fix style and *add them to LAMMPS*. See the discussion above for how  $I$  can be specified with a wildcard asterisk to effectively specify multiple values.

If a value begins with “v\_”, a variable name must follow which has been previously defined in the input script. It must be an *atom-style variable*. Atom-style variables can reference thermodynamic keywords and various per-atom attributes, or invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of generating per-atom quantities to reduce.

If the *replace* keyword is used, two indices *vec1* and *vec2* are specified, where each index ranges from 1 to the number of input values. The *replace* keyword can only be used if the *mode* is *min* or *max*. It works as follows. A min/max is computed as usual on the *vec2* input vector. The index  $N$  of that value within *vec2* is also stored. Then, instead of performing a min/max on the *vec1* input vector, the stored index is used to select the  $N$ th element of the *vec1* vector.

Thus, for example, if you wish to use this compute to find the bond with maximum stretch, you can do it as follows:

```
compute 1 all property/local batom1 batom2
compute 2 all bond/local dist
compute 3 all reduce max c_1[1] c_1[2] c_2 replace 1 3 replace 2 3
thermo_style custom step temp c_3[1] c_3[2] c_3[3]
```

The first two input values in the compute reduce command are vectors with the IDs of the 2 atoms in each bond, using the *compute property/local* command. The last input value is bond distance, using the *compute bond/local* command. Instead of taking the max of the two atom ID vectors, which does not yield useful information in this context, the *replace* keywords will extract the atom IDs for the two atoms in the bond of maximum stretch. These atom IDs and the bond stretch will be printed with thermodynamic output.

If a single input is specified this compute produces a global scalar value. If multiple inputs are specified, this compute produces a global vector of values, the length of which is equal to the number of inputs specified.

As discussed below, for the *sum*, *sumabs*, and *sumsq* modes, the value(s) produced by this compute are all “extensive”, meaning their value scales linearly with the number of atoms involved. If normalized values are desired, this compute can be accessed by the *thermo\_style custom* command with *thermo\_modify norm yes* set as an option. Or it can be accessed by a *variable* that divides by the appropriate atom count.

### 18.83.4 Output info

This compute calculates a global scalar if a single input value is specified or a global vector of length  $N$ , where  $N$  is the number of inputs, and which can be accessed by indices 1 to  $N$ . These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

All the scalar or vector values calculated by this compute are “intensive”, except when the *sum*, *sumabs*, or *sumsq* modes are used on per-atom or local vectors, in which case the calculated values are “extensive”.

The scalar or vector values will be in whatever *units* the quantities being reduced are in.

### 18.83.5 Restrictions

none

### 18.83.6 Related commands

*compute*, *fix*, *variable*

### 18.83.7 Default

none

## 18.84 compute reduce/chunk command

### 18.84.1 Syntax

```
compute ID group-ID reduce/chunk chunkID mode input1 input2 ...
```

- ID, group-ID are documented in *compute* command
- reduce/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command
- mode = *sum* or *min* or *max*
- one or more inputs can be listed
- input = c\_ID, c\_ID[N], f\_ID, f\_ID[N], v\_ID

```
c_ID = per-atom vector calculated by a compute with ID
c_ID[I] = Ith column of per-atom array calculated by a compute with ID, I can
↔include wildcard (see below)
f_ID = per-atom vector calculated by a fix with ID
f_ID[I] = Ith column of per-atom array calculated by a fix with ID, I can include
↔wildcard (see below)
v_name = per-atom vector calculated by an atom-style variable with name
```

## 18.84.2 Examples

```
compute 1 all reduce/chunk mychunk min c_cluster
```

## 18.84.3 Description

Define a calculation that reduces one or more per-atom vectors into per-chunk values. This can be useful for diagnostic output. Or when used in conjunction with the `compute chunk/spread/atom` command it can be used to create per-atom values that induce a new set of chunks with a second `compute chunk/atom` command. An example is given below.

In LAMMPS, chunks are collections of atoms defined by a `compute chunk/atom` command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as `chunkID`. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the `compute chunk/atom` and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

For each atom, this compute accesses its chunk ID from the specified `chunkID` compute. The per-atom value from an input contributes to a per-chunk value corresponding the the chunk ID.

The reduction operation is specified by the `mode` setting and is performed over all the per-atom values from the atoms in each chunk. The `sum` option adds the pre-atom values to a per-chunk total. The `min` or `max` options find the minimum or maximum value of the per-atom values for each chunk.

Note that only atoms in the specified group contribute to the reduction operation. If the `chunkID` compute returns a 0 for the chunk ID of an atom (i.e., the atom is not in a chunk defined by the `compute chunk/atom` command), that atom will also not contribute to the reduction operation. An input that is a compute or fix may define its own group which affects the quantities it returns. For example, a compute with return a zero value for atoms that are not in the group specified for that compute.

Each listed input is operated on independently. Each input can be the result of a `compute` or `fix` or the evaluation of an atom-style *variable*.

Note that for values from a compute or fix, the bracketed index `I` can be specified using a wildcard asterisk with the index to effectively specify multiple values. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the size of the vector (for `mode = scalar`) or the number of columns in the array (for `mode = vector`), then an asterisk with no numeric values means all indices from 1 to  $N$ . A leading asterisk means all indices from 1 to  $n$  (inclusive). A trailing asterisk means all indices from  $n$  to  $N$  (inclusive). A middle asterisk means all indices from  $m$  to  $n$  (inclusive).

Using a wildcard is the same as if the individual columns of the array had been listed one by one. For example, the following two compute reduce/chunk commands are equivalent, since the `compute property/chunk` command creates a per-atom array with 3 columns:

```
compute prop all property/atom vx vy vz
compute 10 all reduce/chunk mychunk max c_prop[*]
compute 10 all reduce/chunk mychunk max c_prop[1] c_prop[2] c_prop[3]
```

Here is an example of using this compute, in conjunction with the `compute chunk/spread/atom` command to identify self-assembled micelles. The commands below can be added to the examples/in.micelle script.

Imagine a collection of polymer chains or small molecules with hydrophobic end groups. All the hydrophobic (HP) atoms are assigned to a group called “phobic”.

These commands will assign a unique cluster ID to all HP atoms within a specified distance of each other. A cluster will contain all HP atoms in a single molecule, but also the HP atoms in nearby molecules (e.g., molecules that have clumped to form a micelle due to the attraction induced by the hydrophobicity). The output of the chunk/reduce command will be a cluster ID per chunk (molecule). Molecules with the same cluster ID are in the same micelle.

```
group phobic type 4      # specific to in.micelle model
compute cluster phobic cluster/atom 2.0
compute cmol all chunk/atom molecule
compute reduce phobic reduce/chunk cmol min c_cluster
```

This per-chunk info could be output in at least two ways:

```
fix 10 all ave/time 1000 1 1000 c_reduce file tmp.phobic mode vector

compute spread all chunk/spread/atom cmol c_reduce
dump 1 all custom 1000 tmp.dump id type mol x y z c_cluster c_spread
dump_modify 1 sort id
```

In the first case, each snapshot in the tmp.phobic file will contain one line per molecule. Molecules with the same value are in the same micelle. In the second case each dump snapshot contains all atoms, each with a final field with the cluster ID of the micelle that the HP atoms of that atom's molecule belong to.

The result from compute chunk/spread/atom can be used to define a new set of chunks, where all the atoms in all the molecules in the same micelle are assigned to the same chunk (i.e., one chunk per micelle).

```
compute micelle all chunk/atom c_spread compress yes
```

Further analysis on a per-micelle basis can now be performed using any of the per-chunk computes listed on the [Howto chunk](#) doc page (e.g., count the number of atoms in each micelle, calculate its center or mass, shape/moments of inertia, and radius of gyration).

```
compute prop all property/chunk micelle count
fix 20 all ave/time 1000 1 1000 c_prop file tmp.micelle mode vector
```

Each snapshot in the tmp.micelle file will have one line per micelle with its count of atoms, plus a first line for a chunk with all the solvent atoms. By the time 50000 steps have elapsed, there are a handful of large micelles.

---

## 18.84.4 Output info

This compute calculates a global vector if a single input value is specified, otherwise a global array is output. The number of columns in the array is the number of inputs provided. The length of the vector or the number of vector elements or array rows = the number of chunks *N<sub>chunk</sub>* as calculated by the specified [compute chunk/atom](#) command. The vector or array can be accessed by any command that uses global values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom values for the vector or each column of the array will be in whatever *units* the corresponding input value is in. The vector or array values are “intensive”.

## 18.84.5 Restrictions

none

## 18.84.6 Related commands

*compute chunk/atom, compute reduce, compute chunk/spread/atom*

## 18.84.7 Default

none

# 18.85 compute rigid/local command

## 18.85.1 Syntax

```
compute ID group-ID rigid/local rigidID input1 input2 ...
```

- ID, group-ID are documented in *compute* command
- rigid/local = style name of this compute command
- rigidID = ID of fix rigid/small command or one of its variants
- input = one or more rigid body attributes

```
possible attributes = id, mol, mass,
                    x, y, z, xu, yu, zu, ix, iy, iz
                    vx, vy, vz, fx, fy, fz,
                    omegax, omegay, omegaz,
                    angmomx, angmomy, angmomz,
                    quatw, quati, quatj, quatk,
                    tqx, tqy, tqz,
                    inertiax, inertiay, inertiaz
id = atom ID of atom within body which owns body properties
mol = molecule ID used to define body in fix rigid/small command
mass = total mass of body
x,y,z = center of mass coords of body
xu,yu,zu = unwrapped center of mass coords of body
ix,iy,iz = box image that the center of mass is in
vx,vy,vz = center of mass velocities
fx,fy,fz = force of center of mass
omegax,omegay,omegaz = angular velocity of body
angmomx,angmomy,angmomz = angular momentum of body
quatw,quati,quatj,quatk = quaternion components for body
tqx,tqy,tqz = torque on body
inertiax,inertiay,inertiaz = diagonalized moments of inertia of body
```

## 18.85.2 Examples

```
compute 1 all rigid/local myRigid mol x y z
```

---

## 18.85.3 Description

Define a computation that simply stores rigid body attributes for rigid bodies defined by the *fix rigid/small* command or one of its NVE, NVT, NPT, NPH variants. The data is stored as local data so it can be accessed by other *output commands* that process local data, such as the *compute reduce* or *dump local* commands.

Note that this command only works with the *fix rigid/small* command or its variants, not the *fix rigid* command and its variants. The ID of the *fix rigid/small* command used to define rigid bodies must be specified as *rigidID*. The *fix rigid* command is typically used to define a handful of (potentially very large) rigid bodies. It outputs similar per-body information as this command directly from the *fix* as global data; see the *fix rigid* page for details

The local data stored by this command is generated by looping over all the atoms owned on a processor. If the atom is not in the specified *group-ID* or is not part of a rigid body it is skipped. If it is not the atom within a body that is assigned to store the body information it is skipped (only one atom per body is so assigned). If it is the assigned atom, then the info for that body is output. This means that information for N bodies is generated. N may be less than the # of bodies defined by the *fix rigid* command, if the atoms in some bodies are not in the *group-ID*.

---

**Note:** Which atom in a body owns the body info is determined internal to LAMMPS; it's the one nearest the geometric center of the body. Typically you should avoid this complication, by defining the group associated with this *fix* to include/exclude entire bodies.

---

Note that as atoms and bodies migrate from processor to processor, there will be no consistent ordering of the entries within the local vector or array from one timestep to the next.

Here is an example of how to use this compute to dump rigid body info to a file:

```
compute 1 all rigid/local myRigid mol x y z fx fy fz
dump 1 all local 1000 tmp.dump index c_1[1] c_1[2] c_1[3] c_1[4] c_1[5] c_1[6] c_1[7]
```

---

This section explains the rigid body attributes that can be specified.

The *id* attribute is the atom-ID of the atom which owns the rigid body, which is assigned by the *fix rigid/small* command.

The *mol* attribute is the molecule ID of the rigid body. It should be the molecule ID which all of the atoms in the body belong to, since that is how the *fix rigid/small* command defines its rigid bodies.

The *mass* attribute is the total mass of the rigid body.

There are two options for outputting the coordinates of the center of mass (COM) of the body. The *x*, *y*, *z* attributes write the COM “unscaled”, in the appropriate distance *units* (Å, sigma, etc). Use *xu*, *yu*, *zu* if you want the COM “unwrapped” by the image flags for each body. Unwrapped means that if the body COM has passed through a periodic boundary one or more times, the value is generated what the COM coordinate would be if it had not been wrapped back into the periodic box.

The image flags for the body can be generated directly using the *ix*, *iy*, *iz* attributes. For periodic dimensions, they specify which image of the simulation box the COM is considered to be in. An image of 0 means it is inside the box as defined. A value of 2 means add 2 box lengths to get the true value. A value of -1 means subtract 1 box length to get the true value. LAMMPS updates these flags as the rigid body COMs cross periodic boundaries during the simulation.

The *vx*, *vy*, *vz*, *fx*, *fy*, *fz* attributes are components of the COM velocity and force on the COM of the body.

The *omegax*, *omegay*, and *omegaz* attributes are the angular velocity components of the body in the system frame around its COM.

The *angmomx*, *angmomy*, and *angmomz* attributes are the angular momentum components of the body in the system frame around its COM.

The *quatw*, *quati*, *quatj*, and *quatk* attributes are the components of the 4-vector quaternion representing the orientation of the rigid body. See the *set* command for an explanation of the quaternion vector.

The *tqx*, *tqy*, *tqz* attributes are components of the torque acting on the body around its COM.

The *inertiax*, *inertiay*, *inertiaz* attributes are components of diagonalized inertia tensor for the body, i.e the 3 moments of inertia for the body around its principal axes, as computed internally by LAMMPS.

---

## 18.85.4 Output info

This compute calculates a local vector or local array depending on the number of keywords. The length of the vector or number of rows in the array is the number of rigid bodies. If a single keyword is specified, a local vector is produced. If two or more keywords are specified, a local array is produced where the number of columns = the number of keywords. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector or array values will be in whatever *units* the corresponding attribute is in:

- id,mol = unitless
- mass = mass units
- x,y,z and xy,yu,zu = distance units
- vx,vy,vz = velocity units
- fx,fy,fz = force units
- omegax,omegay,omegaz = radians/time units
- angmomx,angmomy,angmomz = mass\*distance<sup>2</sup>/time units
- quatw,quati,quatj,quatk = unitless
- tqx,tqy,tqz = torque units
- inertiax,inertiay,inertiaz = mass\*distance<sup>2</sup> units

## 18.85.5 Restrictions

This compute is part of the RIGID package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 18.85.6 Related commands

*dump local, compute reduce*

## 18.85.7 Default

none

## 18.86 compute saed command

### 18.86.1 Syntax

```
compute ID group-ID saed lambda type1 type2 ... typeN keyword value ...
```

- ID, group-ID are documented in *compute* command
- saed = style name of this compute command
- lambda = wavelength of incident radiation (length units)
- type1 type2 ... typeN = chemical symbol of each atom type (see valid options below)
- zero or more keyword/value pairs may be appended
- keyword = *Kmax* or *Zone* or *dR\_Ewald* or *c* or *manual* or *echo*

*Kmax* value = Maximum distance explored from reciprocal space origin  
(inverse length units)

*Zone* values = z1 z2 z3

z1,z2,z3 = Zone axis of incident radiation. If z1=z2=z3=0 all  
reciprocal space will be meshed up to *Kmax*

*dR\_Ewald* value = Thickness of Ewald sphere slice intercepting  
reciprocal space (inverse length units)

*c* values = c1 c2 c3

c1,c2,c3 = parameters to adjust the spacing of the reciprocal  
lattice nodes in the h, k, and l directions respectively

*manual* = flag to use manual spacing of reciprocal lattice points  
based on the values of the *c* parameters

*echo* = flag to provide extra output for debugging purposes

### 18.86.2 Examples

```
compute 1 all saed 0.0251 Al 0 Kmax 1.70 Zone 0 0 1 dR_Ewald 0.01 c 0.5 0.5 0.5
compute 2 all saed 0.0251 Ni Kmax 1.70 Zone 0 0 0 c 0.05 0.05 0.05 manual echo

fix 1 all saed/vtk 1 1 1 c_1 file Al203_001.saed
fix 2 all saed/vtk 1 1 1 c_2 file Ni_000.saed
```



### 18.86.3 Description

Define a computation that calculates electron diffraction intensity as described in (*Coleman*) on a mesh of reciprocal lattice nodes defined by the entire simulation domain (or manually) using simulated radiation of wavelength  $\lambda$ .

The electron diffraction intensity  $I$  at each reciprocal lattice point is computed from the structure factor  $F$  using the equations:

$$I = \frac{F^* F}{N}$$

$$F(\mathbf{k}) = \sum_{j=1}^N f_j(\theta) \exp(2\pi i \mathbf{k} \cdot \mathbf{r}_j)$$

Here,  $\mathbf{K}$  is the location of the reciprocal lattice node,  $\mathbf{r}_j$  is the position of each atom,  $f_j$  are atomic scattering factors.

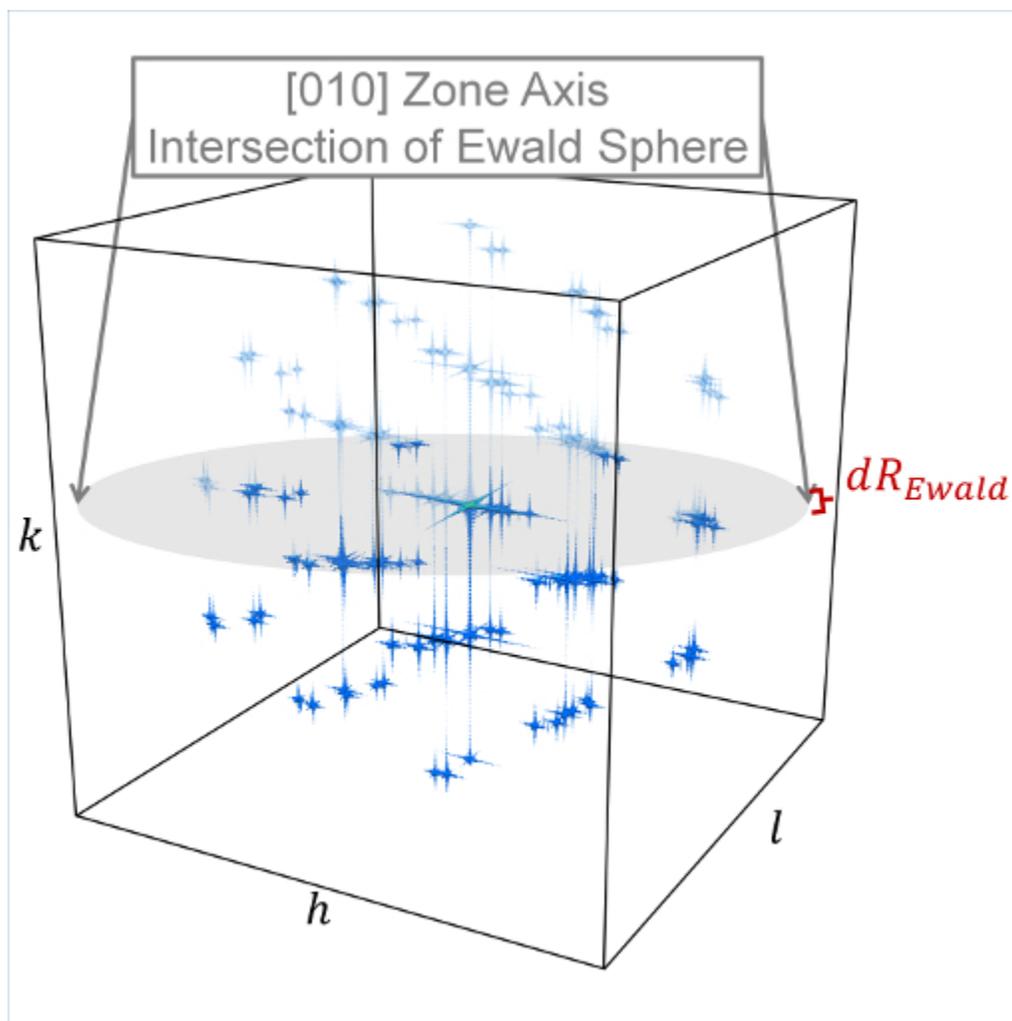
Diffraction intensities are calculated on a three-dimensional mesh of reciprocal lattice nodes. The mesh spacing is defined either (a) by the entire simulation domain or (b) manually using selected values as shown in the 2D diagram below.



For a mesh defined by the simulation domain, a rectilinear grid is constructed with spacing  $c \cdot \text{inv}(A)$  along each reciprocal lattice axis. Where  $A$  are the vectors corresponding to the edges of the simulation cell. If one or two directions has non-periodic boundary conditions, then the spacing in these directions is defined from the average of the (inversed) box lengths with periodic boundary conditions. Meshes defined by the simulation domain must contain at least one periodic boundary.

If the *manual* flag is included, the mesh of reciprocal lattice nodes will be defined using the  $c$  values for the spacing along each reciprocal lattice axis. Note that manual mapping of the reciprocal space mesh is good for comparing diffraction results from multiple simulations; however it can reduce the likelihood that Bragg reflections will be satisfied unless small spacing parameters ( $< 0.05 \text{ \AA}^{-1}$ ) are implemented. Meshes with manual spacing do not require a periodic boundary.

The limits of the reciprocal lattice mesh are determined by the use of the *Kmax*, *Zone*, and *dR\_Ewald* parameters. The rectilinear mesh created about the origin of reciprocal space is terminated at the boundary of a sphere of radius *Kmax* centered at the origin. If *Zone* parameters  $z1 = z2 = z3 = 0$  are used, diffraction intensities are computed throughout the entire spherical volume - note this can greatly increase the cost of computation. Otherwise, *Zone* parameters will denote the  $z1 = h$ ,  $z2 = k$ , and  $z3 = \ell$  (in a global sense) zone axis of an intersecting Ewald sphere. Diffraction intensities will only be computed at the intersection of the reciprocal lattice mesh and a *dR\_Ewald* thick surface of the Ewald sphere. See the example 3D intensity data and the intersection of a [010] zone axis in the below image.



The atomic scattering factors,  $f_j$ , accounts for the reduction in diffraction intensity due to Compton scattering. Compute saed uses analytical approximations of the atomic scattering factors that vary for each atom type (type1 type2 ... typeN) and angle of diffraction. The analytic approximation is computed using the formula (*Brown*):

$$f_j \left( \frac{\sin(\theta)}{\lambda} \right) = \sum_i^5 a_i \exp \left( -b_i \frac{\sin^2(\theta)}{\lambda^2} \right)$$

Coefficients parameterized by (*Fox*) are assigned for each atom type designating the chemical symbol and charge of each atom type. Valid chemical symbols for compute saed are:

H	He	Li	Be	B	C	N	O	F	Ne	Na	Mg	Al	Si	P	S	Cl	Ar	K	Ca
Sc	Ti	V	Cr	Mn	Fe	Co	Ni	Cu	Zn	Ga	Ge	As	Se	Br	Kr	Rb	Sr	Y	Zr
Nb	Mo	Tc	Ru	Rh	Pd	Ag	Cd	In	Sn	Sb	Te	I	Xe	Cs	Ba	La	Ce	Pr	Nd
Pm	Sm	Eu	Gd	Tb	Dy	Ho	Er	Tm	Yb	Lu	Hf	Ta	W	Re	Os	Ir	Pt	Au	Hg
Tl	Pb	Bi	Po	At	Rn	Fr	Ra	Ac	Th	Pa	U	Np	Pu	Am	Cm	Bk	Cf		

If the *echo* keyword is specified, compute saed will provide extra reporting information to the screen.

## 18.86.4 Output info

This compute calculates a global vector. The length of the vector is the number of reciprocal lattice nodes that are explored by the mesh. The entries of the global vector are the computed diffraction intensities as described above.

The vector can be accessed by any command that uses global values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

All array values calculated by this compute are “intensive”.

## 18.86.5 Restrictions

This compute is part of the DIFFRACTION package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The compute\_saed command does not work for triclinic cells.

## 18.86.6 Related commands

*fix saed\_vtk*, *compute xrd*

## 18.86.7 Default

The option defaults are Kmax = 1.70, Zone 1 0 0, c 1 1 1, dR\_Ewald = 0.01.

---

**(Coleman)** Coleman, Spearot, Capolungo, MSMSE, 21, 055020 (2013).

**(Brown)** Brown et al. International Tables for Crystallography Volume C: Mathematical and Chemical Tables, 554-95 (2004).

**(Fox)** Fox, O’Keefe, Tabbernor, Acta Crystallogr. A, 45, 786-93 (1989).

## 18.87 compute slice command

### 18.87.1 Syntax

```
compute ID group-ID slice Nstart Nstop Nskip input1 input2 ...
```

- ID, group-ID are documented in *compute* command
- slice = style name of this compute command
- Nstart = starting index within input vector(s)
- Nstop = stopping index within input vector(s)
- Nskip = extract every Nskip elements from input vector(s)
- input = c\_ID, c\_ID[N], f\_ID, f\_ID[N]

```
c_ID = global vector calculated by a compute with ID
c_ID[I] = Ith column of global array calculated by a compute with ID
f_ID = global vector calculated by a fix with ID
f_ID[I] = Ith column of global array calculated by a fix with ID
v_name = vector calculated by an vector-style variable with name
```

## 18.87.2 Examples

```
compute 1 all slice 1 100 10 c_msdmol[4]
compute 1 all slice 301 400 1 c_msdmol[4] v_myVec
```

## 18.87.3 Description

Define a calculation that “slices” one or more vector inputs into smaller vectors, one per listed input. The inputs can be global quantities; they cannot be per-atom or local quantities. *Computes* and *fixes* and vector-style *variables* can generate such global quantities. The group specified with this command is ignored.

The values extracted from the input vector(s) are determined by the *Nstart*, *Nstop*, and *Nskip* parameters. The elements of an input vector of length N are indexed from 1 to N. Starting at element *Nstart*, every Mth element is extracted, where  $M = Nskip$ , until element *Nstop* is reached. The extracted quantities are stored as a vector, which is typically shorter than the input vector.

Each listed input is operated on independently to produce one output vector. Each listed input must be a global vector or column of a global array calculated by another *compute* or *fix*.

If an input value begins with “c\_”, a compute ID must follow which has been previously defined in the input script and which generates a global vector or array. See the individual *compute* doc page for details. If no bracketed integer is appended, the vector calculated by the compute is used. If a bracketed integer is appended, the Ith column of the array calculated by the compute is used. Users can also write code for their own compute styles and *add them to LAMMPS*.

If a value begins with “f\_”, a fix ID must follow which has been previously defined in the input script and which generates a global vector or array. See the individual *fix* page for details. Note that some fixes only produce their values on certain timesteps, which must be compatible with when compute slice references the values, else an error results. If no bracketed integer is appended, the vector calculated by the fix is used. If a bracketed integer is appended, the Ith column of the array calculated by the fix is used. Users can also write code for their own fix style and *add them to LAMMPS*.

If an input value begins with “v\_”, a variable name must follow which has been previously defined in the input script. Only vector-style variables can be referenced. See the *variable* command for details. Note that variables of style *vector* define a formula which can reference individual atom properties or thermodynamic keywords, or they can invoke other computes, fixes, or variables when they are evaluated, so this is a very general means of specifying quantities to slice.

If a single input is specified this compute produces a global vector, even if the length of the vector is 1. If multiple inputs are specified, then a global array of values is produced, with the number of columns equal to the number of inputs specified.

## 18.87.4 Output info

This compute calculates a global vector if a single input value is specified or a global array with N columns where N is the number of inputs. The length of the vector or the number of rows in the array is equal to the number of values extracted from each input vector. These values can be used by any command that uses global vector or array values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector or array values calculated by this compute are simply copies of values generated by computes or fixes or variables that are input vectors to this compute. If there is a single input vector of intensive and/or extensive values, then each value in the vector of values calculated by this compute will be “intensive” or “extensive”, depending on the corresponding input value. If there are multiple input vectors, and all the values in them are intensive, then the array values calculated by this compute are “intensive”. If there are multiple input vectors, and any value in them is extensive, then the array values calculated by this compute are “extensive”. Values produced by a variable are treated as intensive.

The vector or array values will be in whatever *units* the input quantities are in.

## 18.87.5 Restrictions

none

## 18.87.6 Related commands

*compute*, *fix*, *compute reduce*

## 18.87.7 Default

none

# 18.88 compute smd/contact/radius command

## 18.88.1 Syntax

```
compute ID group-ID smd/contact/radius
```

- ID, group-ID are documented in *compute* command
- smd/contact/radius = style name of this compute command

## 18.88.2 Examples

```
compute 1 all smd/contact/radius
```

### 18.88.3 Description

Define a computation which outputs the contact radius, i.e., the radius used to prevent particles from penetrating each other. The contact radius is used only to prevent particles belonging to different physical bodies from penetrating each other. It is used by the contact pair styles, e.g., `smd/hertz` and `smd/tri_surface`.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

The value of the contact radius will be 0.0 for particles not in the specified compute group.

### 18.88.4 Output info

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle vector values will be in distance *units*.

### 18.88.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 18.88.6 Related commands

*dump custom* `smd/hertz` `smd/tri_surface`

### 18.88.7 Default

none

## 18.89 compute smd/damage command

### 18.89.1 Syntax

```
compute ID group-ID smd/damage
```

- ID, group-ID are documented in *compute* command
- smd/damage = style name of this compute command

## 18.89.2 Examples

```
compute 1 all smd/damage
```

## 18.89.3 Description

Define a computation that calculates the damage status of SPH particles according to the damage model which is defined via the SMD SPH pair styles, e.g., the maximum plastic strain failure criterion.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

### Output Info:

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle values are dimensionless and in the range of zero to one.

## 18.89.4 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the “Build

## 18.89.5 Related commands

*smd/plastic\_strain, smd/tlsph\_stress*

## 18.89.6 Default

none

# 18.90 compute smd/hourglass/error command

## 18.90.1 Syntax

```
compute ID group-ID smd/hourglass/error
```

- ID, group-ID are documented in [compute](#) command
- smd/hourglass/error = style name of this compute command

## 18.90.2 Examples

```
compute 1 all smd/hourglass/error
```

## 18.90.3 Description

Define a computation which outputs the error of the approximated relative separation with respect to the actual relative separation of the particles *i* and *j*. Ideally, if the deformation gradient is exact, and there exists a unique mapping between all particles' positions within the neighborhood of the central node and the deformation gradient, the approximated relative separation will coincide with the actual relative separation of the particles *i* and *j* in the deformed configuration. This compute is only really useful for debugging the hourglass control mechanism which is part of the Total-Lagrangian SPH pair style.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

### Output Info:

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle vector values will be dimensionless. See [units](#).

## 18.90.4 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This quantity will be computed only for particles which interact with `tlsph` pair style.

## 18.90.5 Related commands

*smd/tlsph\_defgrad*

## 18.90.6 Default

# 18.91 compute smd/internal/energy command

## 18.91.1 Syntax

```
compute ID group-ID smd/internal/energy
```

- ID, group-ID are documented in [compute](#) command
- smd/smd/internal/energy = style name of this compute command



## 18.91.2 Examples

```
compute 1 all smd/internal/energy
```

## 18.91.3 Description

Define a computation which outputs the per-particle enthalpy, i.e., the sum of potential energy and heat.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

## 18.91.4 Output Info

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle vector values will be given in *units* of energy.

## 18.91.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. This compute can only be used for particles which interact via the updated Lagrangian or total Lagrangian SPH pair styles.

## 18.91.6 Related commands

none

## 18.91.7 Default

# 18.92 compute smd/plastic/strain command

## 18.92.1 Syntax

```
compute ID group-ID smd/plastic/strain
```

- ID, group-ID are documented in [compute](#) command
- smd/plastic/strain = style name of this compute command

## 18.92.2 Examples

```
compute 1 all smd/plastic/strain
```

## 18.92.3 Description

Define a computation that outputs the equivalent plastic strain per particle. This command is only meaningful if a material model with plasticity is defined.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

### Output Info:

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle values will be given dimensionless. See [units](#).

## 18.92.4 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. This compute can only be used for particles which interact via the updated Lagrangian or total Lagrangian SPH pair styles.

## 18.92.5 Related commands

*smd/plastic/strain/rate*, *smd/tlsph/strain/rate*, *smd/tlsph/strain*

## 18.92.6 Default

none

# 18.93 compute smd/plastic/strain/rate command

## 18.93.1 Syntax

```
compute ID group-ID smd/plastic/strain/rate
```

- ID, group-ID are documented in [compute](#) command
- smd/plastic/strain/rate = style name of this compute command

## 18.93.2 Examples

```
compute 1 all smd/plastic/strain/rate
```

## 18.93.3 Description

Define a computation that outputs the time rate of the equivalent plastic strain. This command is only meaningful if a material model with plasticity is defined.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

### Output Info:

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle values will be given in *units* of one over time.

## 18.93.4 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. This compute can only be used for particles which interact via the updated Lagrangian or total Lagrangian SPH pair styles.

## 18.93.5 Related commands

*smd/plastic/strain*, *smd/tlsph/strain/rate*, *smd/tlsph/strain*

## 18.93.6 Default

none

# 18.94 compute smd/rho command

## 18.94.1 Syntax

```
compute ID group-ID smd/rho
```

- ID, group-ID are documented in [compute](#) command
- smd/rho = style name of this compute command

## 18.94.2 Examples

```
compute 1 all smd/rho
```

## 18.94.3 Description

Define a computation that calculates the per-particle mass density. The mass density is the mass of a particle which is constant during the course of a simulation, divided by its volume, which can change due to mechanical deformation.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

## 18.94.4 Output info

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle values will be in *units* of mass over volume.

## 18.94.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 18.94.6 Related commands

*compute smd/vol*

## 18.94.7 Default

none

# 18.95 compute smd/tlsph/defgrad command

## 18.95.1 Syntax

```
compute ID group-ID smd/tlsph/defgrad
```

- ID, group-ID are documented in [compute](#) command
- smd/tlsph/defgrad = style name of this compute command

## 18.95.2 Examples

```
compute 1 all smd/tlsph/defgrad
```

## 18.95.3 Description

Define a computation that calculates the deformation gradient. It is only meaningful for particles which interact according to the Total-Lagrangian SPH pair style.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

## 18.95.4 Output info

This compute outputs a per-particle vector of vectors (tensors), which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options.

The per-particle vector values will be given dimensionless. See [units](#). The per-particle vector has 10 entries. The first nine entries correspond to the *xx*, *xy*, *xz*, *yx*, *yy*, *yz*, *zx*, *zy*, *zz* components of the asymmetric deformation gradient tensor. The tenth entry is the determinant of the deformation gradient.

## 18.95.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. TThis compute can only be used for particles which interact via the total Lagrangian SPH pair style.

## 18.95.6 Related commands

*smd/hourglass/error*

## 18.95.7 Default

none

## 18.96 compute smd/tlsph/dt command

### 18.96.1 Syntax

```
compute ID group-ID smd/tlsph/dt
```

- ID, group-ID are documented in [compute](#) command
- smd/tlsph/dt = style name of this compute command

## 18.96.2 Examples

```
compute 1 all smd/tlsph/dt
```

## 18.96.3 Description

Define a computation that outputs the CFL-stable time increment per particle. This time increment is essentially given by the speed of sound, divided by the SPH smoothing length. Because both the speed of sound and the smoothing length typically change during the course of a simulation, the stable time increment needs to be re-computed every time step. This calculation is performed automatically in the relevant SPH pair styles and this compute only serves to make the stable time increment accessible for output purposes.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.96.4 Output info

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle values will be given in *units* of time.

## 18.96.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This compute can only be used for particles interacting with the Total-Lagrangian SPH pair style.

## 18.96.6 Related commands

*smd/adjust/dt*

## 18.96.7 Default

none

# 18.97 compute smd/tlsph/num/neighs command

## 18.97.1 Syntax

```
compute ID group-ID smd/tlsph/num/neighs
```

- ID, group-ID are documented in [compute](#) command
- smd/tlsph/num/neighs = style name of this compute command

## 18.97.2 Examples

```
compute 1 all smd/tlsph/num/neighs
```

## 18.97.3 Description

Define a computation that calculates the number of particles inside of the smoothing kernel radius for particles interacting via the Total-Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.97.4 Output info

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle values are dimensionless. See [units](#).

## 18.97.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This quantity will be computed only for particles which interact with the Total-Lagrangian pair style.

## 18.97.6 Related commands

*smd/ulsph/num/neighs*

## 18.97.7 Default

none

# 18.98 compute smd/tlsph/shape command

## 18.98.1 Syntax

```
compute ID group-ID smd/tlsph/shape
```

- ID, group-ID are documented in [compute](#) command
- smd/tlsph/shape = style name of this compute command

## 18.98.2 Examples

```
compute 1 all smd/tlsph/shape
```

## 18.98.3 Description

Define a computation that outputs the current shape of the volume associated with a particle as a rotated ellipsoid. It is only meaningful for particles which interact according to the Total-Lagrangian SPH pair style.

See [this PDF guide](#) to use Smooth Mach Dynamics in LAMMPS.

## 18.98.4 Output info

This compute calculates a per-particle vector of vectors, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle vector has 7 entries. The first three entries correspond to the lengths of the ellipsoid's axes and have units of length. These axis values are computed as the contact radius times the xx, yy, or zz components of the Green-Lagrange strain tensor associated with the particle. The next 4 values are quaternions (order: q, x, y, z) which describe the spatial rotation of the particle relative to its initial state.

## 18.98.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This quantity will be computed only for particles which interact with the Total-Lagrangian SPH pair style.

## 18.98.6 Related commands

*smd/contact/radius*

## 18.98.7 Default

none

# 18.99 compute smd/tlsph/strain command

## 18.99.1 Syntax

```
compute ID group-ID smd/tlsph/strain
```

- ID, group-ID are documented in [compute](#) command
- smd/tlsph/strain = style name of this compute command



## 18.99.2 Examples

```
compute 1 all smd/tlsph/strain
```

## 18.99.3 Description

Define a computation that calculates the Green-Lagrange strain tensor for particles interacting via the Total-Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.99.4 Output info

This compute calculates a per-particle vector of vectors (tensors), which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options.

The per-particle tensor values will be given dimensionless. See [units](#).

The per-particle vector has 6 entries, corresponding to the xx, yy, zz, xy, xz, yz components of the symmetric strain tensor.

## 18.99.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This quantity will be computed only for particles which interact with the Total-Lagrangian SPH pair style.

## 18.99.6 Related commands

*smd/tlsph/strain/rate*, *smd/tlsph/stress*

## 18.99.7 Default

none

## 18.100 compute smd/tlsph/strain/rate command

### 18.100.1 Syntax

```
compute ID group-ID smd/tlsph/strain/rate
```

- ID, group-ID are documented in [compute](#) command
- smd/tlsph/strain/rate = style name of this compute command

## 18.100.2 Examples

```
compute 1 all smd/tlsph/strain/rate
```

## 18.100.3 Description

Define a computation that calculates the rate of the strain tensor for particles interacting via the Total-Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.100.4 Output info

This compute calculates a per-particle vector of vectors (tensors), which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options.

The values will be given in *units* of one over time.

The per-particle vector has 6 entries, corresponding to the xx, yy, zz, xy, xz, yz components of the symmetric strain rate tensor.

## 18.100.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This quantity will be computed only for particles which interact with Total-Lagrangian SPH pair style.

## 18.100.6 Related commands

*compute smd/tlsph/strain*, *compute smd/tlsph/stress*

## 18.100.7 Default

none

# 18.101 compute smd/tlsph/stress command

## 18.101.1 Syntax

```
compute ID group-ID smd/tlsph/stress
```

- ID, group-ID are documented in [compute](#) command
- smd/tlsph/stress = style name of this compute command

## 18.101.2 Examples

```
compute 1 all smd/tlsph/stress
```

## 18.101.3 Description

Define a computation that outputs the Cauchy stress tensor for particles interacting via the Total-Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.101.4 Output info

This compute calculates a per-particle vector of vectors (tensors), which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options.

The values will be given in *units* of pressure.

The per-particle vector has 7 entries. The first six entries correspond to the xx, yy, zz, xy, xz and yz components of the symmetric Cauchy stress tensor. The seventh entry is the second invariant of the stress tensor, i.e., the von Mises equivalent stress.

## 18.101.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This quantity will be computed only for particles which interact with the Total-Lagrangian SPH pair style.

## 18.101.6 Related commands

*compute smd/tlsph/strain*, *compute smd/tlsph/strain/rate*

## 18.101.7 Default

none

# 18.102 compute smd/triangle/vertices command

## 18.102.1 Syntax

```
compute ID group-ID smd/triangle/vertices
```

- ID, group-ID are documented in [compute](#) command
- smd/triangle/vertices = style name of this compute command

## 18.102.2 Examples

```
compute 1 all smd/triangle/vertices
```

## 18.102.3 Description

Define a computation that returns the coordinates of the vertices corresponding to the triangle-elements of a mesh created by the *fix smd/wall\_surface*.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.102.4 Output info

This compute returns a per-particle vector of vectors, which can be accessed by any command that uses per-particle values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The per-particle vector has nine entries,  $(x1/y1/z1)$ ,  $(x2/y2/z2)$ , and  $(x3/y3/z3)$  corresponding to the first, second, and third vertex of each triangle.

It is only meaningful to use this compute for a group of particles which is created via the *fix smd/wall\_surface* command.

The output of this compute can be used with the `dump2vtk_tris` tool to generate a VTK representation of the `smd/wall_surface` mesh for visualization purposes.

The values will be given in *units* of distance.

## 18.102.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 18.102.6 Related commands

*fix smd/move/tri/surf*, *fix smd/wall\_surface*

## 18.102.7 Default

none

## 18.103 compute smd/ulsph/effm command

### 18.103.1 Syntax

```
compute ID group-ID smd/ulsph/effm
```

- ID, group-ID are documented in *compute* command
- smd/ulsph/effm = style name of this compute command

## 18.103.2 Examples

```
compute 1 all smd/ulsph/effm
```

## 18.103.3 Description

Define a computation that outputs the effective shear modulus for particles interacting via the updated Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.103.4 Output info

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle vector contains the current effective per atom shear modulus as computed by the *pair smd/ulsph* pair style.

## 18.103.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. This compute can only be used for particles which interact with the updated Lagrangian SPH pair style.

## 18.103.6 Related commands

*pair smd/ulsph*

## 18.103.7 Default

none

# 18.104 compute smd/ulsph/num/neighs command

## 18.104.1 Syntax

```
compute ID group-ID smd/ulsph/num/neighs
```

- ID, group-ID are documented in [compute](#) command
- smd/ulsph/num/neighs = style name of this compute command

## 18.104.2 Examples

```
compute 1 all smd/ulsph/num/neighs
```

## 18.104.3 Description

Define a computation that returns the number of neighbor particles inside of the smoothing kernel radius for particles interacting via the updated Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.104.4 Output info

This compute returns a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle values will be given dimensionless, see [units](#).

## 18.104.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. This compute can only be used for particles which interact with the updated Lagrangian SPH pair style.

## 18.104.6 Related commands

*compute smd/tlsph/num/neighs*

## 18.104.7 Default

none

# 18.105 compute smd/ulsph/strain command

## 18.105.1 Syntax

```
compute ID group-ID smd/ulsph/strain
```

- ID, group-ID are documented in [compute](#) command
- smd/ulsph/strain = style name of this compute command

## 18.105.2 Examples

```
compute 1 all smd/ulsph/strain
```

## 18.105.3 Description

Define a computation that outputs the logarithmic strain tensor. for particles interacting via the updated Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.105.4 Output info

This compute calculates a per-particle tensor, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle vector has 6 entries, corresponding to the xx, yy, zz, xy, xz, yz components of the symmetric strain rate tensor.

The per-particle tensor values will be given dimensionless, see [units](#).

## 18.105.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. This compute can only be used for particles which interact with the updated Lagrangian SPH pair style.

## 18.105.6 Related commands

*compute smd/tlsph/strain*

## 18.105.7 Default

none

# 18.106 compute smd/ulsph/strain/rate command

## 18.106.1 Syntax

```
compute ID group-ID smd/ulsph/strain/rate
```

- ID, group-ID are documented in [compute](#) command
- smd/ulsph/strain/rate = style name of this compute command

## 18.106.2 Examples

```
compute 1 all smd/ulsph/strain/rate
```

## 18.106.3 Description

Define a computation that outputs the rate of the logarithmic strain tensor for particles interacting via the updated Lagrangian SPH pair style.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.106.4 Output info

This compute calculates a per-particle vector of vectors (tensors), which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options.

The values will be given in *units* of one over time.

The per-particle vector has 6 entries, corresponding to the xx, yy, zz, xy, xz, yz components of the symmetric strain rate tensor.

## 18.106.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This compute can only be used for particles which interact with the updated Lagrangian SPH pair style.

## 18.106.6 Related commands

*compute smd/tlsph/strain/rate*

## 18.106.7 Default

none

## 18.107 compute smd/ulsph/stress command

### 18.107.1 Syntax

```
compute ID group-ID smd/ulsph/stress
```

- ID, group-ID are documented in [compute](#) command
- smd/ulsph/stress = style name of this compute command



## 18.107.2 Examples

```
compute 1 all smd/ulsph/stress
```

## 18.107.3 Description

Define a computation that outputs the Cauchy stress tensor.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.107.4 Output info

This compute calculates a per-particle vector of vectors (tensors), which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) doc page for an overview of LAMMPS output options.

The values will be given in *units* of pressure.

The per-particle vector has 7 entries. The first six entries correspond to the xx, yy, zz, xy, xz, yz components of the symmetric Cauchy stress tensor. The seventh entry is the second invariant of the stress tensor, i.e., the von Mises equivalent stress.

## 18.107.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info. This compute can only be used for particles which interact with the updated Lagrangian SPH pair style.

## 18.107.6 Related commands

*compute smd/ulsph/strain, compute smd/ulsph/strain/rate compute smd/tlspH/stress*

## 18.107.7 Default

none

## 18.108 compute smd/vol command

### 18.108.1 Syntax

```
compute ID group-ID smd/vol
```

- ID, group-ID are documented in [compute](#) command
- smd/vol = style name of this compute command

## 18.108.2 Examples

```
compute 1 all smd/vol
```

## 18.108.3 Description

Define a computation that provides the per-particle volume and the sum of the per-particle volumes of the group for which the fix is defined.

See [this PDF guide](#) to using Smooth Mach Dynamics in LAMMPS.

## 18.108.4 Output info

This compute calculates a per-particle vector, which can be accessed by any command that uses per-particle values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-particle vector values will be given in *units* of volume.

Additionally, the compute returns a scalar, which is the sum of the per-particle volumes of the group for which the fix is defined.

## 18.108.5 Restrictions

This compute is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 18.108.6 Related commands

*compute smd/rho*

## 18.108.7 Default

none

## 18.109 compute sna/atom command

## 18.110 compute snad/atom command

## 18.111 compute snav/atom command

## 18.112 compute snap command

## 18.113 compute sna/grid command

## 18.114 compute sna/grid/local command

### 18.114.1 Syntax

```

compute ID group-ID sna/atom rcutfac rfac0 twojmax R_1 R_2 ... w_1 w_2 ... keyword_
↪values ...
compute ID group-ID snad/atom rcutfac rfac0 twojmax R_1 R_2 ... w_1 w_2 ... keyword_
↪values ...
compute ID group-ID snav/atom rcutfac rfac0 twojmax R_1 R_2 ... w_1 w_2 ... keyword_
↪values ...
compute ID group-ID snap rcutfac rfac0 twojmax R_1 R_2 ... w_1 w_2 ... keyword values ...
compute ID group-ID snap rcutfac rfac0 twojmax R_1 R_2 ... w_1 w_2 ... keyword values ...
compute ID group-ID sna/grid nx ny nz rcutfac rfac0 twojmax R_1 R_2 ... w_1 w_2 ...
↪keyword values ...
compute ID group-ID sna/grid/local nx ny nz rcutfac rfac0 twojmax R_1 R_2 ... w_1 w_2 ...
↪ keyword values ...

```

- ID, group-ID are documented in *compute* command
- sna/atom = style name of this compute command
- rcutfac = scale factor applied to all cutoff radii (positive real)
- rfac0 = parameter in distance to angle conversion ( $0 < \text{rcutfac} < 1$ )
- twojmax = band limit for bispectrum components (non-negative integer)
- R\_1, R\_2,... = list of cutoff radii, one for each type (distance units)
- w\_1, w\_2,... = list of neighbor weights, one for each type
- nx, ny, nz = number of grid points in x, y, and z directions (positive integer)
- zero or more keyword/value pairs may be appended
- keyword = *rmin0* or *switchflag* or *bzeroflag* or *quadraticflag* or *chem* or *bnormflag* or *wselfallflag* or *bikflag* or *switchinnerflag* or *sinner* or *dinner* or *dgradflag*
  - rmin0* value = parameter in distance to angle conversion (distance units)
  - switchflag* value = 0 or 1
    - 0 = do not use switching function
    - 1 = use switching function
  - bzeroflag* value = 0 or 1
    - 0 = do not subtract B0

```

    1 = subtract B0
quadraticflag value = 0 or 1
    0 = do not generate quadratic terms
    1 = generate quadratic terms
chem values = nelements elementlist
    nelements = number of SNAP elements
    elementlist = ntypes integers in range [0, nelements)
bnormflag value = 0 or 1
    0 = do not normalize
    1 = normalize bispectrum components
wselfallflag value = 0 or 1
    0 = self-contribution only for element of central atom
    1 = self-contribution for all elements
switchinnerflag value = 0 or 1
    0 = do not use inner switching function
    1 = use inner switching function
sinner values = sinnerlist
    sinnerlist = ntypes values of Sinner (distance units)
dinner values = dinnerlist
    dinnerlist = ntypes values of Dinner (distance units)
bikflag value = 0 or 1 (only implemented for compute snap)
    0 = descriptors are summed over atoms of each type
    1 = descriptors are listed separately for each atom
dgradflag value = 0 or 1 (only implemented for compute snap)
    0 = descriptor gradients are summed over atoms of each type
    1 = descriptor gradients are listed separately for each atom pair

```

## 18.114.2 Examples

```

compute b all sna/atom 1.4 0.99363 6 2.0 2.4 0.75 1.0 rmin0 0.0
compute db all sna/atom 1.4 0.95 6 2.0 1.0
compute vb all sna/atom 1.4 0.95 6 2.0 1.0
compute snap all snap 1.4 0.95 6 2.0 1.0
compute snap all snap 1.0 0.99363 6 3.81 3.83 1.0 0.93 chem 2 0 1
compute snap all snap 1.0 0.99363 6 3.81 3.83 1.0 0.93 switchinnerflag 1 sinner 1.35 1.6_
↪dinner 0.25 0.3
compute bgrid all sna/grid/local 200 200 200 1.4 0.95 6 2.0 1.0

```

## 18.114.3 Description

Define a computation that calculates a set of quantities related to the bispectrum components of the atoms in a group. These computes are used primarily for calculating the dependence of energy, force, and stress components on the linear coefficients in the *snap pair\_style*, which is useful when training a SNAP potential to match target data.

Bispectrum components of an atom are order parameters characterizing the radial and angular distribution of neighbor atoms. The detailed mathematical definition is given in the paper by Thompson et al. (*Thompson*)

The position of a neighbor atom  $i'$  relative to a central atom  $i$  is a point within the 3D ball of radius  $R_{ii'} = rcutfac (R_i + R_{i'})$

Bartok et al. (*Bartok*), proposed mapping this 3D ball onto the 3-sphere, the surface of the unit ball in a four-dimensional

space. The radial distance  $r$  within  $R_{ii'}$  is mapped on to a third polar angle  $\theta_0$  defined by,

$$\theta_0 = \text{rfac0} \frac{r - r_{\text{min0}}}{R_{ii'} - r_{\text{min0}}} \pi$$

In this way, all possible neighbor positions are mapped on to a subset of the 3-sphere. Points south of the latitude  $\theta_0 = \text{rfac0} \pi$  are excluded.

The natural basis for functions on the 3-sphere is formed by the representatives of  $SU(2)$ , the matrices  $U_{m,m'}^j(\theta, \phi, \theta_0)$ . These functions are better known as  $D_{m,m'}^j$ , the elements of the Wigner  $D$ -matrices (*Meremianin, Varshalovich, Mason*) The density of neighbors on the 3-sphere can be written as a sum of Dirac-delta functions, one for each neighbor, weighted by species and radial distance. Expanding this density function as a generalized Fourier series in the basis functions, we can write each Fourier coefficient as

$$w_{m,m'}^j = U_{m,m'}^j(0, 0, 0) + \sum_{r_{ii'} < R_{ii'}} f_c(r_{ii'}) w_{\mu_{i'}} U_{m,m'}^j(\theta_0, \theta, \phi)$$

The  $w_{\mu_{i'}}$  neighbor weights are dimensionless numbers that depend on  $\mu_{i'}$ , the SNAP element of atom  $i'$ , while the central atom is arbitrarily assigned a unit weight. The function  $f_c(r)$  ensures that the contribution of each neighbor atom goes smoothly to zero at  $R_{ii'}$ :

$$f_c(r) = \frac{1}{2} \left( \cos\left(\pi \frac{r - r_{\text{min0}}}{R_{ii'} - r_{\text{min0}}}\right) + 1 \right), r \leq R_{ii'}$$

$$= 0, r > R_{ii'}$$

The expansion coefficients  $w_{m,m'}^j$  are complex-valued and they are not directly useful as descriptors, because they are not invariant under rotation of the polar coordinate frame. However, the following scalar triple products of expansion coefficients can be shown to be real-valued and invariant under rotation (*Bartok*).

$$B_{j_1, j_2, j} = \sum_{m_1, m'_1 = -j_1}^{j_1} \sum_{m_2, m'_2 = -j_2}^{j_2} \sum_{m, m' = -j}^j (w_{m, m'}^j)^* H_{\begin{matrix} j_1 m_1 m'_1 \\ j_2 m_2 m'_2 \\ j m m' \end{matrix}} w_{m_1, m'_1}^{j_1} w_{m_2, m'_2}^{j_2}$$

The constants  $H_{j_1 m_1 m'_1, j_2 m_2 m'_2}^{j m m'}$  are coupling coefficients, analogous to Clebsch-Gordan coefficients for rotations on the 2-sphere. These invariants are the components of the bispectrum and these are the quantities calculated by the compute *sna/atom*. They characterize the strength of density correlations at three points on the 3-sphere. The  $j_2=0$  subset form the power spectrum, which characterizes the correlations of two points. The lowest-order components describe the coarsest features of the density function, while higher-order components reflect finer detail. Each bispectrum component contains terms that depend on the positions of up to 4 atoms (3 neighbors and the central atom).

Compute *sna/datom* calculates the derivative of the bispectrum components summed separately for each LAMMPS atom type:

$$- \sum_{i' \in I} \frac{\partial B_{j_1, j_2, j}^{i'}}{\partial \mathbf{r}_i}$$

The sum is over all atoms  $i'$  of atom type  $I$ . For each atom  $i$ , this compute evaluates the above expression for each direction, each atom type, and each bispectrum component. See section below on output for a detailed explanation.

Compute *snav/atom* calculates the virial contribution due to the derivatives:

$$-\mathbf{r}_i \otimes \sum_{i' \in I} \frac{\partial B_{j_1, j_2, j}^{i'}}{\partial \mathbf{r}_i}$$

Again, the sum is over all atoms  $i'$  of atom type  $I$ . For each atom  $i$ , this compute evaluates the above expression for each of the six virial components, each atom type, and each bispectrum component. See section below on output for a detailed explanation.

Compute *snap* calculates a global array containing information related to all three of the above per-atom computes *sna/atom*, *snad/atom*, and *snav/atom*. The first row of the array contains the summation of *sna/atom* over all atoms, but broken out by type. The last six rows of the array contain the summation of *snav/atom* over all atoms, broken out by type. In between these are  $3*N$  rows containing the same values computed by *snad/atom* (these are already summed over all atoms and broken out by type). The element in the last column of each row contains the potential energy, force, or stress, according to the row. These quantities correspond to the user-specified reference potential that must be subtracted from the target data when fitting SNAP. The potential energy calculation uses the built in compute *thermo\_pe*. The stress calculation uses a compute called *snap\_press* that is automatically created behind the scenes, according to the following command:

```
compute snap_press all pressure NULL virial
```

See section below on output for a detailed explanation of the data layout in the global array.

The compute *sna/grid* and *sna/grid/local* commands calculate bispectrum components for a regular grid of points. These are calculated from the local density of nearby atoms  $i'$  around each grid point, as if there was a central atom  $i$  at the grid point. This is useful for characterizing fine-scale structure in a configuration of atoms, and it is used in the [MALA package](#) to build machine-learning surrogates for finite-temperature Kohn-Sham density functional theory (*Ellis et al.*) Neighbor atoms not in the group do not contribute to the bispectrum components of the grid points. The distance cutoff  $R_{ii'}$  assumes that  $i$  has the same type as the neighbor atom  $i'$ .

Compute *sna/grid* calculates a global array containing bispectrum components for a regular grid of points. The grid is aligned with the current box dimensions, with the first point at the box origin, and forming a regular 3d array with  $n_x$ ,  $n_y$ , and  $n_z$  points in the x, y, and z directions. For triclinic boxes, the array is congruent with the periodic lattice vectors a, b, and c. The array contains one row for each of the  $n_x \times n_y \times n_z$  grid points, looping over the index for  $i_x$  fastest, then  $i_y$ , and  $i_z$  slowest. Each row of the array contains the x, y, and z coordinates of the grid point, followed by the bispectrum components. See section below on output for a detailed explanation of the data layout in the global array.

Compute *sna/grid/local* calculates bispectrum components of a regular grid of points similarly to compute *sna/grid* described above. However, because the array is local, it contains only rows for grid points that are local to the processor sub-domain. The global grid of  $n_x \times n_y \times n_z$  points is still laid out in space the same as for *sna/grid*, but grid points are strictly partitioned, so that every grid point appears in one and only one local array. The array contains one row for each of the local grid points, looping over the global index  $i_x$  fastest, then  $i_y$ , and  $i_z$  slowest. Each row of the array contains the global indexes  $i_x$ ,  $i_y$ , and  $i_z$  first, followed by the x, y, and z coordinates of the grid point, followed by the bispectrum components. See section below on output for a detailed explanation of the data layout in the global array.

The value of all bispectrum components will be zero for atoms not in the group. Neighbor atoms not in the group do not contribute to the bispectrum of atoms in the group.

The neighbor list needed to compute this quantity is constructed each time the calculation is performed (i.e. each time a snapshot of atoms is dumped). Thus it can be inefficient to compute/dump this quantity too frequently.

The argument *rcutfac* is a scale factor that controls the ratio of atomic radius to radial cutoff distance.

The argument *rfac0* and the optional keyword *rmin0* define the linear mapping from radial distance to polar angle  $\theta_0$  on the 3-sphere, given above.

The argument *twojmax* defines which bispectrum components are generated. See section below on output for a detailed explanation of the number of bispectrum components and the ordered in which they are listed.

The keyword *switchflag* can be used to turn off the switching function  $f_c(r)$ .

The keyword *bzeroflag* determines whether or not  $B_0$ , the bispectrum components of an atom with no neighbors, are subtracted from the calculated bispectrum components. This optional keyword normally only affects compute *sna/atom*. However, when *quadraticflag* is on, it also affects *snad/atom* and *snav/atom*.

The keyword *quadraticflag* determines whether or not the quadratic combinations of bispectrum quantities are generated. These are formed by taking the outer product of the vector of bispectrum components with itself. See section below on output for a detailed explanation of the number of quadratic terms and the ordered in which they are listed.

The keyword *chem* activates the explicit multi-element variant of the SNAP bispectrum components. The argument *nelements* specifies the number of SNAP elements that will be handled. This is followed by *elementlist*, a list of integers of length *ntypes*, with values in the range  $[0, \text{nelements})$ , which maps each LAMMPS type to one of the SNAP elements. Note that multiple LAMMPS types can be mapped to the same element, and some elements may be mapped by no LAMMPS type. However, in typical use cases (training SNAP potentials) the mapping from LAMMPS types to elements is one-to-one.

The explicit multi-element variant invoked by the *chem* keyword partitions the density of neighbors into partial densities for each chemical element. This is described in detail in the paper by *Cusentino et al.* The bispectrum components are indexed on ordered triplets of elements:

$$B_{j_1, j_2, j}^{\kappa\lambda\mu} = \sum_{m_1, m'_1 = -j_1}^{j_1} \sum_{m_2, m'_2 = -j_2}^{j_2} \sum_{m, m' = -j}^j (u_{j, m, m'}^\mu)^* H_{\substack{j_1 m_1 m'_1 \\ j_2 m_2 m'_2}}^{j m m'} u_{j_1, m_1, m'_1}^\kappa u_{j_2, m_2, m'_2}^\lambda$$

where  $u_{j, m, m'}^\mu$  is an expansion coefficient for the partial density of neighbors of element  $\mu$

$$u_{j, m, m'}^\mu = w_{\mu_i \mu}^{self} U^{j, m, m'}(0, 0, 0) + \sum_{r_{ii'} < R_{ii'}} \delta_{\mu \mu_i'} f_c(r_{ii'}) w_{\mu_i'} U^{j, m, m'}(\theta_0, \theta, \phi)$$

where  $w_{\mu_i \mu}^{self}$  is the self-contribution, which is either 1 or 0 (see keyword *wselallflag* below),  $\delta_{\mu \mu_i'}$  indicates that the sum is only over neighbor atoms of element  $\mu$ , and all other quantities are the same as those appearing in the original equation for  $u_{m, m'}^j$  given above.

The keyword *wselallflag* defines the rule used for the self-contribution. If *wselallflag* is on, then  $w_{\mu_i \mu}^{self} = 1$ . If it is off then  $w_{\mu_i \mu}^{self} = 0$ , except in the case of  $\mu_i = \mu$ , when  $w_{\mu_i \mu}^{self} = 1$ . When the *chem* keyword is not used, this keyword has no effect.

The keyword *bnormflag* determines whether or not the bispectrum component  $B_{j_1, j_2, j}$  is divided by a factor of  $2j + 1$ . This normalization simplifies force calculations because of the following symmetry relation

$$\frac{B_{j_1, j_2, j}}{2j + 1} = \frac{B_{j, j_2, j_1}}{2j_1 + 1} = \frac{B_{j_1, j, j_2}}{2j_2 + 1}$$

This option is typically used in conjunction with the *chem* keyword, and LAMMPS will generate a warning if both *chem* and *bnormflag* are not both set or not both unset.

The keyword *switchinnerflag* with value 1 activates an additional radial switching function similar to  $f_c(r)$  above, but acting to switch off smoothly contributions from neighbor atoms at short separation distances. This is useful when SNAP is used in combination with a simple repulsive potential. For a neighbor atom at distance  $r$ , its contribution is scaled by a multiplicative factor  $f_{inner}(r)$  defined as follows:

$$\begin{aligned} &= 0, r \leq S_{inner} - D_{inner} \\ f_{inner}(r) &= \frac{1}{2} \left( 1 - \cos\left(\frac{\pi}{2} \left( 1 + \frac{r - S_{inner}}{D_{inner}} \right) \right) \right), S_{inner} - D_{inner} < r \leq S_{inner} + D_{inner} \\ &= 1, r > S_{inner} + D_{inner} \end{aligned}$$

where the switching region is centered at  $S_{inner}$  and it extends a distance  $D_{inner}$  to the left and to the right of this. With this option, additional keywords *sinner* and *dinner* must be used, each followed by *ntypes* values for  $S_{inner}$  and  $D_{inner}$ , respectively. When the central atom and the neighbor atom have different types, the values of  $S_{inner}$  and  $D_{inner}$  are the arithmetic means of the values for both types.

The keywords *bikflag* and *dgradflag* are only used by compute *snap*. The keyword *bikflag* determines whether or not to list the descriptors of each atom separately, or sum them together and list in a single row. If *bikflag* is set to 0 then a single bispectrum row is used, which contains the per-atom bispectrum descriptors  $B_{i, k}$  summed over all atoms  $i$  to produce  $B_k$ . If *bikflag* is set to 1 this is replaced by a separate per-atom bispectrum row for each atom. In this case, the entries in the final column for these rows are set to zero.

The keyword *dgradflag* determines whether to sum atom gradients or list them separately. If *dgradflag* is set to 0, the bispectrum descriptor gradients w.r.t. atom *j* are summed over all atoms *i* of type *I* (similar to *snad/atom* above). If *dgradflag* is set to 1, gradients are listed separately for each pair of atoms. Each row corresponds to a single term  $\frac{\partial B_{i,k}}{\partial r_j^a}$  where  $r_j^a$  is the *a*-th position coordinate of the atom with global index *j*. This also changes the number of columns to be equal to the number of bispectrum components, with 3 additional columns representing the indices *i*, *j*, and *a*, as explained more in the Output info section below. The option *dgradflag=1* requires that *bikflag=1*.

---

**Note:** Using *dgradflag* = 1 produces a global array with  $N + 3N^2 + 1$  rows which becomes expensive for systems with more than 1000 atoms.

---

---

**Note:** If you have a bonded system, then the settings of *special\_bonds* command can remove pairwise interactions between atoms in the same bond, angle, or dihedral. This is the default setting for the *special\_bonds* command, and means those pairwise interactions do not appear in the neighbor list. Because this fix uses the neighbor list, it also means those pairs will not be included in the calculation. One way to get around this, is to write a dump file, and use the *rerun* command to compute the bispectrum components for snapshots in the dump file. The rerun script can use a *special\_bonds* command that includes all pairs in the neighbor list.

---

#### 18.114.4 Output info

Compute *snad/atom* calculates a per-atom array, each column corresponding to a particular bispectrum component. The total number of columns and the identity of the bispectrum component contained in each column depend of the value of *twojmax*, as described by the following piece of python code:

```
for j1 in range(0,twojmax+1):
    for j2 in range(0,j1+1):
        for j in range(j1-j2,min(twojmax,j1+j2)+1,2):
            if (j>=j1): print j1/2.,j2/2.,j/2.
```

For even  $twojmax = 2(m-1)$ ,  $K = m(m+1)(2m+1)/6$ , the *m*-th pyramidal number. For odd  $twojmax = 2m-1$ ,  $K = m(m+1)(m+2)/3$ , twice the *m*-th tetrahedral number.

---

**Note:** the *diagonal* keyword allowing other possible choices for the number of bispectrum components was removed in 2019, since all potentials use the value of 3, corresponding to the above set of bispectrum components.

---

Compute *snad/atom* evaluates a per-atom array. The columns are arranged into *ntypes* blocks, listed in order of atom type *I*. Each block contains three sub-blocks corresponding to the *x*, *y*, and *z* components of the atom position. Each of these sub-blocks contains *K* columns for the *K* bispectrum components, the same as for compute *snad/atom*

Compute *snav/atom* evaluates a per-atom array. The columns are arranged into *ntypes* blocks, listed in order of atom type *I*. Each block contains six sub-blocks corresponding to the *xx*, *yy*, *zz*, *yz*, *xz*, and *xy* components of the virial tensor in Voigt notation. Each of these sub-blocks contains *K* columns for the *K* bispectrum components, the same as for compute *snad/atom*

Compute *snap* evaluates a global array. The columns are arranged into *ntypes* blocks, listed in order of atom type *I*. Each block contains one column for each bispectrum component, the same as for compute *snad/atom*. A final column contains the corresponding energy, force component on an atom, or virial stress component. The rows of the array appear in the following order:

- 1 row: *snad/atom* quantities summed for all atoms of type *I*



- $3*N$  rows: *snad/atom* quantities, with derivatives w.r.t.  $x$ ,  $y$ , and  $z$  coordinate of atom  $i$  appearing in consecutive rows. The atoms are sorted based on atom ID.
- 6 rows: *snav/atom* quantities summed for all atoms of type  $I$

For example, if  $K=30$  and  $ntypes=1$ , the number of columns in the per-atom arrays generated by *sna/atom*, *snad/atom*, and *snav/atom* are 30, 90, and 180, respectively. With *quadratic* value=1, the numbers of columns are 930, 2790, and 5580, respectively. The number of columns in the global array generated by *snap* are 31, and 931, respectively, while the number of rows is  $1+3*N+6$ , where  $N$  is the total number of atoms.

Compute *sna/grid* evaluates a global array. The array contains one row for each of the  $n_x \times n_y \times n_z$  grid points, looping over the index for  $i_x$  fastest, then  $i_y$ , and  $i_z$  slowest. Each row of the array contains the  $x$ ,  $y$ , and  $z$  coordinates of the grid point, followed by the bispectrum components.

Compute *sna/grid/local* evaluates a local array. The array contains one row for each of the local grid points, looping over the global index  $i_x$  fastest, then  $i_y$ , and  $i_z$  slowest. Each row of the array contains the global indexes  $i_x$ ,  $i_y$ , and  $i_z$  first, followed by the  $x$ ,  $y$ , and  $z$  coordinates of the grid point, followed by the bispectrum components.

If the *quadratic* keyword value is set to 1, then additional columns are generated, corresponding to the products of all distinct pairs of bispectrum components. If the number of bispectrum components is  $K$ , then the number of distinct pairs is  $K(K+1)/2$ . For compute *sna/atom* these columns are appended to existing  $K$  columns. The ordering of quadratic terms is upper-triangular,  $(1,1), (1,2), \dots, (1,K), (2,1), \dots, (K-1,K-1), (K-1,K), (K,K)$ . For computes *snad/atom* and *snav/atom* each set of  $K(K+1)/2$  additional columns is inserted directly after each of sub-block of linear terms i.e. linear and quadratic terms are contiguous. So the nesting order from inside to outside is bispectrum component, linear then quadratic, vector/tensor component, type.

If the *chem* keyword is used, then the data is arranged into  $N_{elem}^3$  sub-blocks, each sub-block corresponding to a particular chemical labeling  $\kappa\lambda\mu$  with the last label changing fastest. Each sub-block contains  $K$  bispectrum components. For the purposes of handling contributions to force, virial, and quadratic combinations, these  $N_{elem}^3$  sub-blocks are treated as a single block of  $KN_{elem}^3$  columns.

If the *bik* keyword is set to 1, the structure of the snap array is expanded. The first  $N$  rows of the snap array correspond to  $B_{i,k}$  instead of a single row summed over atoms  $i$ . In this case, the entries in the final column for these rows are set to zero. Also, each row contains only non-zero entries for the columns corresponding to the type of that atom. This is not true in the case of *dgradflag* keyword = 1 (see below).

If the *dgradflag* keyword is set to 1, this changes the structure of the global array completely. Here the *snad/atom* quantities are replaced with rows corresponding to descriptor gradient components on single atoms:

$$\frac{\partial B_{i,k}}{\partial r_j^a}$$

where  $r_j^a$  is the  $a$ -th position coordinate of the atom with global index  $j$ . The rows are organized in chunks, where each chunk corresponds to an atom with global index  $j$ . The rows in an atom  $j$  chunk correspond to atoms with global index  $i$ . The total number of rows for these descriptor gradients is therefore  $3N^2$ . The number of columns is equal to the number of bispectrum components, plus 3 additional left-most columns representing the global atom indices  $i$ ,  $j$ , and Cartesian direction  $a$  (0, 1, 2, for  $x$ ,  $y$ ,  $z$ ). The first 3 columns of the first  $N$  rows belong to the reference potential force components. The remaining  $K$  columns contain the  $B_{i,k}$  per-atom descriptors corresponding to the non-zero entries obtained when *bikflag* = 1. The first column of the last row, after the first  $N+3N^2$  rows, contains the reference potential energy. The virial components are not used with this option. The total number of rows is therefore  $N+3N^2+1$  and the number of columns is  $K+3$ .

These values can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options. To see how this command can be used within a Python workflow to train SNAP potentials, see the examples in [FitSNAP](#).

### 18.114.5 Restrictions

These computes are part of the ML-SNAP package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 18.114.6 Related commands

*pair\_style snap*

### 18.114.7 Default

The optional keyword defaults are *rmin0* = 0, *switchflag* = 1, *bzeroflag* = 1, *quadraticflag* = 0, *bnormflag* = 0, *wselfallflag* = 0, *switchinnerflag* = 0,

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## 18.115 compute sph/e/atom command

### 18.115.1 Syntax

```
compute ID group-ID sph/e/atom
```

- ID, group-ID are documented in *compute* command
- sph/e/atom = style name of this compute command

### 18.115.2 Examples

```
compute 1 all sph/e/atom
```

### 18.115.3 Description

Define a computation that calculates the per-atom internal energy for each atom in a group.

The internal energy is the energy associated with the internal degrees of freedom of an SPH particle, i.e. a Smooth-Particle Hydrodynamics particle.

See [this PDF guide](#) to using SPH in LAMMPS.

The value of the internal energy will be 0.0 for atoms not in the specified compute group.

### 18.115.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values will be in energy *units*.

### 18.115.5 Restrictions

This compute is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 18.115.6 Related commands

*dump custom*

### 18.115.7 Default

none

## 18.116 compute sph/rho/atom command

### 18.116.1 Syntax

```
compute ID group-ID sph/rho/atom
```

- ID, group-ID are documented in *compute* command
- sph/rho/atom = style name of this compute command

## 18.116.2 Examples

```
compute 1 all sph/rho/atom
```

### 18.116.3 Description

Define a computation that calculates the per-atom SPH density for each atom in a group, i.e. a Smooth-Particle Hydrodynamics density.

The SPH density is the mass density of an SPH particle, calculated by kernel function interpolation using “pair style sph/rhosum”.

See [this PDF guide](#) to using SPH in LAMMPS.

The value of the SPH density will be 0.0 for atoms not in the specified compute group.

### 18.116.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values will be in mass/volume *units*.

### 18.116.5 Restrictions

This compute is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 18.116.6 Related commands

*dump custom*

### 18.116.7 Default

none

## 18.117 compute sph/t/atom command

### 18.117.1 Syntax

```
compute ID group-ID sph/t/atom
```

- ID, group-ID are documented in [compute](#) command
- sph/t/atom = style name of this compute command

## 18.117.2 Examples

```
compute 1 all sph/t/atom
```

## 18.117.3 Description

Define a computation that calculates the per-atom internal temperature for each atom in a group.

The internal temperature is the ratio of internal energy over the heat capacity associated with the internal degrees of freedom of an SPH particles, i.e. a Smooth-Particle Hydrodynamics particle.

$$T_{int} = E_{int}/C_{V,int}$$

See [this PDF guide](#) to using SPH in LAMMPS.

The value of the internal energy will be 0.0 for atoms not in the specified compute group.

## 18.117.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values will be in temperature *units*.

## 18.117.5 Restrictions

This compute is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 18.117.6 Related commands

*dump custom*

## 18.117.7 Default

none

# 18.118 compute spin command

## 18.118.1 Syntax

```
compute ID group-ID spin
```

- ID, group-ID are documented in *compute* command
- spin = style name of this compute command

## 18.118.2 Examples

```
compute out_mag all spin
```

## 18.118.3 Description

Define a computation that calculates magnetic quantities for a system of atoms having spins.

This compute calculates the following 6 magnetic quantities:

- the three first quantities are the x,y and z coordinates of the total magnetization,
- the fourth quantity is the norm of the total magnetization,
- The fifth quantity is the magnetic energy (in eV),
- The sixth one is referred to as the spin temperature, according to the work of (*Nurdin*).

The simplest way to output the results of the compute spin calculation is to define some of the quantities as variables, and to use the thermo and thermo\_style commands, for example:

```
compute out_mag      all spin

variable mag_z       equal c_out_mag[3]
variable mag_norm    equal c_out_mag[4]
variable temp_mag    equal c_out_mag[6]

thermo               10
thermo_style         custom step v_mag_z v_mag_norm v_temp_mag
```

This series of commands evaluates the total magnetization along z, the norm of the total magnetization, and the magnetic temperature. Three variables are assigned to those quantities. The thermo and thermo\_style commands print them every 10 timesteps.

## 18.118.4 Output info

The array values are “intensive”. The array values will be in metal units (*units*).

## 18.118.5 Restrictions

The *spin* compute is part of the SPIN package. This compute is only enabled if LAMMPS was built with this package. See the *Build package* page for more info. The atom\_style has to be “spin” for this compute to be valid.

### Related commands:

none

## 18.118.6 Default

none

(Nurdin) Nurdin and Schotte Phys Rev E, 61(4), 3579 (2000)

## 18.119 compute stress/atom command

## 18.120 compute centroid/stress/atom command

### 18.120.1 Syntax

```
compute ID group-ID style temp-ID keyword ...
```

- ID, group-ID are documented in *compute* command
- style = *stress/atom* or *centroid/stress/atom*
- temp-ID = ID of compute that calculates temperature, can be NULL if not needed
- zero or more keywords may be appended
- keyword = *ke* or *pair* or *bond* or *angle* or *dihedral* or *improper* or *kspace* or *fix* or *virial*

### 18.120.2 Examples

```
compute 1 mobile stress/atom NULL
compute 1 mobile stress/atom myRamp
compute 1 all stress/atom NULL pair bond
compute 1 all centroid/stress/atom NULL bond dihedral improper
```

### 18.120.3 Description

Define a computation that computes per-atom stress tensor for each atom in a group. In case of compute *stress/atom*, the tensor for each atom is symmetric with 6 components and is stored as a 6-element vector in the following order: *xx*, *yy*, *zz*, *xy*, *xz*, *yz*. In case of compute *centroid/stress/atom*, the tensor for each atom is asymmetric with 9 components and is stored as a 9-element vector in the following order: *xx*, *yy*, *zz*, *xy*, *xz*, *yz*, *yx*, *zx*, *zy*. See the *compute pressure* command if you want the stress tensor (pressure) of the entire system.

The stress tensor for atom *I* is given by the following formula, where *a* and *b* take on values *x*, *y*, *z* to generate the components of the tensor:

$$S_{ab} = -mv_a v_b - W_{ab}$$

The first term is a kinetic energy contribution for atom *I*. See details below on how the specified *temp-ID* can affect the velocities used in this calculation. The second term is the virial contribution due to intra and intermolecular interactions, where the exact computation details are determined by the compute style.

In case of compute *stress/atom*, the virial contribution is:

$$\begin{aligned}
 W_{ab} = & \frac{1}{2} \sum_{n=1}^{N_p} (r_{1_a} F_{1_b} + r_{2_a} F_{2_b}) + \frac{1}{2} \sum_{n=1}^{N_b} (r_{1_a} F_{1_b} + r_{2_a} F_{2_b}) \\
 & + \frac{1}{3} \sum_{n=1}^{N_a} (r_{1_a} F_{1_b} + r_{2_a} F_{2_b} + r_{3_a} F_{3_b}) + \frac{1}{4} \sum_{n=1}^{N_d} (r_{1_a} F_{1_b} + r_{2_a} F_{2_b} + r_{3_a} F_{3_b} + r_{4_a} F_{4_b}) \\
 & + \frac{1}{4} \sum_{n=1}^{N_i} (r_{1_a} F_{1_b} + r_{2_a} F_{2_b} + r_{3_a} F_{3_b} + r_{4_a} F_{4_b}) + \text{Kspace}(r_{i_a}, F_{i_b}) + \sum_{n=1}^{N_f} r_{i_a} F_{i_b}
 \end{aligned}$$

The first term is a pairwise energy contribution where  $n$  loops over the  $N_p$  neighbors of atom  $I$ ,  $\mathbf{r}_1$  and  $\mathbf{r}_2$  are the positions of the 2 atoms in the pairwise interaction, and  $\mathbf{F}_1$  and  $\mathbf{F}_2$  are the forces on the 2 atoms resulting from the pairwise interaction. The second term is a bond contribution of similar form for the  $N_b$  bonds which atom  $I$  is part of. There are similar terms for the  $N_a$  angle,  $N_d$  dihedral, and  $N_i$  improper interactions atom  $I$  is part of. There is also a term for the KSpace contribution from long-range Coulombic interactions, if defined. Finally, there is a term for the  $N_f$  *fixes* that apply internal constraint forces to atom  $I$ . Currently, only the *fix shake* and *fix rigid* commands contribute to this term. As the coefficients in the formula imply, a virial contribution produced by a small set of atoms (e.g. 4 atoms in a dihedral or 3 atoms in a Tersoff 3-body interaction) is assigned in equal portions to each atom in the set. E.g. 1/4 of the dihedral virial to each of the 4 atoms, or 1/3 of the fix virial due to SHAKE constraints applied to atoms in a water molecule via the *fix shake* command. As an exception, the virial contribution from constraint forces in *fix rigid* on each atom is computed from the constraint force acting on the corresponding atom and its position, i.e. the total virial is not equally distributed.

In case of compute *centroid/stress/atom*, the virial contribution is:

$$\begin{aligned}
 W_{ab} = & \sum_{n=1}^{N_p} r_{I O_a} F_{I_b} + \sum_{n=1}^{N_b} r_{I O_a} F_{I_b} + \sum_{n=1}^{N_a} r_{I O_a} F_{I_b} + \sum_{n=1}^{N_d} r_{I O_a} F_{I_b} + \sum_{n=1}^{N_i} r_{I O_a} F_{I_b} \\
 & + \text{Kspace}(r_{i_a}, F_{i_b}) + \sum_{n=1}^{N_f} r_{i_a} F_{i_b}
 \end{aligned}$$

As with compute *stress/atom*, the first, second, third, fourth and fifth terms are pairwise, bond, angle, dihedral and improper contributions, but instead of assigning the virial contribution equally to each atom, only the force  $\mathbf{F}_I$  acting on atom  $I$  due to the interaction and the relative position  $\mathbf{r}_{I O}$  of the atom  $I$  to the geometric center of the interacting atoms, i.e. centroid, is used. As the geometric center is different for each interaction, the  $\mathbf{r}_{I O}$  also differs. The sixth term, Kspace contribution, is computed identically to compute *stress/atom*. The seventh term is handed differently depending on if the constraint forces are due to *fix shake* or *fix rigid*. In case of SHAKE constraints, each distance constraint is handed as a pairwise interaction. E.g. in case of a water molecule, two OH and one HH distance constraints are treated as three pairwise interactions. In case of *fix rigid*, all constraint forces in the molecule are treated as a single many-body interaction with a single centroid position. In case of water molecule, the formula expression would become identical to that of the three-body angle interaction. Although the total system virial is the same as compute *stress/atom*, compute *centroid/stress/atom* is known to result in more consistent heat flux values for angle, dihedrals, improper and constraint force contributions when computed via *compute heat/flux*.

If no extra keywords are listed, the kinetic contribution all of the virial contribution terms are included in the per-atom stress tensor. If any extra keywords are listed, only those terms are summed to compute the tensor. The *virial* keyword means include all terms except the kinetic energy *ke*.

Note that the stress for each atom is due to its interaction with all other atoms in the simulation, not just with other atoms in the group.

Details of how compute *stress/atom* obtains the virial for individual atoms for either pairwise or many-body potentials, and including the effects of periodic boundary conditions is discussed in (*Thompson*). The basic idea for many-body potentials is to treat each component of the force computation between a small cluster of atoms in the same manner as in the formula above for bond, angle, dihedral, etc interactions. Namely the quantity  $\mathbf{r} \cdot \mathbf{F}$  is summed over the atoms in



the interaction, with the  $r$  vectors unwrapped by periodic boundaries so that the cluster of atoms is close together. The total contribution for the cluster interaction is divided evenly among those atoms.

Details of how compute *centroid/stress/atom* obtains the virial for individual atoms are given in ([Surblys2019](#)) and ([Surblys2021](#)), where the idea is that the virial of the atom  $I$  is the result of only the force  $\mathbf{F}_I$  on the atom due to the interaction and its positional vector  $\mathbf{r}_{I0}$ , relative to the geometric center of the interacting atoms, regardless of the number of participating atoms. The periodic boundary treatment is identical to that of compute *stress/atom*, and both of them reduce to identical expressions for two-body interactions, i.e. computed values for contributions from bonds and two-body pair styles, such as *Lennard-Jones*, will be the same, while contributions from angles, dihedrals and impropers will be different.

The *dihedral\_style charmm* style calculates pairwise interactions between 1-4 atoms. The virial contribution of these terms is included in the pair virial, not the dihedral virial.

The KSpace contribution is calculated using the method in ([Heyes](#)) for the Ewald method and by the methodology described in ([Sirk](#)) for PPPM. The choice of KSpace solver is specified by the *kpace\_style pppm* command. Note that for PPPM, the calculation requires 6 extra FFTs each timestep that per-atom stress is calculated. Thus it can significantly increase the cost of the PPPM calculation if it is needed on a large fraction of the simulation timesteps.

The *temp-ID* argument can be used to affect the per-atom velocities used in the kinetic energy contribution to the total stress. If the kinetic energy is not included in the stress, than the temperature compute is not used and can be specified as NULL. If the kinetic energy is included and you wish to use atom velocities as-is, then *temp-ID* can also be specified as NULL. If desired, the specified temperature compute can be one that subtracts off a bias to leave each atom with only a thermal velocity to use in the formula above, e.g. by subtracting a background streaming velocity. See the doc pages for individual *compute commands* to determine which ones include a bias.

Note that as defined in the formula, per-atom stress is the negative of the per-atom pressure tensor. It is also really a stress\*volume formulation, meaning the computed quantity is in units of pressure\*volume. It would need to be divided by a per-atom volume to have units of stress (pressure), but an individual atom's volume is not well defined or easy to compute in a deformed solid or a liquid. See the *compute voronoi/atom* command for one possible way to estimate a per-atom volume.

Thus, if the diagonal components of the per-atom stress tensor are summed for all atoms in the system and the sum is divided by  $dV$ , where  $d$  = dimension and  $V$  is the volume of the system, the result should be  $-P$ , where  $P$  is the total pressure of the system.

These lines in an input script for a 3d system should yield that result. I.e. the last 2 columns of thermo output will be the same:

```
compute      peratom all stress/atom NULL
compute      p all reduce sum c_peratom[1] c_peratom[2] c_peratom[3]
variable     press equal -(c_p[1]+c_p[2]+c_p[3])/(3*vol)
thermo_style custom step temp etotal press v_press
```

**Note:** The per-atom stress does not include any Lennard-Jones tail corrections to the pressure added by the *pair\_modify tail yes* command, since those are contributions to the global system pressure.

The compute *stress/atom* can be used in a number of ways. Here is an example to compute a 1-d pressure profile in z-direction across the complete simulation box. You will need to adjust the number of bins and the selections for time averaging to your specific simulation. This assumes that the dimensions of the simulation cell does not change.

```
# set number of bins
variable nbins index 20
variable fraction equal 1.0/v_nbins
```

(continues on next page)

(continued from previous page)

```
# define bins as chunks
compute cchunk all chunk/atom bin/1d x lower ${fraction} units reduced
compute stress all stress/atom NULL
# apply conversion to pressure early since we have no variable style for processing.
↪ chunks
variable press atom -(c_stress[1]+c_stress[2]+c_stress[3])/(3.0*vol*${fraction})
compute binpress all reduce/chunk cchunk sum v_press
fix avg all ave/time 10 40 400 c_binpress mode vector file ave_stress.txt
```

### 18.120.4 Output info

Compute *stress/atom* calculates a per-atom array with 6 columns, which can be accessed by indices 1-6 by any command that uses per-atom values from a compute as input. Compute *centroid/stress/atom* produces a per-atom array with 9 columns, but otherwise can be used in an identical manner to compute *stress/atom*. See the *Howto output* page for an overview of LAMMPS output options.

The ordering of the 6 columns for *stress/atom* is as follows: xx, yy, zz, xy, xz, yz. The ordering of the 9 columns for *centroid/stress/atom* is as follows: xx, yy, zz, xy, xz, yz, yx, zx, zy.

The per-atom array values will be in pressure\*volume *units* as discussed above.

### 18.120.5 Restrictions

Currently, compute *centroid/stress/atom* does not support pair styles with many-body interactions (*EAM* is an exception, since its computations are performed pairwise), nor granular pair styles with pairwise forces which are not aligned with the vector between the pair of particles. All bond styles are supported. All angle, dihedral, improper styles are supported with the exception of INTEL and KOKKOS variants of specific styles. It also does not support models with long-range Coulombic or dispersion forces, i.e. the *kpace\_style* command in LAMMPS. It also does not implement the following fixes which add rigid-body constraints: *fix rigid/\** and the OpenMP accelerated version of *fix rigid/small*, while all other *fix rigid/\*small* are implemented.

LAMMPS will generate an error if one of these options is included in your model. Extension of centroid stress calculations to these force and fix styles is planned for the future.

### 18.120.6 Related commands

*compute pe*, *compute pressure*

### 18.120.7 Default

none

(Heyes) Heyes, Phys Rev B, 49, 755 (1994).

(Sirk) Sirk, Moore, Brown, J Chem Phys, 138, 064505 (2013).

(Thompson) Thompson, Plimpton, Mattson, J Chem Phys, 131, 154107 (2009).

(Surblys2019) Surblys, Matsubara, Kikugawa, Ohara, Phys Rev E, 99, 051301(R) (2019).

(Surblys2021) Surblys, Matsubara, Kikugawa, Ohara, J Appl Phys 130, 215104 (2021).

## 18.121 compute stress/mop command

## 18.122 compute stress/mop/profile command

### 18.122.1 Syntax

```
compute ID group-ID style dir args keywords ...
```

- ID, group-ID are documented in *compute* command
- style = *stress/mop* or *stress/mop/profile*
- dir = x or y or z is the direction normal to the plane
- args = argument specific to the compute style
- keywords = *kin* or *conf* or *total* (one of more can be specified)

*stress/mop* args = pos

pos = *lower* or *center* or *upper* or coordinate value (distance units) is the position of the plane

*stress/mop/profile* args = origin delta

origin = *lower* or *center* or *upper* or coordinate value (distance units) is the position of the first plane

delta = value (distance units) is the distance between planes

### 18.122.2 Examples

```
compute 1 all stress/mop x lower total
compute 1 liquid stress/mop z 0.0 kin conf
fix 1 all ave/time 10 1000 10000 c_1[*] file mop.time
fix 1 all ave/time 10 1000 10000 c_1[2] file mop.time

compute 1 all stress/mop/profile x lower 0.1 total
compute 1 liquid stress/mop/profile z 0.0 0.25 kin conf
fix 1 all ave/time 500 20 10000 c_1[*] ave running overwrite file mopp.time mode vector
```

### 18.122.3 Description

Compute *stress/mop* and compute *stress/mop/profile* define computations that calculate components of the local stress tensor using the method of planes (*Todd*). Specifically in compute *stress/mop* calculates 3 components are computed in directions *dir,x*; *dir,y*; and *dir,z*; where *dir* is the direction normal to the plane, while in compute *stress/mop/profile* the profile of the stress is computed.

Contrary to methods based on histograms of atomic stress (i.e., using *compute stress/atom*), the method of planes is compatible with mechanical balance in heterogeneous systems and at interfaces (*Todd*).

The stress tensor is the sum of a kinetic term and a configurational term, which are given respectively by Eq. (21) and Eq. (16) in (*Todd*). For the kinetic part, the algorithm considers that atoms have crossed the plane if their positions at times  $t - \Delta t$  and  $t$  are one on either side of the plane, and uses the velocity at time  $t - \Delta t/2$  given by the velocity Verlet algorithm.

Between one and three keywords can be used to indicate which contributions to the stress must be computed: kinetic stress (*kin*), configurational stress (*conf*), and/or total stress (*total*).

NOTE 1: The configurational stress is computed considering all pairs of atoms where at least one atom belongs to group *group-ID*.

NOTE 2: The local stress does not include any Lennard-Jones tail corrections to the stress added by the *pair\_modify tail yes* command, since those are contributions to the global system pressure.

NOTE 3: The local stress profile generated by compute *stress/mop/profile* is similar to that obtained by compute *stress/cartesian*. A key difference is that compute *stress/mop/profile* considers particles crossing a set of planes, while compute *stress/cartesian* computes averages for a set of small volumes. More information on the similarities and differences can be found in (*Ikeshoji*).

### 18.122.4 Output info

Compute *stress/mop* calculates a global vector (indices starting at 1), with 3 values for each declared keyword (in the order the keywords have been declared). For each keyword, the stress tensor components are ordered as follows: *stress\_dir,x*, *stress\_dir,y*, and *stress\_dir,z*.

Compute *stress/mop/profile* instead calculates a global array, with 1 column giving the position of the planes where the stress tensor was computed, and with 3 columns of values for each declared keyword (in the order the keywords have been declared). For each keyword, the profiles of stress tensor components are ordered as follows: *stress\_dir,x*; *stress\_dir,y*; and *stress\_dir,z*.

The values are in pressure *units*.

The values produced by this compute can be accessed by various *output commands*. For instance, the results can be written to a file using the *fix ave/time* command. Please see the example in the *examples/PACKAGES/mop* folder.

### 18.122.5 Restrictions

These styles are part of the EXTRA-COMPUTE package. They are only enabled if LAMMPS is built with that package. See the *Build package* doc page on for more info.

The method is only implemented for 3d orthogonal simulation boxes whose size does not change in time, and axis-aligned planes.

The method only works with two-body pair interactions, because it requires the class method *pair->single()* to be implemented. In particular, it does not work with more than two-body pair interactions, intra-molecular interactions, and long range (*k-space*) interactions.

### 18.122.6 Related commands

*compute stress/atom*, *compute pressure*, *compute stress/cartesian*, *compute stress/cylinder*, *compute stress/spherical*

## 18.122.7 Default

none

(**Todd**) B. D. Todd, Denis J. Evans, and Peter J. Daivis: “Pressure tensor for inhomogeneous fluids”, Phys. Rev. E 52, 1627 (1995).

(**Ikeshoji**) Ikeshoji, Hafskjold, Furuholt, Mol Sim, 29, 101-109, (2003).

## 18.123 compute stress/cartesian command

## 18.124 compute stress/cylinder command

## 18.125 compute stress/spherical command

### 18.125.1 Syntax

```
compute ID group-ID style args
```

- ID, group-ID are documented in *compute* command
- style = stress/cartesian or stress/spherical or stress/cylinder
- args = argument specific to the compute style

*stress/cartesian* args = dim bin\_width

dim = x or y or z

bin\_width = width of the bin

one or two dim/bin\_width pairs may be appended

*stress/cylinder* args = zlo zh Rmax bin\_width keyword

zlo = minimum z-boundary for cylinder

zhi = maximum z-boundary for cylinder

Rmax = maximum radius to perform calculation to

bin\_width = width of radial bins to use for calculation

keyword = ke (zero or one can be specified)

ke = yes or no

*stress/spherical*

x0, y0, z0 = origin of the spherical coordinate system

bin\_width = width of spherical shells

Rmax = maximum radius of spherical shells

### 18.125.2 Examples

```
compute 1 all stress/cartesian x 0.1
compute 1 all stress/cartesian y 0.25 z 0.1
compute 1 all stress/cylinder -10.0 10.0 15.0 0.25
compute 1 all stress/cylinder -10.0 10.0 15.0 0.25 ke no
compute 1 all stress/spherical 0 0 0 0.1 10
```

### 18.125.3 Description

Compute *stress/cartesian*, compute *stress/cylinder*, and compute *stress/spherical* define computations that calculate profiles of the diagonal components of the local stress tensor in the specified coordinate system. The stress tensor is split into a kinetic contribution  $P^k$  and a virial contribution  $P^v$ . The sum gives the total stress tensor  $P = P^k + P^v$ . These computes can for example be used to calculate the diagonal components of the local stress tensor of interfaces with flat, cylindrical, or spherical symmetry. These computes obeys momentum balance through fluid interfaces. They use the Irving–Kirkwood contour, which is the straight line between particle pairs.

The *stress/cartesian* computes the stress profile along one or two Cartesian coordinates, as described in (*Ikeshoji*). The compute *stress/cylinder* computes the stress profile along the radial direction in cylindrical coordinates, as described in (*Addington*). The compute *stress/spherical* computes the stress profile along the radial direction in spherical coordinates, as described in (*Ikeshoji*).

### 18.125.4 Output info

The output columns for *stress/cartesian* are the position of the center of the local volume in the first and second dimensions, number density,  $P_{xx}^k$ ,  $P_{yy}^k$ ,  $P_{zz}^k$ ,  $P_{xx}^v$ ,  $P_{yy}^v$ , and  $P_{zz}^v$ . There are 8 columns when one dimension is specified and 9 columns when two dimensions are specified. The number of bins (rows) is  $(L_1/b_1)(L_2/b_2)$ , where  $L_1$  and  $L_2$  are the lengths of the simulation box in the specified dimensions and  $b_1$  and  $b_2$  are the specified bin widths. When only one dimension is specified, the number of bins (rows) is  $L_1/b_1$ .

The default output columns for *stress/cylinder* are the radius to the center of the cylindrical shell, number density,  $P_{rr}^k$ ,  $P_{\phi\phi}^k$ ,  $P_{zz}^k$ ,  $P_{rr}^v$ ,  $P_{\phi\phi}^v$ , and  $P_{zz}^v$ . When the keyword *ke* is set to *no*, the kinetic contributions are not calculated, and consequently there are only 5 columns: the position of the center of the cylindrical shell, the number density,  $P_{rr}^v$ ,  $P_{\phi\phi}^v$ , and  $P_{zz}^v$ . The number of bins (rows) is  $R_{\max}/b$ , where  $b$  is the specified bin width.

The output columns for *stress/spherical* are the position of the center of the spherical shell, the number density,  $P_{rr}^k$ ,  $P_{\theta\theta}^k$ ,  $P_{\phi\phi}^k$ ,  $P_{rr}^v$ ,  $P_{\theta\theta}^v$ , and  $P_{\phi\phi}^v$ . There are 8 columns and the number of bins (rows) is  $R_{\max}/b$ , where  $b$  is the specified bin width.

This array can be output with *fix ave/time*,

```
compute p all stress/cartesian x 0.1
fix 2 all ave/time 100 1 100 c_p[*] file dump_p.out mode vector
```

The values calculated by this compute are “intensive”. The stress values will be in pressure *units*. The number density values are in inverse volume *units*.

NOTE 1: The local stress does not include any Lennard-Jones tail corrections to the stress added by the *pair\_modify tail yes* command, since those are contributions to the global system pressure.

NOTE 2: The local stress profiles generated by these computes are similar to those obtained by the *method-of-planes (MOP)*. A key difference is that compute *stress/mop/profile <compute\_stress\_mop>* considers particles crossing a set of planes, while *stress/cartesian* computes averages for a set of small volumes. More information on the similarities and differences can be found in (*Ikeshoji*).

### 18.125.5 Restrictions

These computes calculate the stress tensor contributions for pair styles only (i.e., no bond, angle, dihedral, etc. contributions, and in the presence of bonded interactions, the result will be incorrect due to exclusions for special bonds) and requires pairwise force calculations not available for most many-body pair styles. Note that  $k$ -space calculations are also excluded.

These computes are part of the EXTRA-COMPUTE package. They are only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

### 18.125.6 Related commands

*compute stress/atom*, *compute pressure*, *compute stress/mop/profile*

### 18.125.7 Default

The keyword default for ke in style *stress/cylinder* is yes.

---

**(Ikeshoji)** Ikeshoji, Hafskjold, Furuholt, Mol Sim, 29, 101-109, (2003).

**(Addington)** Addington, Long, Gubbins, J Chem Phys, 149, 084109 (2018).

## 18.126 compute force/tally command

### 18.127 compute heat/flux/tally command

### 18.128 compute heat/flux/virial/tally command

### 18.129 compute pe/tally command

### 18.130 compute pe/mol/tally command

## 18.131 compute stress/tally command

### 18.131.1 Syntax

```
compute ID group-ID style group2-ID
```

- ID, group-ID are documented in *compute* command
- style = *force/tally* or *heat/flux/tally* or *heat/flux/virial/tally* or *pe/tally* or *pe/mol/tally* or *stress/tally*
- group2-ID = group ID of second (or same) group

## 18.131.2 Examples

```
compute 1 lower force/tally upper
compute 1 left pe/tally right
compute 1 lower stress/tally lower
compute 1 subregion heat/flux/tally all
compute 1 liquid heat/flux/virial/tally solid
```

## 18.131.3 Description

Define a computation that calculates properties between two groups of atoms by accumulating them from pairwise non-bonded computations. Except for *heat/flux/virial/tally*, the two groups can be the same. This is similar to *compute group/group* only that the data is accumulated directly during the non-bonded force computation. The computes *force/tally*, *pe/tally*, *stress/tally*, and *heat/flux/tally* are primarily provided as example how to program additional, more sophisticated computes using the tally callback mechanism. Compute *pe/mol/tally* is one such style, that can—through using this mechanism—separately tally intermolecular and intramolecular energies. Something that would otherwise be impossible without integrating this as a core functionality into the base classes of LAMMPS.

Compute *heat/flux/tally* obtains the heat flux (strictly speaking, heat flow) inside the first group, which is the sum of the convective contribution due to atoms in the first group and the virial contribution due to interaction between the first and second groups:

$$\mathbf{Q} = \sum_{i \in \text{group 1}} e_i \mathbf{v}_i + \frac{1}{2} \sum_{i \in \text{group 1}} \sum_{\substack{j \in \text{group 2} \\ j \neq i}} (\mathbf{F}_{ij} \cdot \mathbf{v}_j) \mathbf{r}_{ij}$$

When the second group in *heat/flux/tally* is set to “all”, the resulting values will be identical to that obtained by *compute heat/flux*, provided only pairwise interactions exist.

Compute *heat/flux/virial/tally* obtains the total virial heat flux (strictly speaking, heat flow) into the first group due to interaction with the second group, and is defined as:

$$Q = \frac{1}{2} \sum_{i \in \text{group 1}} \sum_{j \in \text{group 2}} \mathbf{F}_{ij} \cdot (\mathbf{v}_i + \mathbf{v}_j)$$

Although, the *heat/flux/virial/tally* compute does not include the convective term, it can be used to obtain the total heat flux over control surfaces, when there are no particles crossing over, such as is often in solid–solid and solid–liquid interfaces. This would be identical to the method of planes method. Note that the *heat/flux/virial/tally* compute is distinctly different from the *heat/flux* and *heat/flux/tally* computes, that are essentially volume averaging methods. The following example demonstrates the difference:

```
# System with only pairwise interactions.
# Non-periodic boundaries in the x direction.
# Has LeftLiquid and RightWall groups along x direction.

# Heat flux over the solid-liquid interface
compute hflow_hfvt RightWall heat/flux/virial/tally LeftLiquid
variable hflux_hfvt equal c_hflow_hfvt/(ly*lz)

# x component of approximate heat flux vector inside the liquid region,
# two approaches.
#
```

(continues on next page)



(continued from previous page)

```

compute myKE all ke/atom
compute myPE all pe/atom
compute myStress all stress/atom NULL virial
compute hflow_hf LeftLiquid heat/flux myKE myPE myStress
variable hflux_hf equal c_hflow_hf[1]/${volLiq}
#
compute hflow_hft LeftLiquid heat/flux/tally all
variable hflux_hft equal c_hflow_hft[1]/${volLiq}

# Pressure over the solid-liquid interface, three approaches.
#
compute force_gg RightWall group/group LeftLiquid
variable press_gg equal c_force_gg[1]/(ly*lz)
#
compute force_ft RightWall force/tally LeftLiquid
compute rforce_ft RightWall reduce sum c_force_ft[1]
variable press_ft equal c_rforce_ft/(ly*lz)
#
compute rforce_hfvt all reduce sum c_hflow_hfvt[1]
variable press_hfvt equal c_rforce_hfvt/(ly*lz)

```

The pairwise contributions are computed via a callback that the compute registers with the non-bonded pairwise force computation. This limits the use to systems that have no bonds, no Kspace, and no many-body interactions. On the other hand, the computation does not have to compute forces or energies a second time and thus can be much more efficient. The callback mechanism allows to write more complex pairwise property computations.

#### 18.131.4 Output info

- Compute *pe/tally* calculates a global scalar (the energy) and a per atom scalar (the contributions of the single atom to the global scalar).
- Compute *pe/mol/tally* calculates a global four-element vector containing (in this order): *evdwl* and *ecoul* for intramolecular pairs and *evdwl* and *ecoul* for intermolecular pairs. Since molecules are identified by their molecule IDs, the partitioning does not have to be related to molecules, but the energies are tallied into the respective slots depending on whether the molecule IDs of a pair are the same or different.
- Compute *force/tally* calculates a global scalar (the force magnitude) and a per atom 3-element vector (force contribution from each atom).
- Compute *stress/tally* calculates a global scalar (average of the diagonal elements of the stress tensor) and a per atom vector (the six elements of stress tensor contributions from the individual atom).
- As in *compute heat/flux*, compute *heat/flux/tally* calculates a global vector of length 6, where the first three components are the *x*, *y*, *z* components of the full heat flow vector, and the next three components are the corresponding components of just the convective portion of the flow (i.e., the first term in the equation for *Q*).
- Compute *heat/flux/virial/tally* calculates a global scalar (heat flow) and a per atom three-element vector (contribution to the force acting over atoms in the first group from individual atoms in both groups).

Both the scalar and vector values calculated by this compute are “extensive”.

### 18.131.5 Restrictions

This compute is part of the TALLY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Not all pair styles can be evaluated in a pairwise mode as required by this compute. For example, 3-body and other many-body potentials, such as *Tersoff* and *Stillinger-Weber* cannot be used. *EAM* potentials only include the pair potential portion of the EAM interaction when used by this compute, not the embedding term. Also bonded or Kspace interactions do not contribute to this compute.

When used with dynamic groups, a *run 0* command needs to be inserted in order to initialize the dynamic groups before accessing the computes.

### 18.131.6 Related commands

- *compute group/group*
- *compute heat/flux*

### 18.131.7 Default

none

## 18.132 compute tdpd/cc/atom command

### 18.132.1 Syntax

```
compute ID group-ID tdpd/cc/atom index
```

- ID, group-ID are documented in *compute* command
- tdpd/cc/atom = style name of this compute command
- index = index of chemical species (1 to Nspecies)

### 18.132.2 Examples

```
compute 1 all tdpd/cc/atom 2
```

### 18.132.3 Description

Define a computation that calculates the per-atom chemical concentration of a specified species for each tDPD particle in a group.

The chemical concentration of each species is defined as the number of molecules carried by a tDPD particle for dilute solution. For more details see (*Li2015*).

### 18.132.4 Output info

This compute calculates a per-atom vector, which can be accessed by any command that uses per-atom values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The per-atom vector values will be in the units of chemical species per unit mass.

### 18.132.5 Restrictions

This compute is part of the DPD-MESO package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 18.132.6 Related commands

*pair\_style tdpd*

### 18.132.7 Default

none

---

(Li2015) Li, Yazdani, Tartakovsky, Karniadakis, J Chem Phys, 143: 014101 (2015). DOI: 10.1063/1.4923254

## 18.133 compute temp command

Accelerator Variants: *temp/kk*

### 18.133.1 Syntax

```
compute ID group-ID temp
```

- ID, group-ID are documented in [compute](#) command
- temp = style name of this compute command

### 18.133.2 Examples

```
compute 1 all temp  
compute myTemp mobile temp
```

### 18.133.3 Description

Define a computation that calculates the temperature of a group of atoms. A compute of this style can be used by any command that computes a temperature (e.g., *thermo\_modify*, *fix temp/rescale*, *fix npt*)

The temperature is calculated by the formula

$$\text{KE} = \frac{\text{dim}}{2} N k_B T,$$

where KE = total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ ), dim = 2 or 3 is the dimensionality of the simulation,  $N$  is the number of atoms in the group,  $k_B$  is the Boltzmann constant, and  $T$  is the absolute temperature.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the *compute\_modify* command if this is not the case.

This compute subtracts out degrees-of-freedom due to fixes that constrain molecular motion, such as *fix shake* and *fix rigid*. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the *extra* option of the *compute\_modify* command.

A compute of this style with the ID of “thermo\_temp” is created when LAMMPS starts up, as if this command were in the input script:

```
compute thermo_temp all temp
```

See the “thermo\_style” command for more details.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostating.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 18.133.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length six (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.133.5 Restrictions

none

### 18.133.6 Related commands

*compute temp/partial, compute temp/region, compute pressure*

### 18.133.7 Default

none

## 18.134 compute temp/asphere command

### 18.134.1 Syntax

```
compute ID group-ID temp/asphere keyword value ...
```

- ID, group-ID are documented in *compute* command
- temp/asphere = style name of this compute command
- zero or more keyword/value pairs may be appended
- keyword = *bias* or *dof*

*bias* value = bias-ID

bias-ID = ID of a temperature compute that removes a velocity bias

*dof* value = *all* or *rotate*

*all* = compute temperature of translational and rotational degrees of freedom

*rotate* = compute temperature of just rotational degrees of freedom

### 18.134.2 Examples

```
compute 1 all temp/asphere
compute myTemp mobile temp/asphere bias tempCOM
compute myTemp mobile temp/asphere dof rotate
```

### 18.134.3 Description

Define a computation that calculates the temperature of a group of aspherical particles, including a contribution from both their translational and rotational kinetic energy. This differs from the usual *compute temp* command, which assumes point particles with only translational kinetic energy.

Only finite-size particles (aspherical or spherical) can be included in the group. For 3d finite-size particles, each has six degrees of freedom (three translational, three rotational). For 2d finite-size particles, each has three degrees of freedom (two translational, one rotational).

**Note:** This choice for degrees of freedom (DOF) assumes that all finite-size aspherical or spherical particles in your model will freely rotate, sampling all their rotational DOF. It is possible to use a combination of interaction potentials

and fixes that induce no torque or otherwise constrain some of all of your particles so that this is not the case. Then there are fewer DOF and you should use the `compute_modify extra` command to adjust the DOF accordingly.

---

For example, an aspherical particle with all three of its shape parameters the same is a sphere. If it does not rotate, then it should have 3 DOF instead of 6 in 3d (or two instead of three in 2d). A uniaxial aspherical particle has two of its three shape parameters the same. If it does not rotate around the axis perpendicular to its circular cross section, then it should have 5 DOF instead of 6 in 3d. The latter is the case for uniaxial ellipsoids in a *GayBerne model* since there is no induced torque around the optical axis. It will also be the case for biaxial ellipsoids when exactly two of the semiaxes have the same length and the corresponding relative well depths are equal.

The translational kinetic energy is computed the same as is described by the `compute temp` command. The rotational kinetic energy is computed as  $\frac{1}{2}I\omega^2$ , where  $I$  is the inertia tensor for the aspherical particle and  $\omega$  is its angular velocity, which is computed from its angular momentum.

---

**Note:** For *2d models*, particles are treated as ellipsoids, not ellipses, meaning their moments of inertia will be the same as in 3d.

---

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  and  $\omega^2$  are replaced by  $v_x v_y$  and  $\omega_x \omega_y$  for the  $xy$  component, and the appropriate elements of the moment of inertia tensor are used. The six components of the vector are ordered  $xx, yy, zz, xy, xz, yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the `compute_modify` command if this is not the case.

This compute subtracts out translational degrees-of-freedom due to fixes that constrain molecular motion, such as *fix shake* and *fix rigid*. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the *extra* option of the `compute_modify` command.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostatting.

---

The keyword/value option pairs are used in the following ways.

For the *bias* keyword, *bias-ID* refers to the ID of a temperature compute that removes a “bias” velocity from each atom. This allows compute temp/sphere to compute its thermal temperature after the translational kinetic energy components have been altered in a prescribed way (e.g., to remove a flow velocity profile). Thermostats that use this compute will work with this bias term. See the doc pages for individual computes that calculate a temperature and the doc pages for fixes that perform thermostatting for more details.

For the *dof* keyword, a setting of *all* calculates a temperature that includes both translational and rotational degrees of freedom. A setting of *rotate* calculates a temperature that includes only rotational degrees of freedom.

---

### 18.134.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.134.5 Restrictions

This compute is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This compute requires that atoms store angular momentum and a quaternion as defined by the [atom\\_style ellipsoid](#) command.

All particles in the group must be finite-size. They cannot be point particles, but they can be aspherical or spherical as defined by their shape attribute.

### 18.134.6 Related commands

*compute temp*

### 18.134.7 Default

none

## 18.135 compute temp/body command

### 18.135.1 Syntax

```
compute ID group-ID temp/body keyword value ...
```

- ID, group-ID are documented in [compute](#) command
- temp/body = style name of this compute command
- zero or more keyword/value pairs may be appended
- keyword = *bias* or *dof*
  - bias* value = bias-ID
    - bias-ID = ID of a temperature compute that removes a velocity bias
  - dof* value = *all* or *rotate*
    - all = compute temperature of translational and rotational degrees of freedom
    - rotate = compute temperature of just rotational degrees of freedom

## 18.135.2 Examples

```
compute 1 all temp/body
compute myTemp mobile temp/body bias tempCOM
compute myTemp mobile temp/body dof rotate
```

---

## 18.135.3 Description

Define a computation that calculates the temperature of a group of body particles, including a contribution from both their translational and rotational kinetic energy. This differs from the usual `compute temp` command, which assumes point particles with only translational kinetic energy.

Only body particles can be included in the group. For 3d particles, each has 6 degrees of freedom (3 translational, 3 rotational). For 2d body particles, each has 3 degrees of freedom (2 translational, 1 rotational).

---

**Note:** This choice for degrees of freedom (DOF) assumes that all body particles in your model will freely rotate, sampling all their rotational DOF. It is possible to use a combination of interaction potentials and fixes that induce no torque or otherwise constrain some of all of your particles so that this is not the case. Then there are less DOF and you should use the `compute_modify extra` command to adjust the DOF accordingly.

---

The translational kinetic energy is computed the same as is described by the `compute temp` command. The rotational kinetic energy is computed as  $\frac{1}{2}I\omega^2$ , where  $I$  is the moment of inertia tensor for the spherical particle and  $\omega$  is its angular velocity, which is computed from its angular momentum.

A kinetic energy tensor, stored as a 6-element vector, is also calculated by this compute. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  and  $\omega^2$  are replaced by  $v_x v_y$  and  $\omega_x \omega_y$  for the math:xy component, and the appropriate elements of the inertia tensor are used. The six components of the vector are ordered  $xx, yy, zz, xy, xz, yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the `dynamic` option of the `compute_modify` command if this is not the case.

This compute subtracts out translational degrees-of-freedom due to fixes that constrain molecular motion, such as `fix shake` and `fix rigid`. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the `extra` option of the `compute_modify` command.

See the [Howto thermostat](#) page for a discussion of different ways to compute temperature and perform thermostatting.

---

The keyword/value option pairs are used in the following ways.

For the `bias` keyword, `bias-ID` refers to the ID of a temperature compute that removes a “bias” velocity from each atom. This allows compute temp/sphere to compute its thermal temperature after the translational kinetic energy components have been altered in a prescribed way (e.g., to remove a flow velocity profile). Thermostats that use this compute will work with this bias term. See the doc pages for individual computes that calculate a temperature and the doc pages for fixes that perform thermostatting for more details.

For the `dof` keyword, a setting of `all` calculates a temperature that includes both translational and rotational degrees of freedom. A setting of `rotate` calculates a temperature that includes only rotational degrees of freedom.

---



### 18.135.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.135.5 Restrictions

This compute is part of the BODY package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This compute requires that atoms store angular momentum and a quaternion as defined by the [atom\\_style body](#) command.

### 18.135.6 Related commands

*compute temp*

### 18.135.7 Default

none

## 18.136 compute temp/chunk command

### 18.136.1 Syntax

```
compute ID group-ID temp/chunk chunkID value1 value2 ... keyword value ...
```

- ID, group-ID are documented in [compute](#) command
- temp/chunk = style name of this compute command
- chunkID = ID of [compute chunk/atom](#) command
- zero or more values can be listed as value1,value2,etc.
- value = *temp* or *kecom* or *internal*

```
temp = temperature of each chunk
kecom = kinetic energy of each chunk based on velocity of center of mass
internal = internal kinetic energy of each chunk
```

- zero or more keyword/value pairs may be appended
- keyword = *com* or *bias* or *adof* or *cdof*

```
com value = yes or no
  yes = subtract center-of-mass velocity from each chunk before calculating
  ↪ temperature
  no = do not subtract center-of-mass velocity
```

```
bias value = bias-ID
  bias-ID = ID of a temperature compute that removes a velocity bias
adof value = dof_per_atom
  dof_per_atom = define this many degrees-of-freedom per atom
cdof value = dof_per_chunk
  dof_per_chunk = define this many degrees-of-freedom per chunk
```

### 18.136.2 Examples

```
compute 1 fluid temp/chunk molchunk
compute 1 fluid temp/chunk molchunk temp internal
compute 1 fluid temp/chunk molchunk bias tpartial adof 2.0
```

### 18.136.3 Description

Define a computation that calculates the temperature of a group of atoms that are also in chunks, after optionally subtracting out the center-of-mass velocity of each chunk. By specifying optional values, it can also calculate the per-chunk temperature or energies of the multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as *chunkID*. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

The temperature is calculated by the formula

$$\text{KE} = \frac{\text{DOF}}{2} k_B T,$$

where KE is the total kinetic energy of all atoms assigned to chunks (sum of  $\frac{1}{2}mv^2$ ), DOF is the the total number of degrees of freedom for those atoms,  $k_B$  is Boltzmann constant, and  $T$  is the absolute temperature.

The DOF is calculated as  $N \times \text{adof} + N_{\text{chunk}} \times \text{cdof}$ , where  $N$  is the number of atoms contributing to the kinetic energy, *adof* is the number of degrees of freedom per atom, and *cdof* is the number of degrees of freedom per chunk. By default, *adof* = 2 or 3 = dimensionality of system, as set via the *dimension* command, and *cdof* = 0.0. This gives the usual formula for temperature.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the *xy* component, and so on. The six components of the vector are ordered *xx*, *yy*, *zz*, *xy*, *xz*, *yz*.

Note that the number of atoms contributing to the temperature is calculated each time the temperature is evaluated since it is assumed the atoms may be dynamically assigned to chunks. Thus there is no need to use the *dynamic* option of the *compute\_modify* command for this compute style.

If any optional values are specified, then per-chunk quantities are also calculated and stored in a global array, as described below.

The *temp* value calculates the temperature for each chunk by the formula

$$\text{KE} = \frac{\text{DOF}}{2} k_B T,$$

where KE is the total kinetic energy of the chunk of atoms (sum of  $\frac{1}{2}mv^2$ ), DOF is the total number of degrees of freedom for all atoms in the chunk,  $k_B$  is the Boltzmann constant, and  $T$  is the absolute temperature.

The number of degrees of freedom (DOF) in this case is calculated as  $N \times adof + cdof$ , where  $N$  is the number of atoms in the chunk,  $adof$  is the number of degrees of freedom per atom, and  $cdof$  is the number of degrees of freedom per chunk. By default,  $cdof = 2$  or  $3 =$  dimensionality of system, as set via the *dimension* command, and  $cdof = 0.0$ . This gives the usual formula for temperature.

The *kecom* value calculates the kinetic energy of each chunk as if all its atoms were moving with the velocity of the center-of-mass of the chunk.

The *internal* value calculates the internal kinetic energy of each chunk. The internal KE is summed over the atoms in the chunk using an internal “thermal” velocity for each atom, which is its velocity minus the center-of-mass velocity of the chunk.

Note that currently the global and per-chunk temperatures calculated by this compute only include translational degrees of freedom for each atom. No rotational degrees of freedom are included for finite-size particles. Also no degrees of freedom are subtracted for any velocity bias or constraints that are applied, such as *compute temp/partial*, or *fix shake* or *fix rigid*. This is because those degrees of freedom (e.g., a constrained bond) could apply to sets of atoms that are both included and excluded from a specific chunk, and hence the concept is somewhat ill-defined. In some cases, you can use the *adof* and *cdof* keywords to adjust the calculated degrees of freedom appropriately, as explained below.

Note that the per-chunk temperature calculated by this compute and the *fix ave/chunk temp* command can be different. This compute calculates the temperature for each chunk for a single snapshot. *Fix ave/chunk* can do that but can also time average those values over many snapshots, or it can compute a temperature as if the atoms in the chunk on different timesteps were collected together as one set of atoms to calculate their temperature. This compute allows the center-of-mass velocity of each chunk to be subtracted before calculating the temperature; *fix ave/chunk* does not.

**Note:** Only atoms in the specified group contribute to the calculations performed by this compute. The *compute chunk/atom* command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

The simplest way to output the per-chunk results of the *compute temp/chunk* calculation to a file is to use the *fix ave/time* command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk all temp/chunk cc1 temp
fix 1 all ave/time 100 1 100 c_myChunk[1] file tmp.out mode vector
```

The keyword/value option pairs are used in the following ways.

The *com* keyword can be used with a value of *yes* to subtract the velocity of the center-of-mass for each chunk from the velocity of the atoms in that chunk, before calculating either the global or per-chunk temperature. This can be useful if the atoms are streaming or otherwise moving collectively, and you wish to calculate only the thermal temperature.

For the *bias* keyword, *bias-ID* refers to the ID of a temperature compute that removes a “bias” velocity from each atom. This also allows calculation of the global or per-chunk temperature using only the thermal temperature of atoms in each chunk after the translational kinetic energy components have been altered in a prescribed way (e.g., to remove a velocity profile). It also applies to the calculation of the other per-chunk values, such as *kecom* or *internal*, which involve the center-of-mass velocity of each chunk, which is calculated after the velocity bias is removed from each atom. Note that the temperature compute will apply its bias globally to the entire system, not on a per-chunk basis.

The *adof* and *cdof* keywords define the values used in the degree of freedom (DOF) formulas used for the global or per-chunk temperature, as described above. They can be used to calculate a more appropriate temperature for some kinds of chunks. Here are three examples:

If spatially binned chunks contain some number of water molecules and *fix shake* is used to make each molecule rigid, then you could calculate a temperature with six degrees of freedom (DOF) (three translational, three rotational) per molecule by setting *adof* to 2.0.

If *compute temp/partial* is used with the *bias* keyword to only allow the x component of velocity to contribute to the temperature, then *adof* = 1.0 would be appropriate.

If each chunk consists of a large molecule, with some number of its bonds constrained by *fix shake* or the entire molecule by *fix rigid/small*, *adof* = 0.0 and *cdof* could be set to the remaining degrees of freedom for the entire molecule (entire chunk in this case; i.e., 6 for 3d, or 3 for 2d, for a rigid molecule).

---

## 18.136.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

This compute also optionally calculates a global array, if one or more of the optional values are specified. The number of rows in the array is the number of chunks *Nchunk* as calculated by the specified *compute chunk/atom* command. The number of columns is the number of specified values (1 or more). These values can be accessed by any command that uses global array values from a compute as input. Again, see the *Howto output* doc page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”. The array values are “intensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*. The array values will be in temperature *units* for the *temp* value, and in energy *units* for the *kecom* and *internal* values.

## 18.136.5 Restrictions

The *com* and *bias* keywords cannot be used together.

## 18.136.6 Related commands

*compute temp, fix ave/chunk temp*

## 18.136.7 Default

The option defaults are *com* no, no *bias*, *adof* = dimensionality of the system (2 or 3), and *cdof* = 0.0.

# 18.137 compute temp/com command

## 18.137.1 Syntax

```
compute ID group-ID temp/com
```

- ID, group-ID are documented in *compute* command
- temp/com = style name of this compute command

## 18.137.2 Examples

```
compute 1 all temp/com
compute myTemp mobile temp/com
```

## 18.137.3 Description

Define a computation that calculates the temperature of a group of atoms, after subtracting out the center-of-mass velocity of the group. This is useful if the group is expected to have a non-zero net velocity for some reason. A compute of this style can be used by any command that computes a temperature, (e.g., *thermo\_modify*, *fix temp/rescale*, *fix npt*).

After the center-of-mass velocity has been subtracted from each atom, the temperature is calculated by the formula

$$\text{KE} = \frac{\text{dim}}{2} N k_B T,$$

where KE is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ ), dim = 2 or 3 is the dimensionality of the simulation,  $N$  is number of atoms in the group,  $k_B$  is the Boltzmann constant, and  $T$  is the absolute temperature.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the *compute\_modify* command if this is not the case.

The removal of the center-of-mass velocity by this fix is essentially computing the temperature after a “bias” has been removed from the velocity of the atoms. If this compute is used with a fix command that performs thermostating then this bias will be subtracted from each atom, thermostating of the remaining thermal velocity will be performed, and the bias will be added back in. Thermostating fixes that work in this way include *fix nvt*, *fix temp/rescale*, *fix temp/berendsen*, and *fix langevin*.

This compute subtracts out degrees-of-freedom due to fixes that constrain molecular motion, such as *fix shake* and *fix rigid*. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the *extra* option of the *compute\_modify* command.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostating.

## 18.137.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.137.5 Restrictions

none

### 18.137.6 Related commands

*compute temp*

### 18.137.7 Default

none

## 18.138 compute temp/cs command

### 18.138.1 Syntax

```
compute ID group-ID temp/cs group1 group2
```

- ID, group-ID are documented in *compute* command
- temp/cs = style name of this compute command
- group1 = group-ID of either cores or shells
- group2 = group-ID of either shells or cores

### 18.138.2 Examples

```
compute oxygen_c-s all temp/cs 0_core 0_shell  
compute core_shells all temp/cs cores shells
```

### 18.138.3 Description

Define a computation that calculates the temperature of a system based on the center-of-mass velocity of atom pairs that are bonded to each other. This compute is designed to be used with the adiabatic core/shell model of (*Mitchell and Finchham*). See the *Howto coreshell* page for an overview of the model as implemented in LAMMPS. Specifically, this compute enables correct temperature calculation and thermostating of core/shell pairs where it is desirable for the internal degrees of freedom of the core/shell pairs to not be influenced by a thermostat. A compute of this style can be used by any command that computes a temperature via *fix\_modify* (e.g., *fix temp/rescale*, *fix npt*).

Note that this compute does not require all ions to be polarized, hence defined as core/shell pairs. One can mix core/shell pairs and ions without a satellite particle if desired. The compute will consider the non-polarized ions according to the physical system.

For this compute, core and shell particles are specified by two respective group IDs, which can be defined using the *group* command. The number of atoms in the two groups must be the same and there should be one bond defined between a pair of atoms in the two groups. Non-polarized ions which might also be included in the treated system should not be included into either of these groups, they are taken into account by the *group-ID* (second argument) of the compute.

The temperature is calculated by the formula

$$\text{KE} = \frac{\text{dim}}{2} N k_B T,$$

where KE is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ ), dim = 2 or 3 is the dimensionality of the simulation,  $N$  is the number of atoms in the group,  $k_B$  is the Boltzmann constant, and  $T$  is the absolute temperature. Note that the velocity of each core or shell atom used in the KE calculation is the velocity of the center-of-mass (COM) of the core/shell pair the atom is part of.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ . In contrast to the temperature, the velocity of each core or shell atom is taken individually.

The change this fix makes to core/shell atom velocities is essentially computing the temperature after a “bias” has been removed from the velocity of the atoms. This “bias” is the velocity of the atom relative to the center-of-mass velocity of the core/shell pair. If this compute is used with a fix command that performs thermostating then this bias will be subtracted from each atom, thermostating of the remaining center-of-mass velocity will be performed, and the bias will be added back in. This means the thermostating will effectively be performed on the core/shell pairs, instead of on the individual core and shell atoms. Thermostating fixes that work in this way include *fix nvt*, *fix temp/rescale*, *fix temp/berendsen*, and *fix langevin*.

The internal energy of core/shell pairs can be calculated by the *compute temp/chunk* command, if chunks are defined as core/shell pairs. See the *Howto coreshell* doc page for more discussion on how to do this.

### 18.138.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.138.5 Restrictions

The number of core/shell pairs contributing to the temperature is assumed to be constant for the duration of the run. No fixes should be used which generate new molecules or atoms during a simulation.

### 18.138.6 Related commands

*compute temp*, *compute temp/chunk*

## 18.138.7 Default

none

---

(Mitchell and Finchham) Mitchell, Finchham, J Phys Condensed Matter, 5, 1031-1038 (1993).

## 18.139 compute temp/deform command

Accelerator Variants: *temp/deform/kk*

### 18.139.1 Syntax

```
compute ID group-ID temp/deform
```

- ID, group-ID are documented in *compute* command
- temp/deform = style name of this compute command

### 18.139.2 Examples

```
compute myTemp all temp/deform
```

### 18.139.3 Description

Define a computation that calculates the temperature of a group of atoms, after subtracting out a streaming velocity induced by the simulation box changing size and/or shape, for example in a non-equilibrium MD (NEMD) simulation. The size/shape change is induced by use of the *fix deform* command. A compute of this style is created by the *fix nvt/sllod* command to compute the thermal temperature of atoms for thermostating purposes. A compute of this style can also be used by any command that computes a temperature (e.g., *thermo\_modify*, *fix temp/rescale*, *fix npt*).

The deformation *fix* changes the box size and/or shape over time, so each atom in the simulation box can be thought of as having a “streaming” velocity. For example, if the box is being sheared in *x*, relative to *y*, then atoms at the bottom of the box (low *y*) have a small *x* velocity, while atoms at the top of the box (high *y*) have a large *x* velocity. This position-dependent streaming velocity is subtracted from each atom’s actual velocity to yield a thermal velocity, which is then used to compute the temperature.

---

**Note:** *Fix deform* has an option for remapping either atom coordinates or velocities to the changing simulation box. When using this compute in conjunction with a deforming box, *fix deform* should NOT remap atom positions, but rather should let atoms respond to the changing box by adjusting their own velocities (or let *fix deform* remap the atom velocities; see its *remap* option). If *fix deform* does remap atom positions, then they appear to move with the box but their velocity is not changed, and thus they do NOT have the streaming velocity assumed by this compute. LAMMPS will warn you if *fix deform* is defined and its *remap* setting is not consistent with this compute.

---

After the streaming velocity has been subtracted from each atom, the temperature is calculated by the formula

$$KE = \frac{\text{dim}}{2} N k_B T,$$



where KE is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ , dim = 2 or 3 is the dimensionality of the simulation,  $N$  is the number of atoms in the group,  $k_B$  is the Boltzmann constant, and  $T$  is the temperature. Note that  $v$  in the kinetic energy formula is the atom's velocity.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the `compute_modify` command if this is not the case.

The removal of the box deformation velocity component by this fix is essentially computing the temperature after a “bias” has been removed from the velocity of the atoms. If this compute is used with a fix command that performs thermostating then this bias will be subtracted from each atom, thermostating of the remaining thermal velocity will be performed, and the bias will be added back in. Thermostating fixes that work in this way include `fix nvt`, `fix temp/rescale`, `fix temp/berendsen`, and `fix langevin`.

---

**Note:** The temperature calculated by this compute is only accurate if the atoms are indeed moving with a stream velocity profile that matches the box deformation. If not, then the compute will subtract off an incorrect stream velocity, yielding a bogus thermal temperature. You should **not** assume that your atoms are streaming at the same rate the box is deforming. Rather, you should monitor their velocity profiles (e.g., via the `fix ave/chunk` command). You can also compare the results of this compute to `compute temp/profile`, which actually calculates the stream profile before subtracting it. If the two computes do not give roughly the same temperature, then your atoms are not streaming consistent with the box deformation. See the `fix deform` command for more details on ways to get atoms to stream consistently with the box deformation.

---

This compute subtracts out degrees-of-freedom due to fixes that constrain molecular motion, such as `fix shake` and `fix rigid`. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the *extra* option of the `compute_modify` command.

See the [Howto thermostat](#) page for a discussion of different ways to compute temperature and perform thermostating.

### 18.139.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.139.5 Restrictions

none

## 18.139.6 Related commands

*compute temp/ramp, compute temp/profile, fix deform, fix nvt/sllod*

## 18.139.7 Default

none

# 18.140 compute temp/deform/eff command

## 18.140.1 Syntax

```
compute ID group-ID temp/deform/eff
```

- ID, group-ID are documented in *compute* command
- temp/deform/eff = style name of this compute command

## 18.140.2 Examples

```
compute myTemp all temp/deform/eff
```

## 18.140.3 Description

Define a computation that calculates the temperature of a group of nuclei and electrons in the *electron force field* model, after subtracting out a streaming velocity induced by the simulation box changing size and/or shape, for example in a non-equilibrium MD (NEMD) simulation. The size/shape change is induced by use of the *fix deform* command. A compute of this style is created by the *fix nvt/sllod/eff* command to compute the thermal temperature of atoms for thermostating purposes. A compute of this style can also be used by any command that computes a temperature (e.g., *thermo\_modify, fix npt/eff*).

The calculation performed by this compute is exactly like that described by the *compute temp/deform* command, except that the formula for the temperature includes the radial electron velocity contributions, as discussed by the *compute temp/eff* command. Note that only the translational degrees of freedom for each nuclei or electron are affected by the streaming velocity adjustment. The radial velocity component of the electrons is not affected.

## 18.140.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.140.5 Restrictions

This compute is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 18.140.6 Related commands

*compute temp/ramp, fix deform, fix nvt/sllod/eff*

### 18.140.7 Default

none

## 18.141 compute temp/drude command

### 18.141.1 Syntax

```
compute ID group-ID temp/drude
```

- ID, group-ID are documented in *compute* command
- temp/drude = style name of this compute command

### 18.141.2 Examples

```
compute TDRUDE all temp/drude
```

Example input scripts available: `examples/PACKAGES/drude`.

### 18.141.3 Description

Define a computation that calculates the temperatures of core–Drude pairs. This compute is designed to be used with the *thermalized Drude oscillator model*. Polarizable models in LAMMPS are described on the *Howto polarizable* doc page.

Drude oscillators consist of a core particle and a Drude particle connected by a harmonic bond, and the relative motion of these Drude oscillators is usually maintained cold by a specific thermostat that acts on the relative motion of the core–Drude particle pairs. Therefore, because LAMMPS considers Drude particles as normal atoms in its default temperature compute (*compute temp* command), the reduced temperature of the core–Drude particle pairs is not calculated correctly.

By contrast, this compute calculates the temperature of the cores using center-of-mass velocities of the core–Drude pairs, and the reduced temperature of the Drude particles using the relative velocities of the Drude particles with respect to their cores. Non-polarizable atoms are considered as cores. Their velocities contribute to the temperature of the cores.

### 18.141.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6, which can be accessed by indices 1–6, whose components are

1. temperature of the centers of mass (temperature units)
2. temperature of the dipoles (temperature units)
3. number of degrees of freedom of the centers of mass
4. number of degrees of freedom of the dipoles
5. kinetic energy of the centers of mass (energy units)
6. kinetic energy of the dipoles (energy units)

These values can be used by any command that uses global scalar or vector values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

Both the scalar value and the first two values of the vector calculated by this compute are “intensive”. The other four vector values are “extensive”.

### 18.141.5 Restrictions

The number of degrees of freedom contributing to the temperature is assumed to be constant for the duration of the run unless the *fix\_modify* command sets the option *dynamic yes*.

### 18.141.6 Related commands

*fix drude*, *fix langevin/drude*, *fix drude/transform*, *pair\_style thole*, *compute temp*

### 18.141.7 Default

none

## 18.142 compute temp/eff command

### 18.142.1 Syntax

```
compute ID group-ID temp/eff
```

- ID, group-ID are documented in *compute* command
- temp/eff = style name of this compute command

## 18.142.2 Examples

```
compute 1 all temp/eff
compute myTemp mobile temp/eff
```

## 18.142.3 Description

Define a computation that calculates the temperature of a group of nuclei and electrons in the *electron force field* model. A compute of this style can be used by commands that compute a temperature (e.g., *thermo\_modify*, *fix npt/eff*).

The temperature is calculated by the formula

$$KE = \frac{\text{dim}}{2} N k_B T,$$

where KE is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$  for nuclei and sum of  $\frac{1}{2}(mv^2 + \frac{3}{4}ms^2)$  for electrons, where  $s$  includes the radial electron velocity contributions),  $\text{dim} = 2$  or  $3$  is the dimensionality of the simulation,  $N$  is the number of atoms (only total number of nuclei in the eFF (see the *pair\_eff* command) in the group,  $k_B$  is the Boltzmann constant, and  $T$  is the absolute temperature. This expression is summed over all nuclear and electronic degrees of freedom, essentially by setting the kinetic contribution to the heat capacity to  $\frac{3}{2}k$  (where only nuclei contribute). This subtlety is valid for temperatures well below the Fermi temperature, which for densities two to five times the density of liquid hydrogen ranges from 86,000 to 170,000 K.

**Note:** For eFF models, in order to override the default temperature reported by LAMMPS in the thermodynamic quantities reported via the *thermo* command, the user should apply a *thermo\_modify* command, as shown in the following example:

```
compute          effTemp all temp/eff
thermo_style     custom step etotal pe ke temp press
thermo_modify    temp effTemp
```

A six-component kinetic energy tensor is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, etc. For the eFF, again, the radial electronic velocities are also considered.

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the *compute\_modify* command if this is not the case.

This compute subtracts out degrees-of-freedom due to fixes that constrain molecular motion, such as *fix shake* and *fix rigid*. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the *extra* option of the *compute\_modify* command.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostatting.

## 18.142.4 Output info

The scalar value calculated by this compute is “intensive”, meaning it is independent of the number of atoms in the simulation. The vector values are “extensive”, meaning they scale with the number of atoms in the simulation.

## 18.142.5 Restrictions

This compute is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 18.142.6 Related commands

*compute temp/partial, compute temp/region, compute pressure*

## 18.142.7 Default

none

# 18.143 compute temp/partial command

## 18.143.1 Syntax

```
compute ID group-ID temp/partial xflag yflag zflag
```

- ID, group-ID are documented in *compute* command
- temp/partial = style name of this compute command
- xflag,yflag,zflag = 0/1 for whether to exclude/include this dimension

## 18.143.2 Examples

```
compute newT flow temp/partial 1 1 0
```

## 18.143.3 Description

Define a computation that calculates the temperature of a group of atoms, after excluding one or more velocity components. A compute of this style can be used by any command that computes a temperature (e.g. *thermo\_modify*, *fix temp/rescale*, *fix npt*).

The temperature is calculated by the formula

$$KE = \frac{\text{dim}}{2} N k_B T,$$

where KE is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ ), dim = 2 or 3 is the dimensionality of the simulation,  $N$  is the number of atoms in the group,  $k_B$  is the Boltzmann constant, and  $T$  = temperature. The calculation of KE excludes the  $x$ ,  $y$ , or  $z$  dimensions if *xflag*, *yflag*, or *zflag* is 0. The dim parameter is adjusted to give the correct number of degrees of freedom.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the calculation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the `compute_modify` command if this is not the case.

The removal of velocity components by this fix is essentially computing the temperature after a “bias” has been removed from the velocity of the atoms. If this compute is used with a fix command that performs thermostating then this bias will be subtracted from each atom, thermostating of the remaining thermal velocity will be performed, and the bias will be added back in. Thermostating fixes that work in this way include `fix nvt`, `fix temp/rescale`, `fix temp/berendsen`, and `fix langevin`.

This compute subtracts out degrees-of-freedom due to fixes that constrain molecular motion, such as `fix shake` and `fix rigid`. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the *extra* option of the `compute_modify` command.

See the [Howto thermostat](#) page for a discussion of different ways to compute temperature and perform thermostating.

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 18.143.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

#### 18.143.5 Restrictions

none

### 18.143.6 Related commands

*compute temp, compute temp/region, compute pressure*

### 18.143.7 Default

none

## 18.144 compute temp/profile command

### 18.144.1 Syntax

```
compute ID group-ID temp/profile xflag yflag zflag binstyle args
```

- ID, group-ID are documented in *compute* command
- temp/profile = style name of this compute command
- xflag,yflag,zflag = 0/1 for whether to exclude/include this dimension
- binstyle = *x* or *y* or *z* or *xy* or *yz* or *xz* or *xyz*
  - x* arg = Nx
  - y* arg = Ny
  - z* arg = Nz
  - xy* args = Nx Ny
  - yz* args = Ny Nz
  - xz* args = Nx Nz
  - xyz* args = Nx Ny Nz

Nx, Ny, Nz = number of velocity bins in *x*, *y*, *z* dimensions
- zero or more keyword/value pairs may be appended
- keyword = *out*
  - out* value = *tensor* or *bin*

### 18.144.2 Examples

```
compute myTemp flow temp/profile 1 1 1 x 10  
compute myTemp flow temp/profile 1 1 1 x 10 out bin  
compute myTemp flow temp/profile 0 1 1 xyz 20 20 20
```



### 18.144.3 Description

Define a computation that calculates the temperature of a group of atoms, after subtracting out a spatially-averaged center-of-mass velocity field, before computing the kinetic energy. This can be useful for thermostating a collection of atoms undergoing a complex flow (e.g. via a profile-unbiased thermostat (PUT) as described in (Evans)). A compute of this style can be used by any command that computes a temperature (e.g. *thermo\_modify*, *fix temp/rescale*, *fix npt*).

The *xflag*, *yflag*, *zflag* settings determine which components of average velocity are subtracted out.

The *binstyle* setting and its *Nx*, *Ny*, *Nz* arguments determine how bins are setup to perform spatial averaging. “Bins” can be 1d slabs, 2d pencils, or 3d bricks depending on which *binstyle* is used. The simulation box is partitioned conceptually into  $N_x \times N_y \times N_z$  bins. Depending on the *binstyle*, you may only specify one or two of these values; the others are effectively set to 1 (no binning in that dimension). For non-orthogonal (triclinic) simulation boxes, the bins are “tilted” slabs or pencils or bricks that are parallel to the tilted faces of the box. See the *region prism* command for a discussion of the geometry of tilted boxes in LAMMPS.

When a temperature is computed, the center-of-mass velocity for the set of atoms that are both in the compute group and in the same spatial bin is calculated. This bias velocity is then subtracted from the velocities of individual atoms in the bin to yield a thermal velocity for each atom. Note that if there is only one atom in the bin, its thermal velocity will thus be 0.0.

After the spatially-averaged velocity field has been subtracted from each atom, the temperature is calculated by the formula

$$\text{KE} = \left( \frac{\text{dim}}{N} - N_s N_x N_y N_z - \text{extra} \right) \frac{k_B T}{2},$$

where KE is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ ; dim = 2 or 3 is the dimensionality of the simulation;  $N_s = 0, 1, 2, \text{ or } 3$  for streaming velocity subtracted in 0, 1, 2, or 3 dimensions, respectively; *extra* is the number of extra degrees of freedom; *N* is the number of atoms in the group;  $k_B$  is the Boltzmann constant, and *T* is the absolute temperature. The  $N_s N_x N_y N_z$  term is the number of degrees of freedom subtracted to adjust for the removal of the center-of-mass velocity in each direction of the  $N_x * N_y * N_z$  bins, as discussed in the (Evans) paper. The *extra* term defaults to  $\text{dim} - N_s$  and accounts for overall conservation of center-of-mass velocity across the group in directions where streaming velocity is *not* subtracted. This can be altered using the *extra* option of the *compute\_modify* command.

If the *out* keyword is used with a *tensor* value, which is the default, a kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the *xy* component, and so on. The six components of the vector are ordered *xx*, *yy*, *zz*, *xy*, *xz*, *yz*.

If the *out* keyword is used with a *bin* value, the count of atoms and computed temperature for each bin are stored for output, as an array of values, as described below. The temperature of each bin is calculated as described above, where the bias velocity is subtracted and only the remaining thermal velocity of atoms in the bin contributes to the temperature. See the note below for how the temperature is normalized by the degrees-of-freedom of atoms in the bin.

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the *compute\_modify* command if this is not the case.

The removal of the spatially-averaged velocity field by this fix is essentially computing the temperature after a “bias” has been removed from the velocity of the atoms. If this compute is used with a fix command that performs thermostating then this bias will be subtracted from each atom, thermostating of the remaining thermal velocity will be performed, and the bias will be added back in. Thermostating fixes that work in this way include *fix nvt*, *fix temp/rescale*, *fix temp/berendsen*, and *fix langevin*.

This compute subtracts out degrees-of-freedom due to fixes that constrain molecular motion, such as *fix shake* and *fix rigid*. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the *extra* option of the *compute\_modify* command.

---

**Note:** When using the *out* keyword with a value of *bin*, the calculated temperature for each bin includes the degrees-of-freedom adjustment described in the preceding paragraph for fixes that constrain molecular motion, as well as the adjustment due to the *extra* option (which defaults to *dim - Ns* as described above), by fractionally applying them based on the fraction of atoms in each bin. As a result, the bin degrees-of-freedom summed over all bins exactly equals the degrees-of-freedom used in the scalar temperature calculation,  $\sum N_{\text{DOF}_i} = N_{\text{DOF}}$  and the corresponding relation for temperature is also satisfied ( $\sum N_{\text{DOF}_i} T_i = N_{\text{DOF}} T$ ). These relations will break down in cases for which the adjustment exceeds the actual number of degrees of freedom in a bin. This could happen if a bin is empty or in situations in which rigid molecules are non-uniformly distributed, in which case the reported temperature within a bin may not be accurate.

---

See the [Howto thermostat](#) page for a discussion of different ways to compute temperature and perform thermostating. Using this compute in conjunction with a thermostating fix, as explained there, will effectively implement a profile-unbiased thermostat (PUT), as described in ([Evans](#)).

### 18.144.4 Output info

This compute calculates a global scalar (the temperature). Depending on the setting of the *out* keyword, it also calculates a global vector or array. For *out = tensor*, it calculates a vector of length 6 (KE tensor), which can be accessed by indices 1–6. For *out = bin* it calculates a global array which has 2 columns and *N* rows, where *N* is the number of bins. The first column contains the number of atoms in that bin. The second contains the temperature of that bin, calculated as described above. The ordering of rows in the array is as follows. Bins in *x* vary fastest, then *y*, then *z*. Thus for a  $10 \times 10 \times 10$  3d array of bins, there will be 1000 rows. The bin with indices  $(i_x, i_y, i_z) = (2, 3, 4)$  would map to row  $M = 10^2(i_z - 1) + 10(i_y - 1) + i_x = 322$ , where the rows are numbered from 1 to 1000 and the bin indices are numbered from 1 to 10 in each dimension.

These values can be used by any command that uses global scalar or vector or array values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”. The array values are “intensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*. The first column of array values are counts; the values in the second column will be in temperature *units*.

### 18.144.5 Restrictions

You should not use too large a velocity-binning grid, especially in 3d. In the current implementation, the binned velocity averages are summed across all processors, so this will be inefficient if the grid is too large, and the operation is performed every timestep, as it will be for most thermostats.

### 18.144.6 Related commands

*compute temp*, *compute temp/ramp*, *compute temp/deform*, *compute pressure*

## 18.144.7 Default

The option default is out = tensor.

(Evans) Evans and Morriss, Phys Rev Lett, 56, 2172-2175 (1986).

## 18.145 compute temp/ramp command

### 18.145.1 Syntax

```
compute ID group-ID temp/ramp vdim vlo vhi dim clo chi keyword value ...
```

- ID, group-ID are documented in *compute* command
- temp/ramp = style name of this compute command
- vdim = vx or vy or vz
- vlo,vhi = subtract velocities between vlo and vhi (velocity units)
- dim = x or y or z
- clo,chi = lower and upper bound of domain to subtract from (distance units)
- zero or more keyword/value pairs may be appended
- keyword = *units*

*units* value = *lattice* or *box*

### 18.145.2 Examples

```
compute 2nd middle temp/ramp vx 0 8 y 2 12 units lattice
```

### 18.145.3 Description

Define a computation that calculates the temperature of a group of atoms, after subtracting out an ramped velocity profile before computing the kinetic energy. A compute of this style can be used by any command that computes a temperature (e.g. *thermo\_modify*, *fix temp/rescale*, *fix npt*).

The meaning of the arguments for this command which define the velocity ramp are the same as for the *velocity ramp* command which was presumably used to impose the velocity.

After the ramp velocity has been subtracted from the specified dimension for each atom, the temperature is calculated by the formula

$$\text{KE} = \frac{\text{dim}}{2} N k_B T,$$

where KE is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ ), dim = 2 or 3 is the dimensionality of the simulation,  $N$  is the number of atoms in the group,  $k_B$  is the Boltzmann constant, and  $T$  is the absolute temperature.

The *units* keyword determines the meaning of the distance units used for coordinates (*clo*, *chi*) and velocities (*vlo*, *vhi*). A *box* value selects standard distance units as defined by the *units* command (e.g., Å for units = real or metal). A

*lattice* value means the distance units are in lattice spacings (i.e., velocity in lattice spacings per unit time). The *lattice* command must have been previously used to define the lattice spacing.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the *compute\_modify* command if this is not the case.

The removal of the ramped velocity component by this fix is essentially computing the temperature after a “bias” has been removed from the velocity of the atoms. If this compute is used with a fix command that performs thermostating then this bias will be subtracted from each atom, thermostating of the remaining thermal velocity will be performed, and the bias will be added back in. Thermostating fixes that work in this way include *fix nvt*, *fix temp/rescale*, *fix temp/berendsen*, and *fix langevin*.

This compute subtracts out degrees-of-freedom due to fixes that constrain molecular motion, such as *fix shake* and *fix rigid*. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the *extra* option of the *compute\_modify* command.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostating.

### 18.145.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.145.5 Restrictions

none

### 18.145.6 Related commands

*compute temp*, *compute temp/profile*, *compute temp/deform*, *compute pressure*

### 18.145.7 Default

The option default is `units = lattice`.

## 18.146 compute temp/region command

### 18.146.1 Syntax

```
compute ID group-ID temp/region region-ID
```

- ID, group-ID are documented in *compute* command
- temp/region = style name of this compute command
- region-ID = ID of region to use for choosing atoms

### 18.146.2 Examples

```
compute mine flow temp/region boundary
```

### 18.146.3 Description

Define a computation that calculates the temperature of a group of atoms in a geometric region. This can be useful for thermostating one portion of the simulation box. For example, a McDLT simulation where one side is cooled, and the other side is heated. A compute of this style can be used by any command that computes a temperature (e.g., *thermo\_modify*, *fix temp/rescale*).

Note that a *region*-style temperature can be used to thermostat with *fix temp/rescale* or *fix langevin*, but should probably not be used with Nose–Hoover style fixes (*fix nvt*, *fix npt*, or *fix nph*) if the degrees of freedom included in the computed temperature vary with time.

The temperature is calculated by the formula

$$\text{KE} = \frac{\text{dim}}{2} N k_B T,$$

where KE = is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ ), dim = 2 or 3 is the dimensionality of the simulation,  $N$  is the number of atoms in both the group and region,  $k_B$  is the Boltzmann constant, and  $T$  temperature.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ .

The number of atoms contributing to the temperature is calculated each time the temperature is evaluated since it is assumed atoms can enter/leave the region. Thus there is no need to use the *dynamic* option of the *compute\_modify* command for this compute style.

The removal of atoms outside the region by this fix is essentially computing the temperature after a “bias” has been removed, which in this case is the velocity of any atoms outside the region. If this compute is used with a fix command that performs thermostating then this bias will be subtracted from each atom, thermostating of the remaining thermal velocity will be performed, and the bias will be added back in. Thermostating fixes that work in this way include *fix nvt*, *fix temp/rescale*, *fix temp/berendsen*, and *fix langevin*. This means that when this compute is used to calculate the temperature for any of the thermostating fixes via the *fix modify temp* command, the thermostat will operate only on atoms that are currently in the geometric region.

Unlike other compute styles that calculate temperature, this compute does not subtract out degrees-of-freedom due to fixes that constrain motion, such as *fix shake* and *fix rigid*. This is because those degrees of freedom (e.g., a constrained bond) could apply to sets of atoms that straddle the region boundary, and hence the concept is somewhat ill-defined. If

needed the number of subtracted degrees of freedom can be set explicitly using the *extra* option of the *compute\_modify* command.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostatting.

### 18.146.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.146.5 Restrictions

none

### 18.146.6 Related commands

*compute temp*, *compute pressure*

### 18.146.7 Default

none

## 18.147 compute temp/region/eff command

### 18.147.1 Syntax

```
compute ID group-ID temp/region/eff region-ID
```

- ID, group-ID are documented in *compute* command
- temp/region/eff = style name of this compute command
- region-ID = ID of region to use for choosing atoms

### 18.147.2 Examples

```
compute mine flow temp/region/eff boundary
```

### 18.147.3 Description

Define a computation that calculates the temperature of a group of nuclei and electrons in the *electron force field* model, within a geometric region using the electron force field. A compute of this style can be used by commands that compute a temperature (e.g., *thermo\_modify*).

The operation of this compute is exactly like that described by the *compute temp/region* command, except that the formula for the temperature itself includes the radial electron velocity contributions, as discussed by the *compute temp/eff* command.

### 18.147.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.147.5 Restrictions

This compute is part of the EFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 18.147.6 Related commands

*compute temp/region*, *compute temp/eff*, *compute pressure*

### 18.147.7 Default

none

## 18.148 compute temp/rotate command

### 18.148.1 Syntax

```
compute ID group-ID temp/rotate
```

- ID, group-ID are documented in *compute* command
- temp/rotate = style name of this compute command

## 18.148.2 Examples

```
compute Tbead bead temp/rotate
```

## 18.148.3 Description

Define a computation that calculates the temperature of a group of atoms, after subtracting out the center-of-mass velocity and angular velocity of the group. This is useful if the group is expected to have a non-zero net velocity and/or global rotation motion for some reason. A compute of this style can be used by any command that computes a temperature (e.g., *thermo\_modify*, *fix temp/rescale*, *fix nvt*).

After the center-of-mass velocity and angular velocity has been subtracted from each atom, the temperature is calculated by the formula

$$\text{KE} = \frac{\text{dim}}{2} N k_B T,$$

where KE is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ ), dim = 2 or 3 is the dimensionality of the simulation,  $N$  is the number of atoms in the group,  $k_B$  is the Boltzmann constant, and  $T$  is the absolute temperature.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the *compute\_modify* command if this is not the case.

The removal of the center-of-mass velocity and angular velocity by this fix is essentially computing the temperature after a “bias” has been removed from the velocity of the atoms. If this compute is used with a fix command that performs thermostating then this bias will be subtracted from each atom, thermostating of the remaining thermal velocity will be performed, and the bias will be added back in. Thermostating fixes that work in this way include *fix nvt*, *fix temp/rescale*, *fix temp/berendsen*, and *fix langevin*.

This compute subtracts out degrees-of-freedom due to fixes that constrain molecular motion, such as *fix shake* and *fix rigid*. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees-of-freedom can be altered using the *extra* option of the *compute\_modify* command.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostating.

## 18.148.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1-6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.



## 18.148.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 18.148.6 Related commands

*compute temp*

## 18.148.7 Default

none

## 18.149 compute temp/sphere command

### 18.149.1 Syntax

```
compute ID group-ID temp/sphere keyword value ...
```

- ID, group-ID are documented in *compute* command
- temp/sphere = style name of this compute command
- zero or more keyword/value pairs may be appended
- keyword = *bias* or *dof*
  - bias* value = bias-ID
    - bias-ID = ID of a temperature compute that removes a velocity bias
  - dof* value = *all* or *rotate*
    - all* = compute temperature of translational and rotational degrees of freedom
    - rotate* = compute temperature of just rotational degrees of freedom

### 18.149.2 Examples

```
compute 1 all temp/sphere
compute myTemp mobile temp/sphere bias tempCOM
compute myTemp mobile temp/sphere dof rotate
```

### 18.149.3 Description

Define a computation that calculates the temperature of a group of spherical particles, including a contribution from both their translational and rotational kinetic energy. This differs from the usual *compute temp* command, which assumes point particles with only translational kinetic energy.

Both point and finite-size particles can be included in the group. Point particles do not rotate, so they have only three translational degrees of freedom. For 3d spherical particles, each has six degrees of freedom (three translational, three rotational). For 2d spherical particles, each has three degrees of freedom (two translational, one rotational).

**Note:** This choice for degrees of freedom (DOF) assumes that all finite-size spherical particles in your model will freely rotate, sampling all their rotational DOF. It is possible to use a combination of interaction potentials and fixes that induce no torque or otherwise constrain some of all of your particles so that this is not the case. Then there are less DOF and you should use the `compute_modify extra` command to adjust the DOF accordingly.

---

The translational kinetic energy is computed the same as is described by the `compute temp` command. The rotational kinetic energy is computed as  $\frac{1}{2}I\omega^2$ , where  $I$  is the moment of inertia for a sphere and  $\omega$  is the particle's angular velocity.

---

**Note:** For *2d models*, particles are treated as spheres, not disks, meaning their moment of inertia will be the same as in 3d.

---

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute. The formula for the components of the tensor is the same as the above formulas, except that  $v^2$  and  $\omega^2$  are replaced by  $v_x v_y$  and  $\omega_x \omega_y$  for the  $xy$  component. The six components of the vector are ordered  $xx, yy, zz, xy, xz, yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the `dynamic` option of the `compute_modify` command if this is not the case.

This compute subtracts out translational degrees-of-freedom due to fixes that constrain molecular motion, such as `fix shake` and `fix rigid`. This means the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees of freedom can be altered using the `extra` option of the `compute_modify` command.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostatting.

---

The keyword/value option pairs are used in the following ways.

For the `bias` keyword, `bias-ID` refers to the ID of a temperature compute that removes a “bias” velocity from each atom. This allows compute temp/sphere to compute its thermal temperature after the translational kinetic energy components have been altered in a prescribed way (e.g., to remove a flow velocity profile). Thermostats that use this compute will work with this bias term. See the doc pages for individual computes that calculate a temperature and the doc pages for fixes that perform thermostatting for more details.

For the `dof` keyword, a setting of `all` calculates a temperature that includes both translational and rotational degrees of freedom. A setting of `rotate` calculates a temperature that includes only rotational degrees of freedom.

---

#### 18.149.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 6 (KE tensor), which can be accessed by indices 1–6. These values can be used by any command that uses global scalar or vector values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The vector values are “extensive”.

The scalar value will be in temperature *units*. The vector values will be in energy *units*.

### 18.149.5 Restrictions

This fix requires that atoms store torque and angular velocity (omega) and a radius as defined by the *atom\_style sphere* command.

All particles in the group must be finite-size spheres, or point particles with radius = 0.0.

### 18.149.6 Related commands

*compute temp*, *compute temp/asphere*

### 18.149.7 Default

The option defaults are no bias and dof = all.

## 18.150 compute temp/uef command

### 18.150.1 Syntax

```
compute ID group-ID temp/uef
```

- ID, group-ID are documented in *compute* command
- temp/uef = style name of this compute command

### 18.150.2 Examples

```
compute 1 all temp/uef
compute 2 sel temp/uef
```

### 18.150.3 Description

This command is used to compute the kinetic energy tensor in the reference frame of the applied flow field when *fix nvt/uef* or *fix npt/uef* is used. It is not necessary to use this command to compute the scalar value of the temperature. A *compute temp* may be used for that purpose.

Output information for this command can be found in the documentation for *compute temp*.

### 18.150.4 Restrictions

This fix is part of the UEF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This command can only be used when *fix nvt/uef* or *fix npt/uef* is active.

## 18.150.5 Related commands

*compute temp*, *fix nvt/uef*, *compute pressure/uef*

## 18.150.6 Default

none

# 18.151 compute ti command

## 18.151.1 Syntax

```
compute ID group ti keyword args ...
```

- ID, group-ID are documented in *compute* command
- ti = style name of this compute command
- one or more attribute/arg pairs may be appended
- keyword = pair style (lj/cut, gauss, born, etc.) or *tail* or *kspace*

pair style args = atype v\_name1 v\_name2

atype = atom type (see asterisk form below)

v\_name1 = variable with name1 that is energy scale factor and function of lambda

v\_name2 = variable with name2 that is derivative of v\_name1 with respect to lambda

tail args = atype v\_name1 v\_name2

atype = atom type (see asterisk form below)

v\_name1 = variable with name1 that is energy tail correction scale factor and

↪function of lambda

v\_name2 = variable with name2 that is derivative of v\_name1 with respect to lambda

kpace args = atype v\_name1 v\_name2

atype = atom type (see asterisk form below)

v\_name1 = variable with name1 that is K-Space scale factor and function of lambda

v\_name2 = variable with name2 that is derivative of v\_name1 with respect to lambda

## 18.151.2 Examples

```
compute 1 all ti lj/cut 1 v_lj v_dlj coul/long 2 v_c v_dc kspace 1 v_ks v_dks
compute 1 all ti lj/cut 1*3 v_lj v_dlj coul/long * v_c v_dc kspace * v_ks v_dks
```

### 18.151.3 Description

Define a computation that calculates the derivative of the interaction potential with respect to *lambda*, the coupling parameter used in a thermodynamic integration. This derivative can be used to infer a free energy difference resulting from an alchemical simulation, as described in *Eike*.

Typically this compute will be used in conjunction with the *fix adapt* command which can perform alchemical transformations by adjusting the strength of an interaction potential as a simulation runs, as defined by one or more *pair\_style* or *k-space\_style* commands. This scaling is done via a prefactor on the energy, forces, virial calculated by the pair or *k*-space style. The prefactor is often a function of a *lambda* parameter which may be adjusted from 0 to 1 (or vice versa) over the course of a *run*. The time-dependent adjustment is what the *fix adapt* command does.

Assume that the unscaled energy of a *pair\_style* or *k-space\_style* is given by  $U$ . Then the scaled energy is

$$U_s = f(\lambda)U$$

where  $f$  is some function of  $\lambda$ . What this compute calculates is

$$\frac{dU_s}{d\lambda} = U \frac{df(\lambda)}{d\lambda} = \frac{U_s}{f(\lambda)} \frac{df(\lambda)}{d\lambda},$$

which is the derivative of the system's scaled potential energy  $U_s$  with respect to  $\lambda$ .

To perform this calculation, you provide one or more atom types as *atype*. The variable *atype* can be specified in one of two ways. An explicit numeric value can be used, as in the first example above, or a wildcard asterisk can be used in place of or in conjunction with the *atype* argument to select multiple atom types. This takes the form “\*” or “\*n” or “m\*” or “m\*n”. If  $N$  is the number of atom types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $m$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive).

You also specify two functions, as *equal-style variables*. The first is specified as  $v\_name1$ , where *name1* is the name of the variable, and is  $f(\lambda)$  in the notation above. The second is specified as  $v\_name2$ , where *name2* is the name of the variable, and is  $df(\lambda)/d\lambda$  in the notation above (i.e., it is the analytic derivative of  $f$  with respect to  $\lambda$ ). Note that the *name1* variable is also typically given as an argument to the *fix adapt* command.

An alchemical simulation may use several pair potentials together, invoked via the *pair\_style hybrid* or *hybrid/overlay* command. The total  $dU_s/d\lambda$  for the overall system is calculated as the sum of each contributing term as listed by the keywords in the *compute ti* command. Individual pair potentials can be listed, which will be sub-styles in the hybrid case. You can also include a *k*-space term via the *k-space* keyword. You can also include a pairwise long-range tail correction to the energy via the *tail* keyword.

For each term, you can specify a different (or the same) scale factor by the two variables that you list. Again, these will typically correspond to the scale factors applied to these various potentials and the *k*-space contribution via the *fix adapt* command.

More details about the exact functional forms for the computation of  $du/dl$  can be found in the paper by *Eike*.

### 18.151.4 Output info

This compute calculates a global scalar, namely  $dU_s/d\lambda$ . This value can be used by any command that uses a global scalar value from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “extensive”.

The scalar value will be in energy *units*.

### 18.151.5 Restrictions

This compute is part of the EXTRA-COMPUTE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 18.151.6 Related commands

*fix adapt*

### 18.151.7 Default

none

---

(Eike) Eike and Maginn, Journal of Chemical Physics, 124, 164503 (2006).

## 18.152 compute torque/chunk command

### 18.152.1 Syntax

```
compute ID group-ID torque/chunk chunkID
```

- ID, group-ID are documented in *compute* command
- torque/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command

### 18.152.2 Examples

```
compute 1 fluid torque/chunk molchunk
```

### 18.152.3 Description

Define a computation that calculates the torque on multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

This compute calculates the three components of the torque vector for eqch chunk, due to the forces on the individual atoms in the chunk around the center-of-mass of the chunk. The calculation includes all effects due to atoms passing through periodic boundaries.

Note that only atoms in the specified group contribute to the calculation. The *compute chunk/atom* command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

---

**Note:** The coordinates of an atom contribute to the chunk’s torque in “unwrapped” form, by using the image flags associated with each atom. See the *dump custom* command for a discussion of “unwrapped” coordinates. See the Atoms section of the *read\_data* command for a discussion of image flags and how they are set for each atom. You can reset the image flags (e.g., to 0) before invoking this compute by using the *set image* command.

---

The simplest way to output the results of the compute torque/chunk calculation to a file is to use the *fix ave/time* command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk all torque/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk[*] file tmp.out mode vector
```

### 18.152.4 Output info

This compute calculates a global array where the number of rows is equal to the number of chunks *Nchunk* as calculated by the specified *compute chunk/atom* command. The number of columns is three for the *x*, *y*, and *z* components of the torque for each chunk. These values can be accessed by any command that uses global array values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The array values are “intensive”. The array values will be in force-distance *units*.

### 18.152.5 Restrictions

none

### 18.152.6 Related commands

*variable torque() function*

### 18.152.7 Default

none

## 18.153 compute vacf command

### 18.153.1 Syntax

```
compute ID group-ID vacf
```

- ID, group-ID are documented in *compute* command
- vacf = style name of this compute command

## 18.153.2 Examples

```
compute 1 all vacf
compute 1 upper vacf
```

## 18.153.3 Description

Define a computation that calculates the velocity auto-correlation function (VACF), averaged over a group of atoms. Each atom's contribution to the VACF is its current velocity vector dotted into its initial velocity vector at the time the compute was specified.

A vector of four quantities is calculated by this compute. The first three elements of the vector are  $v_x v_{x,0}$  (and similar for the  $y$  and  $z$  components), summed and averaged over atoms in the group, where  $v_x$  is the current  $x$ -component of the velocity of the atom and  $v_{x,0}$  is the initial  $x$ -component of the velocity of the atom. The fourth element of the vector is the total VACF (i.e.,  $(v_x v_{x,0} + v_y v_{y,0} + v_z v_{z,0})$ ), summed and averaged over atoms in the group.

The integral of the VACF versus time is proportional to the diffusion coefficient of the diffusing atoms. This can be computed in the following manner, using the *variable trap()* function:

```
compute      2 all vacf
fix          5 all vector 1 c_2[4]
variable     diff equal dt*trap(f_5)
thermo_style custom step v_diff
```

---

**Note:** If you want the quantities calculated by this compute to be continuous when running from a *restart file*, then you should use the same ID for this compute, as in the original run. This is so that the fix this compute creates to store per-atom quantities will also have the same ID, and thus be initialized correctly with time=0 atom velocities from the restart file.

---

## 18.153.4 Output info

This compute calculates a global vector of length 4, which can be accessed by indices 1–4 by any command that uses global vector values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

The vector values are “intensive”. The vector values will be in velocity<sup>2</sup> *units*.

## 18.153.5 Restrictions

none



## 18.153.6 Related commands

*compute msd*

## 18.153.7 Default

none

# 18.154 compute vcm/chunk command

## 18.154.1 Syntax

```
compute ID group-ID vcm/chunk chunkID
```

- ID, group-ID are documented in *compute* command
- vcm/chunk = style name of this compute command
- chunkID = ID of *compute chunk/atom* command

## 18.154.2 Examples

```
compute 1 fluid vcm/chunk molchunk
```

## 18.154.3 Description

Define a computation that calculates the center-of-mass velocity for multiple chunks of atoms.

In LAMMPS, chunks are collections of atoms defined by a *compute chunk/atom* command, which assigns each atom to a single chunk (or no chunk). The ID for this command is specified as chunkID. For example, a single chunk could be the atoms in a molecule or atoms in a spatial bin. See the *compute chunk/atom* and *Howto chunk* doc pages for details of how chunks can be defined and examples of how they can be used to measure properties of a system.

This compute calculates the  $(x, y, z)$  components of the center-of-mass velocity for each chunk. This is done by summing mass\*velocity for each atom in the chunk and dividing the sum by the total mass of the chunk.

Note that only atoms in the specified group contribute to the calculation. The *compute chunk/atom* command defines its own group; atoms will have a chunk ID = 0 if they are not in that group, signifying they are not assigned to a chunk, and will thus also not contribute to this calculation. You can specify the “all” group for this command if you simply want to include atoms with non-zero chunk IDs.

The simplest way to output the results of the compute vcm/chunk calculation to a file is to use the *fix ave/time* command, for example:

```
compute cc1 all chunk/atom molecule
compute myChunk all vcm/chunk cc1
fix 1 all ave/time 100 1 100 c_myChunk[*] file tmp.out mode vector
```

### 18.154.4 Output info

This compute calculates a global array where the number of rows is the number of chunks *Nchunk* as calculated by the specified `compute chunk/atom` command. The number of columns is 3 for the  $(x, y, z)$  center-of-mass velocity coordinates of each chunk. These values can be accessed by any command that uses global array values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The array values are “intensive”. The array values will be in velocity *units*.

### 18.154.5 Restrictions

none

### 18.154.6 Related commands

none

### 18.154.7 Default

none

## 18.155 compute viscosity/cos command

### 18.155.1 Syntax

```
compute ID group-ID viscosity/cos
```

- ID, group-ID are documented in `compute` command
- viscosity/cos = style name of this compute command

### 18.155.2 Examples

```
units      real
compute   cos all viscosity/cos
variable  V equal c_cos[7]
variable  A equal 0.02E-5 # A/fs^2
variable  density equal density
variable  lz equal lz
variable  reciprocalViscosity equal v_V/{A}/v_density*39.4784/v_lz/v_lz*100 # 1/(Pa*s)
```

### 18.155.3 Description

Define a computation that calculates the velocity amplitude of a group of atoms with an cosine-shaped velocity profile and the temperature of them after subtracting out the velocity profile before computing the kinetic energy. A compute of this style can be used by any command that computes a temperature (e.g., *thermo\_modify*, *fix npt*).

This command together with *fix accelerate/cos* enables viscosity calculation with periodic perturbation method, as described by *Hess*. An acceleration along the  $x$ -direction is applied to the simulation system by using *fix accelerate/cos* command. The acceleration is a periodic function along the  $z$ -direction:

$$a_x(z) = A \cos\left(\frac{2\pi z}{l_z}\right)$$

where  $A$  is the acceleration amplitude,  $l_z$  is the  $z$ -length of the simulation box. At steady state, the acceleration generates a velocity profile:

$$v_x(z) = V \cos\left(\frac{2\pi z}{l_z}\right)$$

The generated velocity amplitude  $V$  is related to the shear viscosity  $\eta$  by

$$V = \frac{A\rho}{\eta} \left(\frac{l_z}{2\pi}\right)^2,$$

and it can be obtained from ensemble average of the velocity profile via

$$V = \frac{\sum_i 2m_i v_{i,x} \cos\left(\frac{2\pi z_i}{l_z}\right)}{\sum_i m_i}$$

where  $m_i$ ,  $v_{i,x}$  and  $z_i$  are the mass,  $x$ -component velocity, and  $z$ -coordinate of a particle, respectively.

After the cosine-shaped collective velocity in the  $x$ -direction has been subtracted for each atom, the temperature is calculated by the formula

$$\text{KE} = \frac{\text{dim}}{2} N k_B T,$$

where KE is the total kinetic energy of the group of atoms (sum of  $\frac{1}{2}mv^2$ ),  $\text{dim} = 2$  or  $3$  is the dimensionality of the simulation,  $N$  is the number of atoms in the group,  $k_B$  is the Boltzmann constant, and  $T$  is the absolute temperature.

A kinetic energy tensor, stored as a six-element vector, is also calculated by this compute for use in the computation of a pressure tensor. The formula for the components of the tensor is the same as the above formula, except that  $v^2$  is replaced by  $v_x v_y$  for the  $xy$  component, and so on. The six components of the vector are ordered  $xx$ ,  $yy$ ,  $zz$ ,  $xy$ ,  $xz$ ,  $yz$ .

The number of atoms contributing to the temperature is assumed to be constant for the duration of the run; use the *dynamic* option of the *compute\_modify* command if this is not the case. However, in order to get meaningful result, the group ID of this compute should be all.

The removal of the cosine-shaped velocity component by this command is essentially computing the temperature after a “bias” has been removed from the velocity of the atoms. If this compute is used with a *fix* command that performs thermostating then this bias will be subtracted from each atom, thermostating of the remaining thermal velocity will be performed, and the bias will be added back in. Thermostating fixes that work in this way include *fix nvt*, *fix temp/rescale*, *fix temp/berendsen*, and *fix langevin*.

This compute subtracts out degrees of freedom due to fixes that constrain molecular motion, such as *fix shake* and *fix rigid*. This means that the temperature of groups of atoms that include these constraints will be computed correctly. If needed, the subtracted degrees of freedom can be altered using the *extra* option of the *compute\_modify* command.

See the *Howto thermostat* page for a discussion of different ways to compute temperature and perform thermostating.

### 18.155.4 Output info

This compute calculates a global scalar (the temperature) and a global vector of length 7, which can be accessed by indices 1–7. The first six elements of the vector are the KE tensor, and the seventh is the cosine-shaped velocity amplitude  $V$ , which can be used to calculate the reciprocal viscosity, as shown in the example. These values can be used by any command that uses global scalar or vector values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options.

The scalar value calculated by this compute is “intensive”. The first six elements of vector values are “extensive”, and the seventh element of vector values is “intensive”.

The scalar value will be in temperature *units*. The first six elements of vector values will be in energy *units*. The seventh element of vector value will be in velocity *units*.

### 18.155.5 Restrictions

This command is only available when LAMMPS was built with the MISC package. Since this compute depends on *fix accelerate/cos* which can only work for 3d systems, it cannot be used for 2d systems.

### 18.155.6 Related commands

*fix accelerate/cos*

### 18.155.7 Default

none

---

(Hess) Hess, B. The Journal of Chemical Physics 2002, 116 (1), 209-217.

## 18.156 compute voronoi/atom command

### 18.156.1 Syntax

```
compute ID group-ID voronoi/atom keyword arg ...
```

- ID, group-ID are documented in *compute* command
- voronoi/atom = style name of this compute command
- zero or more keyword/value pairs may be appended
- keyword = *only\_group* or *surface* or *radius* or *edge\_histo* or *edge\_threshold* or *face\_threshold* or *neighbors* or *peratom*

*only\_group* = no arg

*occupation* = no arg

*surface* arg = sgroup-ID

sgroup-ID = compute the dividing surface between group-ID and sgroup-ID

this keyword adds a third column to the compute output

*radius* arg = v\_r

v\_r = radius atom style variable for a poly-disperse Voronoi tessellation

```

edge_histo arg = maxedge
    maxedge = maximum number of Voronoi cell edges to be accounted in the histogram
edge_threshold arg = minlength
    minlength = minimum length for an edge to be counted
face_threshold arg = minarea
    minarea = minimum area for a face to be counted
neighbors value = yes or no = store list of all neighbors or no
peratom value = yes or no = per-atom quantities accessible or no

```

## 18.156.2 Examples

```

compute 1 all voronoi/atom
compute 2 precipitate voronoi/atom surface matrix
compute 3b precipitate voronoi/atom radius v_r
compute 4 solute voronoi/atom only_group
compute 5 defects voronoi/atom occupation
compute 6 all voronoi/atom neighbors yes

```

## 18.156.3 Description

Define a computation that calculates the Voronoi tessellation of the atoms in the simulation box. The tessellation is calculated using all atoms in the simulation, but non-zero values are only stored for atoms in the group.

By default two per-atom quantities are calculated by this compute. The first is the volume of the Voronoi cell around each atom. Any point in an atom's Voronoi cell is closer to that atom than any other. The second is the number of faces of the Voronoi cell. This is equal to the number of nearest neighbors of the central atom, plus any exterior faces (see note below). If the *peratom* keyword is set to "no", the per-atom quantities are still calculated, but they are not accessible.

If the *only\_group* keyword is specified the tessellation is performed only with respect to the atoms contained in the compute group. This is equivalent to deleting all atoms not contained in the group prior to evaluating the tessellation.

If the *surface* keyword is specified a third quantity per atom is computed: the Voronoi cell surface of the given atom. *surface* takes a group ID as an argument. If a group other than *all* is specified, only the Voronoi cell facets facing a neighbor atom from the specified group are counted towards the surface area.

In the example above, a precipitate embedded in a matrix, only atoms at the surface of the precipitate will have non-zero surface area, and only the outward facing facets of the Voronoi cells are counted (the hull of the precipitate). The total surface area of the precipitate can be obtained by running a "reduce sum" compute on `c_2[3]`.

If the *radius* keyword is specified with an atom style variable as the argument, a poly-disperse Voronoi tessellation is performed. Examples for radius variables are

```

variable r1 atom (type==1)*0.1+(type==2)*0.4
compute radius all property/atom radius
variable r2 atom c_radius

```

Here `v_r1` specifies a per-type radius of 0.1 units for type 1 atoms and 0.4 units for type 2 atoms, and `v_r2` accesses the radius property present in atom\_style sphere for granular models.

The *edge\_histo* keyword activates the compilation of a histogram of number of edges on the faces of the Voronoi cells in the compute group. The argument *maxedge* of the this keyword is the largest number of edges on a single Voronoi cell face expected to occur in the sample. This keyword adds the generation of a global vector with *maxedge*+1 entries.

The last entry in the vector contains the number of faces with more than *maxedge* edges. Since the polygon with the smallest amount of edges is a triangle, entries 1 and 2 of the vector will always be zero.

The *edge\_threshold* and *face\_threshold* keywords allow the suppression of edges below a given minimum length and faces below a given minimum area. Ultra short edges and ultra small faces can occur as artifacts of the Voronoi tessellation. These keywords will affect the neighbor count and edge histogram outputs.

If the *occupation* keyword is specified the tessellation is only performed for the first invocation of the compute and then stored. For all following invocations of the compute the number of atoms in each Voronoi cell in the stored tessellation is counted. In this mode the compute returns a per-atom array with 2 columns. The first column is the number of atoms currently in the Voronoi volume defined by this atom at the time of the first invocation of the compute (note that the atom may have moved significantly). The second column contains the total number of atoms sharing the Voronoi cell of the stored tessellation at the location of the current atom. Numbers in column one can be any positive integer including zero, while column two values will always be greater than zero. Column one data can be used to locate vacancies (the coordinates are given by the atom coordinates at the time step when the compute was first invoked), while column two data can be used to identify interstitial atoms.

If the *neighbors* value is set to yes, then this compute creates a local array with 3 columns. There is one row for each face of each Voronoi cell. The 3 columns are the atom ID of the atom that owns the cell, the atom ID of the atom in the neighboring cell (or zero if the face is external), and the area of the face. The array can be accessed by any command that uses local values from a compute as input. See the [Howto output](#) page for an overview of LAMMPS output options. More specifically, the array can be accessed by a *dump local* command to write a file containing all the Voronoi neighbors in a system:

```
compute 6 all voronoi/atom neighbors yes
dump d2 all local 1 dump.neighbors index c_6[1] c_6[2] c_6[3]
```

If the *face\_threshold* keyword is used, then only faces with areas greater than the threshold are stored.

---

The Voronoi calculation is performed by the freely available [Voro++ package](#), written by Chris Rycroft at UC Berkeley and LBL, which must be installed on your system when building LAMMPS for use with this compute. See instructions on obtaining and installing the Voro++ software in the [src/VORONOI/README](#) file.

---

**Note:** The calculation of Voronoi volumes is performed by each processor for the atoms it owns, and includes the effect of ghost atoms stored by the processor. This assumes that the Voronoi cells of owned atoms are not affected by atoms beyond the ghost atom cut-off distance. This is usually a good assumption for liquid and solid systems, but may lead to underestimation of Voronoi volumes in low density systems. By default, the set of ghost atoms stored by each processor is determined by the cutoff used for *pair\_style* interactions. The cutoff can be set explicitly via the *comm\_modify cutoff* command. The Voronoi cells for atoms adjacent to empty regions will extend into those regions up to the communication cutoff in *x*, *y*, or *z*. In that situation, an exterior face is created at the cutoff distance normal to the *x*, *y*, or *z* direction. For triclinic systems, the exterior face is parallel to the corresponding reciprocal lattice vector.

---

**Note:** The Voro++ package performs its calculation in 3d. This will still work for a 2d LAMMPS simulation, provided all the atoms have the same *z*-coordinate. The Voronoi cell of each atom will be a columnar polyhedron with constant cross-sectional area along the *z*-direction and two exterior faces at the top and bottom of the simulation box. If the atoms do not all have the same *z*-coordinate, then the columnar cells will be accordingly distorted. The cross-sectional area of each Voronoi cell can be obtained by dividing its volume by the *z* extent of the simulation box. Note that you define the *z* extent of the simulation box for 2d simulations when using the *create\_box* or *read\_data* commands.

---

## 18.156.4 Output info

By default, this compute calculates a per-atom array with two columns. In regular dynamic tessellation mode the first column is the Voronoi volume, the second is the neighbor count, as described above (read above for the output data in case the *occupation* keyword is specified). These values can be accessed by any command that uses per-atom values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options. If the *peratom* keyword is set to “no”, the per-atom array is still created, but it is not accessible.

If the *edge\_histo* keyword is used, then this compute generates a global vector of length *maxedge*+1, containing a histogram of the number of edges per face.

If the *neighbors* value is set to *yes*, then this compute calculates a local array with three columns. There is one row for each face of each Voronoi cell.

---

**Note:** Some LAMMPS commands such as the *compute reduce* command can accept either a per-atom or local quantity. If this compute produces both quantities, the command may access the per-atom quantity, even if you want to access the local quantity. This effect can be eliminated by using the *peratom* keyword to turn off the production of the per-atom quantities. For the default value *yes* both quantities are produced. For the value *no*, only the local array is produced.

---

The Voronoi cell volume will be in distance *units* cubed. The Voronoi face area will be in distance *units* squared.

## 18.156.5 Restrictions

This compute is part of the VORONOI package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

It also requires you have a copy of the Voro++ library built and installed on your system. See instructions on obtaining and installing the Voro++ software in the *src/VORONOI/README* file.

## 18.156.6 Related commands

*dump custom*, *dump local*

## 18.156.7 Default

*neighbors* no, *peratom* yes

# 18.157 compute xrd command

## 18.157.1 Syntax

```
compute ID group-ID xrd lambda type1 type2 ... typeN keyword value ...
```

- ID, group-ID are documented in *compute* command
- xrd = style name of this compute command
- lambda = wavelength of incident radiation (length units)
- type1 type2 ... typeN = chemical symbol of each atom type (see valid options below)
- zero or more keyword/value pairs may be appended

- keyword = *2Theta* or *c* or *LP* or *manual* or *echo*

*2Theta* values = *Min2Theta* *Max2Theta*

*Min2Theta*, *Max2Theta* = minimum and maximum 2 theta range to explore (radians or degrees)

*c* values = *c1* *c2* *c3*

*c1*, *c2*, *c3* = parameters to adjust the spacing of the reciprocal lattice nodes in the *h*, *k*, and *l* directions respectively

*LP* value = switch to apply Lorentz-polarization factor

0/1 = off/on

*manual* = flag to use manual spacing of reciprocal lattice points based on the values of the *c* parameters

*echo* = flag to provide extra output for debugging purposes

## 18.157.2 Examples

```
compute 1 all xrd 1.541838 Al 0 2Theta 0.087 0.87 c 1 1 1 LP 1 echo
compute 2 all xrd 1.541838 Al 0 2Theta 10 100 c 0.05 0.05 0.05 LP 1 manual

fix 1 all ave/histo/weight 1 1 1 0.087 0.87 250 c_1[1] c_1[2] mode vector file Rad2Theta.
↪xrd
fix 2 all ave/histo/weight 1 1 1 10 100 250 c_2[1] c_2[2] mode vector file Deg2Theta.xrd
```

## 18.157.3 Description

Define a computation that calculates X-ray diffraction intensity as described in (Coleman) on a mesh of reciprocal lattice nodes defined by the entire simulation domain (or manually) using a simulated radiation of wavelength  $\lambda$ .

The X-ray diffraction intensity,  $I$ , at each reciprocal lattice point,  $k$ , is computed from the structure factor,  $F$ , using the equations:

$$I = L_p(\theta) \frac{F^* F}{N}$$

$$F(\mathbf{k}) = \sum_{j=1}^N f_j(\theta) \exp(2\pi i \mathbf{k} \cdot \mathbf{r}_j)$$

$$L_p(\theta) = \frac{1 + \cos^2(2\theta)}{\cos(\theta) \sin^2(\theta)}$$

$$\frac{\sin(\theta)}{\lambda} = \frac{\|\mathbf{k}\|}{2}$$

Here,  $\mathbf{k}$  is the location of the reciprocal lattice node,  $r_j$  is the position of each atom,  $f_j$  are atomic scattering factors,  $L_p$  is the Lorentz-polarization factor, and  $\theta$  is the scattering angle of diffraction. The Lorentz-polarization factor can be turned off using the optional *LP* keyword.

Diffraction intensities are calculated on a three-dimensional mesh of reciprocal lattice nodes. The mesh spacing is defined either (a) by the entire simulation domain or (b) manually using selected values as shown in the 2D diagram below.



For a mesh defined by the simulation domain, a rectilinear grid is constructed with spacing  $cA^{-1}$  along each reciprocal lattice axis, where  $A$  is a matrix containing the vectors corresponding to the edges of the simulation cell. If one or two directions has non-periodic boundary conditions, then the spacing in these directions is defined from the average of the



(inversed) box lengths with periodic boundary conditions. Meshes defined by the simulation domain must contain at least one periodic boundary.

If the *manual* flag is included, the mesh of reciprocal lattice nodes will be defined using the *c* values for the spacing along each reciprocal lattice axis. Note that manual mapping of the reciprocal space mesh is good for comparing diffraction results from multiple simulations; however, it can reduce the likelihood that Bragg reflections will be satisfied unless small spacing parameters ( $< 0.05 \text{ \AA}^{-1}$ ) are implemented. Meshes with manual spacing do not require a periodic boundary.

The limits of the reciprocal lattice mesh are determined by range of scattering angles explored. The *2Theta* parameter allows the user to reduce the scattering angle range to only the region of interest which reduces the cost of the computation.

The atomic scattering factor,  $f_j$ , accounts for the reduction in diffraction intensity due to Compton scattering. Compute xrd uses analytical approximations of the atomic scattering factors that vary for each atom type (type1 type2 ... typeN) and angle of diffraction. The analytic approximation is computed using the formula (*Colliex*):

$$f_j \left( \frac{\sin(\theta)}{\lambda} \right) = \sum_{i=1}^4 a_i \exp \left( -b_i \frac{\sin^2(\theta)}{\lambda^2} \right) + c$$

Coefficients parameterized by (*Peng*) are assigned for each atom type designating the chemical symbol and charge of each atom type. Valid chemical symbols for compute xrd are:

H	He1-	He	Li	Li1+
Be	Be2+	B	C	Cval
N	O	O1-	F	F1-
Ne	Na	Na1+	Mg	Mg2+
Al	Al3+	Si	Sival	Si4+
P	S	Cl	Cl1-	Ar
K	Ca	Ca2+	Sc	Sc3+
Ti	Ti2+	Ti3+	Ti4+	V
V2+	V3+	V5+	Cr	Cr2+
Cr3+	Mn	Mn2+	Mn3+	Mn4+
Fe	Fe2+	Fe3+	Co	Co2+
Co	Ni	Ni2+	Ni3+	Cu
Cu1+	Cu2+	Zn	Zn2+	Ga
Ga3+	Ge	Ge4+	As	Se
Br	Br1-	Kr	Rb	Rb1+
Sr	Sr2+	Y	Y3+	Zr
Zr4+	Nb	Nb3+	Nb5+	Mo
Mo3+	Mo5+	Mo6+	Tc	Ru
Ru3+	Ru4+	Rh	Rh3+	Rh4+
Pd	Pd2+	Pd4+	Ag	Ag1+
Ag2+	Cd	Cd2+	In	In3+
Sn	Sn2+	Sn4+	Sb	Sb3+
Sb5+	Te	I	I1-	Xe
Cs	Cs1+	Ba	Ba2+	La
La3+	Ce	Ce3+	Ce4+	Pr
Pr3+	Pr4+	Nd	Nd3+	Pm
Pm3+	Sm	Sm3+	Eu	Eu2+
Eu3+	Gd	Gd3+	Tb	Tb3+
Dy	Dy3+	Ho	Ho3+	Er
Er3+	Tm	Tm3+	Yb	Yb2+

continues on next page

Table 1 – continued from previous page

Yb3+	Lu	Lu3+	Hf	Hf4+
Ta	Ta5+	W	W6+	Re
Os	Os4+	Ir	Ir3+	Ir4+
Pt	Pt2+	Pt4+	Au	Au1+
Au3+	Hg	Hg1+	Hg2+	Tl
Tl1+	Tl3+	Pb	Pb2+	Pb4+
Bi	Bi3+	Bi5+	Po	At
Rn	Fr	Ra	Ra2+	Ac
Ac3+	Th	Th4+	Pa	U
U3+	U4+	U6+	Np	Np3+
Np4+	Np6+	Pu	Pu3+	Pu4+
Pu6+	Am	Cm	Bk	Cf

If the *echo* keyword is specified, compute xrd will provide extra reporting information to the screen.

### 18.157.4 Output info

This compute calculates a global array. The number of rows in the array is the number of reciprocal lattice nodes that are explored which by the mesh. The global array has two columns.

The first column contains the diffraction angle in the units (radians or degrees) provided with the *2Theta* values. The second column contains the computed diffraction intensities as described above.

The array can be accessed by any command that uses global values from a compute as input. See the *Howto output* doc page for an overview of LAMMPS output options.

All array values calculated by this compute are “intensive”.

### 18.157.5 Restrictions

This compute is part of the DIFFRACTION package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The compute\_xrd command does not work for triclinic cells.

### 18.157.6 Related commands

*fix ave/histo*, *compute saed*

### 18.157.7 Default

The option defaults are *2Theta* = 1 179 (degrees), *c* = 1 1 1, *LP* = 1, no manual flag, no echo flag.

---

(Coleman) Coleman, Spearot, Capolungo, MSMSE, 21, 055020 (2013).

(Colliex) Colliex et al. International Tables for Crystallography Volume C: Mathematical and Chemical Tables, 249-429 (2004).

(Peng) Peng, Ren, Dudarev, Whelan, Acta Crystallogr. A, 52, 257-76 (1996).

## PAIR STYLES

### 19.1 pair\_style adp command

Accelerator Variants: *adp/kk*, *adp/omp*

#### 19.1.1 Syntax

```
pair_style adp
```

#### 19.1.2 Examples

```
pair_style adp
pair_coeff * * Ta.adp Ta
pair_coeff * * ../potentials/AlCu.adp Al Al Cu
```

#### 19.1.3 Description

Style *adp* computes pairwise interactions for metals and metal alloys using the angular dependent potential (ADP) of (*Mishin*), which is a generalization of the *embedded atom method (EAM) potential*. The LAMMPS implementation is discussed in (*Singh*). The total energy  $E_i$  of an atom  $I$  is given by

$$E_i = F_\alpha \left( \sum_{j \neq i} \rho_\beta(r_{ij}) \right) + \frac{1}{2} \sum_{j \neq i} \phi_{\alpha\beta}(r_{ij}) + \frac{1}{2} \sum_s (\mu_i^s)^2 + \frac{1}{2} \sum_{s,t} (\lambda_i^{st})^2 - \frac{1}{6} \nu_i^2$$

$$\mu_i^s = \sum_{j \neq i} u_{\alpha\beta}(r_{ij}) r_{ij}^s$$

$$\lambda_i^{st} = \sum_{j \neq i} w_{\alpha\beta}(r_{ij}) r_{ij}^s r_{ij}^t$$

$$\nu_i = \sum_s \lambda_i^{ss}$$

where  $F$  is the embedding energy which is a function of the atomic electron density  $\rho$ ,  $\phi$  is a pair potential interaction,  $\alpha$  and  $\beta$  are the element types of atoms  $I$  and  $J$ , and  $s$  and  $t = 1, 2, 3$  and refer to the cartesian coordinates. The  $\mu$  and  $\lambda$  terms represent the dipole and quadruple distortions of the local atomic environment which extend the original EAM framework by introducing angular forces.

Note that unlike for other potentials, cutoffs for ADP potentials are not set in the `pair_style` or `pair_coeff` command; they are specified in the ADP potential files themselves. Likewise, the ADP potential files list atomic masses; thus you do not need to use the `mass` command to specify them.

**ADP potentials are available from:**

- The NIST WWW site at <http://www.ctcms.nist.gov/potentials>. Note that ADP potentials obtained from NIST must be converted into the extended DYNAMO `setfl` format discussed below.
- The OpenKIM Project at <https://openkim.org/browse/models/by-type> provides ADP potentials that can be used directly in LAMMPS with the `kim command` interface.

---

Only a single `pair_coeff` command is used with the `adp` style which specifies an extended DYNAMO `setfl` file, which contains information for  $M$  elements. These are mapped to LAMMPS atom types by specifying  $N$  additional arguments after the filename in the `pair_coeff` command, where  $N$  is the number of LAMMPS atom types:

- filename
- $N$  element names = mapping of extended `setfl` elements to atom types

See the `pair_coeff` page for alternate ways to specify the path for the potential file.

As an example, the `potentials/AlCu.adp` file, included in the `potentials` directory of the LAMMPS distribution, is an extended `setfl` file which has tabulated ADP values for  $w$  elements and their alloy interactions: Cu and Al. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Al, and the fourth to be Cu, you would use the following `pair_coeff` command:

```
pair_coeff * * AlCu.adp Al Al Al Cu
```

The first 2 arguments must be `* *` so as to span all LAMMPS atom types. The first three Al arguments map LAMMPS atom types 1,2,3 to the Al element in the extended `setfl` file. The final Cu argument maps LAMMPS atom type 4 to the Al element in the extended `setfl` file. Note that there is no requirement that your simulation use all the elements specified by the extended `setfl` file.

If a mapping value is specified as NULL, the mapping is not performed. This can be used when an `adp` potential is used as part of the `hybrid` pair style. The NULL values are placeholders for atom types that will be used with other potentials.

`Adp` files in the `potentials` directory of the LAMMPS distribution have an “.adp” suffix. A DYNAMO `setfl` file extended for ADP is formatted as follows. Basically it is the standard `setfl` format with additional tabulated functions `u` and `w` added to the file after the tabulated pair potentials. See the `pair_eam` command for further details on the `setfl` format.

- lines 1,2,3 = comments (ignored)
- line 4:  $N_{\text{elements}}$  Element1 Element2 ... ElementN
- line 5:  $N_{\rho}$ ,  $d_{\rho}$ ,  $N_r$ ,  $d_r$ , cutoff

Following the 5 header lines are  $N_{\text{elements}}$  sections, one for each element, each with the following format:

- line 1 = atomic number, mass, lattice constant, lattice type (e.g. FCC)
- embedding function  $F(\rho)$  ( $N_{\rho}$  values)
- density function  $\rho(r)$  ( $N_r$  values)

Following the  $N_{\text{elements}}$  sections,  $N_r$  values for each pair potential  $\phi(r)$  array are listed for all  $i, j$  element pairs in the same format as other arrays. Since these interactions are symmetric ( $i, j = j, i$ ) only  $\phi$  arrays with  $i \geq j$  are listed, in the following order:

$$i, j = (1, 1), (2, 1), (2, 2), (3, 1), (3, 2), (3, 3), (4, 1), \dots, (N_{\text{elements}}, N_{\text{elements}}).$$

The tabulated values for each  $\phi$  function are listed as  $r * \phi$  (in units of eV-Angstroms), since they are for atom pairs, the same as for *other EAM files*.

After the  $\phi(r)$  arrays, each of the  $u(r)$  arrays are listed in the same order with the same assumptions of symmetry. Directly following the  $u(r)$ , the  $w(r)$  arrays are listed. Note that  $\phi(r)$  is the only array tabulated with a scaling by  $r$ .

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.1.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, no special mixing rules are needed, since the ADP potential files specify alloy interactions explicitly.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in tabulated potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.1.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package.

#### 19.1.6 Related commands

*pair\_coeff*, *pair\_eam*

#### 19.1.7 Default

none

---

**(Mishin)** Mishin, Mehl, and Papaconstantopoulos, Acta Mater, 53, 4029 (2005).

**(Singh)** Singh and Warner, Acta Mater, 58, 5797-5805 (2010),

## 19.2 pair\_style agni command

Accelerator Variants: *agni/omp*

### 19.2.1 Syntax

```
pair_style agni
```

### 19.2.2 Examples

```
pair_style      agni
pair_coeff      * * Al.agni Al
```

### 19.2.3 Description

Style *agni* style computes the many-body vectorial force components for an atom as

$$F_i^u = \sum_t^{N_t} \alpha_t \cdot \exp \left[ -\frac{(d_{i,t}^u)^2}{2l^2} \right]$$

$$d_{i,t}^u = \|V_i^u(\eta) - V_t^u(\eta)\|$$

$$V_i^u(\eta) = \sum_{j \neq i} \frac{r_{ij}^u}{r_{ij}} \cdot e^{-\left(\frac{r_{ij}}{a}\right)^2} \cdot f_d(r_{ij})$$

$$f_d(r_{ij}) = \frac{1}{2} \left[ \cos \left( \frac{\pi r_{ij}}{R_c} \right) + 1 \right]$$

$u$  labels the individual components, i.e.  $x$ ,  $y$  or  $z$ , and  $V$  is the corresponding atomic fingerprint.  $d$  is the Euclidean distance between any two atomic fingerprints. A total of  $N_t$  reference atomic environments are considered to construct the force field file.  $\alpha_t$  and  $l$  are the weight coefficients and length scale parameter of the non-linear regression model.

The method implements the recently proposed machine learning access to atomic forces as discussed extensively in the following publications - (*Botu1*) and (*Botu2*). The premise of the method is to map the atomic environment numerically into a fingerprint, and use machine learning methods to create a mapping to the vectorial atomic forces.

Only a single `pair_coeff` command is used with the *agni* style which specifies an AGNI potential file containing the parameters of the force field for the needed elements. These are mapped to LAMMPS atom types by specifying  $N$  additional arguments after the filename in the `pair_coeff` command, where  $N$  is the number of LAMMPS atom types:

- filename
- $N$  element names = mapping of AGNI elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the force field file.

An AGNI force field is fully specified by the filename which contains the parameters of the force field, i.e., the reference training environments used to construct the machine learning force field. Example force field and input files are provided in the `examples/PACKAGES/agni` directory.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#).

page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 19.2.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

## 19.2.5 Restrictions

Currently, only elemental systems are implemented. Also, the method only provides access to the forces and not energies or stresses. The lack of potential energy data makes this pair style incompatible with several of the *minimizer algorithms* like *cg* or *sd*. It should work with damped dynamics based minimizers like *fire* or *quickmin*. However, one can access the energy via thermodynamic integration of the forces as discussed in (*Botu3*). This pair style is part of the MISC package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The AGNI force field files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use the AGNI potential with any LAMMPS units, but you would need to create your own AGNI potential file with coefficients listed in the appropriate units if your simulation does not use “metal” units.

## 19.2.6 Related commands

*pair\_coeff*

## 19.2.7 Default

none

---

**(Botu1)** V. Botu and R. Ramprasad, Int. J. Quant. Chem., 115(16), 1074 (2015).

**(Botu2)** V. Botu and R. Ramprasad, Phys. Rev. B, 92(9), 094306 (2015).

**(Botu3)** V. Botu, R. Batra, J. Chapman and R. Ramprasad, <https://arxiv.org/abs/1610.02098> (2016).

## 19.3 pair\_style airebo command

Accelerator Variants: *airebo/intel*, *airebo/omp*

## 19.4 pair\_style airebo/morse command

Accelerator Variants: *airebo/morse/intel*, *airebo/morse/omp*

## 19.5 pair\_style rebo command

Accelerator Variants: *rebo/intel*, *rebo/omp*

### 19.5.1 Syntax

```
pair_style style cutoff LJ_flag TORSION_flag cutoff_min
```

- style = *airebo* or *airebo/morse* or *rebo*
- cutoff = LJ or Morse cutoff ( $\sigma$  scale factor) (AIREBO and AIREBO-M only)
- LJ\_flag = 0/1 to turn off/on the LJ or Morse term (AIREBO and AIREBO-M only, optional)
- TORSION\_flag = 0/1 to turn off/on the torsion term (AIREBO and AIREBO-M only, optional)
- cutoff\_min = Start of the transition region of cutoff ( $\sigma$  scale factor) (AIREBO and AIREBO-M only, optional)

### 19.5.2 Examples

```
pair_style airebo 3.0
pair_style airebo 2.5 1 0
pair_coeff * * ../potentials/CH.airebo H C

pair_style airebo/morse 3.0
pair_coeff * * ../potentials/CH.airebo-m H C

pair_style rebo
pair_coeff * * ../potentials/CH.rebo H C
```

### 19.5.3 Description

The *airebo* pair style computes the Adaptive Intermolecular Reactive Empirical Bond Order (AIREBO) Potential of (*Stuart*) for a system of carbon and/or hydrogen atoms. Note that this is the initial formulation of AIREBO from 2000, not the later formulation.

The *airebo/morse* pair style computes the AIREBO-M potential, which is equivalent to AIREBO, but replaces the LJ term with a Morse potential. The Morse potentials are parameterized by high-quality quantum chemistry (MP2) calculations and do not diverge as quickly as particle density increases. This allows AIREBO-M to retain accuracy to much higher pressures than AIREBO (up to 40 GPa for Polyethylene). Details for this potential and its parameterization are given in (*O'Conner*).



The *rebo* pair style computes the Reactive Empirical Bond Order (REBO) Potential of (Brenner). Note that this is the so-called second generation REBO from 2002, not the original REBO from 1990. As discussed below, second generation REBO is closely related to the initial AIREBO; it is just a subset of the potential energy terms with a few slightly different parameters

The AIREBO potential consists of three terms:

$$E = \frac{1}{2} \sum_i \sum_{j \neq i} \left[ E_{ij}^{\text{REBO}} + E_{ij}^{\text{LJ}} + \sum_{k \neq i, j} \sum_{l \neq i, j, k} E_{kijl}^{\text{TORSION}} \right]$$

By default, all three terms are included. For the *airebo* style, if the first two optional flag arguments to the `pair_style` command are included, the LJ and torsional terms can be turned off. Note that both or neither of the flags must be included. If both of the LJ and torsional terms are turned off, it becomes the second-generation REBO potential, with a small caveat on the spline fitting procedure mentioned below. This can be specified directly as `pair_style rebo` with no additional arguments.

The detailed formulas for this potential are given in (Stuart); here we provide only a brief description.

The  $E^{\text{REBO}}$  term has the same functional form as the hydrocarbon REBO potential developed in (Brenner). The coefficients for  $E^{\text{REBO}}$  in AIREBO are essentially the same as Brenner's potential, but a few fitted spline values are slightly different. For most cases the  $E^{\text{REBO}}$  term in AIREBO will produce the same energies, forces and statistical averages as the original REBO potential from which it was derived. The  $E^{\text{REBO}}$  term in the AIREBO potential gives the model its reactive capabilities and only describes short-ranged C-C, C-H and H-H interactions ( $r < 2$  Angstroms). These interactions have strong coordination-dependence through a bond order parameter, which adjusts the attraction between the I,J atoms based on the position of other nearby atoms and thus has 3- and 4-body dependence.

The  $E^{\text{LJ}}$  term adds longer-ranged interactions ( $2 < r < \text{cutoff}$ ) using a form similar to the standard *Lennard Jones potential*. The  $E^{\text{LJ}}$  term in AIREBO contains a series of switching functions so that the short-ranged LJ repulsion ( $1/r^{12}$ ) does not interfere with the energetics captured by the  $E^{\text{REBO}}$  term. The extent of the  $E^{\text{LJ}}$  interactions is determined by the *cutoff* argument to the `pair_style` command which is a scale factor. For each type pair (C-C, C-H, H-H) the cutoff is obtained by multiplying the scale factor by the sigma value defined in the potential file for that type pair. In the standard AIREBO potential,  $\sigma_{CC} = 3.4$  Angstroms, so with a scale factor of 3.0 (the argument in `pair_style`), the resulting  $E^{\text{LJ}}$  cutoff would be 10.2 Angstroms.

By default, the longer-ranged interaction is smoothly switched off between 2.16 and 3.0  $\sigma$ . By specifying *cutoff\_min* in addition to *cutoff*, the switching can be configured to take place between *cutoff\_min* and *cutoff*. *cutoff\_min* can only be specified if all optional arguments are given.

The  $E^{\text{TORSION}}$  term is an explicit 4-body potential that describes various dihedral angle preferences in hydrocarbon configurations.

Only a single `pair_coeff` command is used with the *airebo*, *airebo* or *rebo* style which specifies an AIREBO, REBO, or AIREBO-M potential file with parameters for C and H. Note that as of LAMMPS version 15 May 2019 the *rebo* style in LAMMPS uses its own potential file (CH.rebo). These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of AIREBO elements to atom types

See the *pair\_coeff* page for alternate ways to specify the path for the potential file.

As an example, if your LAMMPS simulation has 4 atom types and you want the first 3 to be C, and the fourth to be H, you would use the following `pair_coeff` command:

```
pair_coeff * * CH.airebo C C C H
```

The first 2 arguments must be \* \* so as to span all LAMMPS atom types. The first three C arguments map LAMMPS atom types 1,2,3 to the C element in the AIREBO file. The final H argument maps LAMMPS atom type 4 to the H element in the SW file. If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *airebo* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

The parameters/coefficients for the AIREBO potentials are listed in the CH.airebo file to agree with the original (*Stuart*) paper. Thus the parameters are specific to this potential and the way it was fit, so modifying the file should be done cautiously.

Similarly the parameters/coefficients for the AIREBO-M potentials are listed in the CH.airebo-m file to agree with the (*O'Connor*) paper. Thus the parameters are specific to this potential and the way it was fit, so modifying the file should be done cautiously. The AIREBO-M Morse potentials were parameterized using a cutoff of 3.0 ( $\sigma$ ). Modifying this cutoff may impact simulation accuracy.

This pair style tallies a breakdown of the total AIREBO potential energy into sub-categories, which can be accessed via the *compute pair* command as a vector of values of length 3. The 3 values correspond to the following sub-categories:

1.  $E_{\text{REBO}}$  = REBO energy
2.  $E_{\text{LJ}}$  = Lennard-Jones energy
3.  $E_{\text{TORSION}}$  = Torsion energy

To print these quantities to the log file (with descriptive column headings) the following commands could be included in an input script:

```
compute 0 all pair airebo
variable REBO      equal c_0[1]
variable LJ        equal c_0[2]
variable TORSION   equal c_0[3]
thermo_style custom step temp epair v_REBO v_LJ v_TORSION
```

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.5.4 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles do not support the *pair\_modify* mix, shift, table, and tail options.

These pair styles do not write their information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

## 19.5.5 Restrictions

These pair styles are part of the MANYBODY package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These pair potentials require the *newton* setting to be “on” for pair interactions.

The CH.airebo and CH.airebo-m potential files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use the AIREBO, AIREBO-M or REBO potential with any LAMMPS units, but you would need to create your own AIREBO or AIREBO-M potential file with coefficients listed in the appropriate units, if your simulation does not use “metal” units.

The pair styles provided here **only** support potential files parameterized for the elements carbon and hydrogen (designated with “C” and “H” in the *pair\_coeff* command. Using potential files for other elements will trigger an error.

## 19.5.6 Related commands

*pair\_coeff*

## 19.5.7 Default

none

---

(**Stuart**) Stuart, Tutein, Harrison, J Chem Phys, 112, 6472-6486 (2000).

(**Brenner**) Brenner, Shenderova, Harrison, Stuart, Ni, Sinnott, J Physics: Condensed Matter, 14, 783-802 (2002).

(**O’Connor**) O’Connor et al., J. Chem. Phys. 142, 024903 (2015).

## 19.6 pair\_style amoeba command

## 19.7 pair\_style hippo command

### 19.7.1 Syntax

```
pair_style style
```

- style = *amoeba* or *hippo*

### 19.7.2 Examples

```
pair_style amoeba
pair_coeff * * protein.prm.amoeba protein.key.amoeba
```

```
pair_style hippo
pair_coeff * * water.prm.hippo water.key.hippo
```

### 19.7.3 Additional info

- [Howto amoeba](#)
- [examples/amoeba](#)
- [tools/amoeba](#)
- [potentials/\\*.amoeba](#)
- [potentials/\\*.hippo](#)

### 19.7.4 Description

The *amoeba* style computes the AMOEBA polarizable field formulated by Jay Ponder’s group at the U Washington at St Louis (*Ren*), (*Shi*). The *hippo* style computes the HIPPO polarizable force field, an extension to AMOEBA, formulated by Josh Rackers and collaborators in the Ponder group (*Rackers*).

These force fields can be used when polarization effects are desired in simulations of water, organic molecules, and biomolecules including proteins, provided that parameterizations (Tinker PRM force field files) are available for the systems you are interested in. Files in the LAMMPS potentials directory with a “amoeba” or “hippo” suffix can be used. The Tinker distribution and website have additional force field files as well.

As discussed on the [Howto amoeba](#) doc page, the intermolecular (non-bonded) portion of the AMOEBA force field contains these terms:

$$U_{amoeba} = U_{multipole} + U_{polar} + U_{hal}$$

while the HIPPO force field contains these terms:

$$U_{hippo} = U_{multipole} + U_{polar} + U_{qxf} + U_{repulsion} + U_{dispersion}$$

Conceptually, these terms compute the following interactions:

- $U_{hal}$  = buffered 14-7 van der Waals with offsets applied to hydrogen atoms
- $U_{repulsion}$  = Pauli repulsion due to rearrangement of electron density
- $U_{dispersion}$  = dispersion between correlated, instantaneous induced dipole moments
- $U_{multipole}$  = electrostatics between permanent point charges, dipoles, and quadrupoles
- $U_{polar}$  = electronic polarization between induced point dipoles
- $U_{qxf}$  = charge transfer effects

Note that the AMOEBA versus HIPPO force fields typically compute the same term differently using their own formulas. The references on this doc page give full details for both force fields.

The formulas for the AMOEBA energy terms are:

$$U_{hal} = \epsilon_{ij} \left( \frac{1.07}{\rho_{ij} + 0.07} \right)^7 \left( \frac{1.12}{\rho_{ij}^7 + 0.12} - 2 \right) U_{multipole} = \vec{M}_i \mathbf{T}_{ij} \vec{M}_j \vec{M} = (q, \mu_{perm}, \Theta) U_{polar} = \frac{1}{2} \vec{\mu}_i^{ind} \vec{E}_i^{perm}$$

The formulas for the HIPPO energy terms are:

$$U_{multipole} = Z_i \frac{1}{r_{ij}} Z_j + Z_i T_{ij}^{damp} \vec{M}_j + Z_j T_{ji}^{damp} \vec{M}_i + \vec{M}_i T_{ij}^{damp} \vec{M}_j \vec{M} = (Q, \mu_{perm}, \Theta) U_{polar} = \frac{1}{2} \vec{\mu}_i^{ind} \vec{E}_i^{perm} U_{qxf} = \epsilon_i e^{-\eta_j r_{ij}}$$

**Note:** The AMOEBA and HIPPO force fields compute long-range charge, dipole, and quadrupole interactions as well as long-range dispersion effects. However, unlike other models with long-range interactions in LAMMPS, this does not

require use of a KSpace style via the *kpace\_style* command. That is because for AMOEBA and HIPPO the long-range computations are intertwined with the pairwise computations. So these pair style include both short- and long-range computations. This means the energy and virial computed by the pair style as well as the “Pair” timing reported by LAMMPS will include the long-range calculations.

---

The implementation of the AMOEBA and HIPPO force fields in LAMMPS was done using F90 code provided by the Ponder group from their [Tinker MD code](#).

The current implementation (July 2022) of AMOEBA in LAMMPS matches the version discussed in (*Ponder*), (*Ren*), and (*Shi*). Likewise the current implementation of HIPPO in LAMMPS matches the version discussed in (*Rackers*).

---

Only a single `pair_coeff` command is used with either the *amoeba* and *hippo* styles which specifies two Tinker files, a PRM and KEY file.

```
pair_coeff * * ../potentials/protein.prm.amoeba ../potentials/protein.key.amoeba
pair_coeff * * ../potentials/water.prm.hippo ../potentials/water.key.hippo
```

Examples of the PRM files are in the potentials directory with an \*.amoeba or \*.hippo suffix. The examples/amoeba directory has examples of both PRM and KEY files.

A Tinker PRM file is composed of sections, each of which has multiple lines. A Tinker KEY file is composed of lines, each of which has a keyword followed by zero or more parameters.

The list of PRM sections and KEY keywords which LAMMPS recognizes are listed on the [Howto amoeba](#) doc page. If not recognized, the section or keyword is skipped.

Note that if the KEY file is specified as NULL, then no file is required; default values for various AMOEBA/HIPPO settings are used. The [Howto amoeba](#) doc page also gives the default settings.

---

New in version TBD.

The *amoeba* and *hippo* pair styles support extraction of two per-atom quantities by the *fix pair* command. This allows the quantities to be output to files by the *dump* or otherwise processed by other LAMMPS commands.

The names of the two quantities are “uind” and “uinp” for the induced dipole moments for each atom. Neither quantity needs to be triggered by the *fix pair* command in order for these pair styles to calculate it.

---

### 19.7.5 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles do not support the *pair\_modify* mix, shift, table, and tail options.

These pair styles do not write their information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

---

## 19.7.6 Restrictions

These pair styles are part of the AMOEBA package. They are only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

The AMOEBA and HIPPO potential (PRM) and KEY files provided with LAMMPS in the potentials and examples/amoeba directories are Tinker files parameterized for Tinker units. Their numeric parameters are converted by LAMMPS to its real units *units*. Thus you can only use these pair styles with real units.

These potentials do not yet calculate per-atom energy or virial contributions.

As explained on the *AMOEBA and HIPPO howto* page, use of these pair styles to run a simulation with the AMOEBA or HIPPO force fields requires several things.

The first is a data file generated by the tools/tinker/tinker2lmp.py conversion script which uses Tinker file force field file input to create a data file compatible with LAMMPS.

The second is use of these commands:

- *atom\_style amoeba*
- *fix property/atom*
- *special\_bonds one/five*

And third, depending on the model being simulated, these commands for intramolecular interactions may also be required:

- *bond\_style class2*
- *angle\_style amoeba*
- *dihedral\_style fourier*
- *improper\_style amoeba*
- *fix amoeba/pitursion*
- *fix amoeba/bitursion*

---

## 19.7.7 Related commands

*atom\_style amoeba*, *bond\_style class2*, *angle\_style amoeba*, *dihedral\_style fourier*, *improper\_style amoeba*, *fix amoeba/pitursion*, *fix amoeba/bitursion*, *special\_bonds one/five*, *fix property/atom*

## 19.7.8 Default

none

---

**(Ponder)** Ponder, Wu, Ren, Pande, Chodera, Schnieders, Haque, Mobley, Lambrecht, DiStasio Jr, M. Head-Gordon, Clark, Johnson, T. Head-Gordon, J Phys Chem B, 114, 2549-2564 (2010).

**(Rackers)** Rackers, Silva, Wang, Ponder, J Chem Theory Comput, 17, 7056-7084 (2021).

**(Ren)** Ren and Ponder, J Phys Chem B, 107, 5933 (2003).

**(Shi)** Shi, Xia, Zhang, Best, Wu, Ponder, Ren, J Chem Theory Comp, 9, 4046, 2013.

## 19.8 pair\_style atm command

### 19.8.1 Syntax

```
pair_style atm cutoff cutoff_triple
```

- cutoff = cutoff for each pair in 3-body interaction (distance units)
- cutoff\_triple = additional cutoff applied to product of 3 pairwise distances (distance units)

### 19.8.2 Examples

```
pair_style atm 4.5 2.5
pair_coeff * * * 0.072

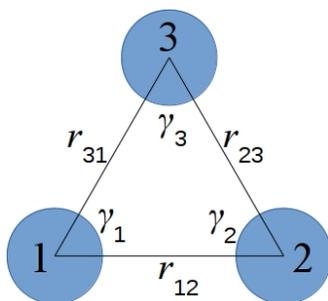
pair_style hybrid/overlay lj/cut 6.5 atm 4.5 2.5
pair_coeff * * lj/cut 1.0 1.0
pair_coeff 1 1 atm 1 0.064
pair_coeff 1 1 atm 2 0.080
pair_coeff 1 2 atm 2 0.100
pair_coeff 2 2 atm 2 0.125
```

### 19.8.3 Description

The *atm* style computes a 3-body *Axilrod-Teller-Muto* potential for the energy  $E$  of a system of atoms as

$$E = \nu \frac{1 + 3 \cos \gamma_1 \cos \gamma_2 \cos \gamma_3}{r_{12}^3 r_{23}^3 r_{31}^3}$$

where  $\nu$  is the three-body interaction strength. The distances between pairs of atoms  $r_{12}$ ,  $r_{23}$ ,  $r_{31}$  and the angles  $\gamma_1$ ,  $\gamma_2$ ,  $\gamma_3$  are as shown in this diagram:



Note that for the interaction between a triplet of atoms  $I, J, K$ , there is no “central” atom. The interaction is symmetric with respect to permutation of the three atoms. Thus the  $\nu$  value is the same for all those permutations of the atom types of  $I, J, K$  and needs to be specified only once, as discussed below.

The *atm* potential is typically used in combination with a two-body potential using the *pair\_style hybrid/overlay* command as in the example above.

The potential for a triplet of atom is calculated only if all 3 distances  $r_{12}$ ,  $r_{23}$ ,  $r_{31}$  between the 3 atoms satisfy  $r_{IJ} < \text{cutoff}$ . In addition, the product of the 3 distances  $r_{12}r_{23}r_{31} < \text{cutoff\_triple}^3$  is required, which excludes from calculation the triplets with small contribution to the interaction.

The following coefficients must be defined for each pair of atoms types via the `pair_coeff` command as in the examples above, or in the restart files read by the `read_restart` commands:

- $K$  = atom type of the third atom (1 to  $N_{\text{types}}$ )
- $\nu$  = prefactor (energy/distance<sup>9</sup> units)

$K$  can be specified in one of two ways. An explicit numeric value can be used, as in the second example above.  $J \leq K$  is required. LAMMPS sets the coefficients for the other 5 symmetric interactions to the same values. E.g. if  $I = 1$ ,  $J = 2$ ,  $K = 3$ , then these 6 values are set to the specified  $\nu$ :  $\nu_{123}$ ,  $\nu_{132}$ ,  $\nu_{213}$ ,  $\nu_{231}$ ,  $\nu_{312}$ ,  $\nu_{321}$ . This enforces the symmetry discussed above.

A wildcard asterisk can be used for  $K$  to set the coefficients for multiple triplets of atom types. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. If  $N$  equals the number of atom types, then an asterisk with no numeric values means all types from 1 to  $N$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $n$  to  $N$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive). Note that only type triplets with  $J \leq K$  are considered; if asterisks imply type triplets where  $K < J$ , they are ignored.

Note that a `pair_coeff` command can override a previous setting for the same  $I, J, K$  triplet. For example, these commands set  $\nu$  for all  $I, J, K$  triplets, then overwrite  $\nu$  for just the  $I, J, K = 2, 3, 4$  triplet:

```
pair_coeff * * * 0.25
pair_coeff 2 3 4 0.1
```

Note that for a simulation with a single atom type, only a single entry is required, e.g.

```
pair_coeff 1 1 1 0.25
```

For a simulation with two atom types, four `pair_coeff` commands will specify all possible  $\nu$  values:

```
pair_coeff 1 1 1 nu1
pair_coeff 1 1 2 nu2
pair_coeff 1 2 2 nu3
pair_coeff 2 2 2 nu4
```

For a simulation with three atom types, ten `pair_coeff` commands will specify all possible  $\nu$  values:

```
pair_coeff 1 1 1 nu1
pair_coeff 1 1 2 nu2
pair_coeff 1 1 3 nu3
pair_coeff 1 2 2 nu4
pair_coeff 1 2 3 nu5
pair_coeff 1 3 3 nu6
pair_coeff 2 2 2 nu7
pair_coeff 2 2 3 nu8
pair_coeff 2 3 3 nu9
pair_coeff 3 3 3 nu10
```

By default the  $\nu$  value for all triplets is set to 0.0. Thus it is not required to provide `pair_coeff` commands that enumerate triplet interactions for all  $K$  types. If some  $I, J, K$  combination is not specified, then there will be no 3-body ATM interactions for that combination and all its permutations. However, as with all pair styles, it is required to specify a `pair_coeff` command for all  $I, J$  combinations, else an error will result.



## 19.8.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style do not support the *pair\_modify* mix, shift, table, and tail options.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file. However, if the *atm* potential is used in combination with other potentials using the *pair\_style hybrid/overlay* command then *pair\_coeff* commands need to be re-specified in the restart input script.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, and *outer* keywords.

## 19.8.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.8.6 Related commands

*pair\_coeff*

## 19.8.7 Default

none

(**Axilrod**) Axilrod and Teller, J Chem Phys, 11, 299 (1943); Muto, Nippon Sugaku-Buturigakkwaishi 17, 629 (1943).

# 19.9 pair\_style awpmd/cut command

## 19.9.1 Syntax

```
pair_style awpmd/cut Rc keyword value ...
```

- Rc = global cutoff, -1 means cutoff of half the shortest box length
- zero or more keyword/value pairs may be appended
- keyword = *hartree* or *dproduct* or *uhf* or *free* or *pbs* or *fix* or *harm* or *ermscale* or *flex\_press*

*hartree* value = none

*dproduct* value = none

*uhf* value = none

*free* value = none

*pbc* value = Plen

Plen = periodic width of electron = -1 or positive value (distance units)

*fix* value = Flen

Flen = fixed width of electron = -1 or positive value (distance units)

*harm* value = width

width = harmonic width constraint

```
ermscale value = factor
  factor = scaling between electron mass and width variable mass
flex_press value = none
```

## 19.9.2 Examples

```
pair_style awpmd/cut -1
pair_style awpmd/cut 40.0 uhf free
pair_coeff * *
pair_coeff 2 2 20.0
```

## 19.9.3 Description

This pair style contains an implementation of the Antisymmetrized Wave Packet Molecular Dynamics (AWPMD) method. Need citation here. Need basic formulas here. Could be links to other documents.

Rc is the cutoff.

The pair\_style command allows for several optional keywords to be specified.

The *hartree*, *dproduct*, and *uhf* keywords specify the form of the initial trial wave function for the system. If the *hartree* keyword is used, then a Hartree multielectron trial wave function is used. If the *dproduct* keyword is used, then a trial function which is a product of two determinants for each spin type is used. If the *uhf* keyword is used, then an unrestricted Hartree-Fock trial wave function is used.

The *free*, *pbc*, and *fix* keywords specify a width constraint on the electron wave packets. If the *free* keyword is specified, then there is no constraint. If the *pbc* keyword is used and *Plen* is specified as -1, then the maximum width is half the shortest box length. If *Plen* is a positive value, then the value is the maximum width. If the *fix* keyword is used and *Flen* is specified as -1, then electrons have a constant width that is read from the data file. If *Flen* is a positive value, then the constant width for all electrons is set to *Flen*.

The *harm* keyword allow oscillations in the width of the electron wave packets. More details are needed.

The *ermscale* keyword specifies a unitless scaling factor between the electron masses and the width variable mass. More details needed.

If the *flex\_press* keyword is used, then a contribution from the electrons is added to the total virial and pressure of the system.

This potential is designed to be used with *atom\_style wavepacket* definitions, in order to handle the description of systems with interacting nuclei and explicit electrons.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- cutoff (distance units)

For *awpmd/cut*, the cutoff coefficient is optional. If it is not used (as in some of the examples above), the default global value specified in the pair\_style command is used.

## 19.9.4 Mixing, shift, table, tail correction, restart, rRESPA info

The *pair\_modify* mix, shift, table, and tail options are not relevant for this pair style.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

## 19.9.5 Restrictions

none

## 19.9.6 Related commands

*pair\_coeff*

## 19.9.7 Default

These are the defaults for the *pair\_style* keywords: *hartree* for the initial wave function, *free* for the wave packet width.

## 19.10 pair\_style beck command

Accelerator Variants: *beck/gpu*, *beck/omp*

### 19.10.1 Syntax

```
pair_style beck Rc
```

- Rc = cutoff for interactions (distance units)

### 19.10.2 Examples

```
pair_style beck 8.0
pair_coeff * * 399.671876712 0.0000867636112694 0.675 4.390 0.0003746
pair_coeff 1 1 399.671876712 0.0000867636112694 0.675 4.390 0.0003746 6.0
```

### 19.10.3 Description

Style *beck* computes interactions based on the potential by (*Beck*), originally designed for simulation of Helium. It includes truncation at a cutoff distance  $R_c$ .

$$E(r) = A \exp[-\alpha r - \beta r^6] - \frac{B}{(r^2 + a^2)^3} \left( 1 + \frac{2.709 + 3a^2}{r^2 + a^2} \right) \quad r < R_c$$

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands.

- $A$  (energy units)
- $B$  (energy-distance<sup>6</sup> units)
- $a$  (distance units)
- $\alpha$  (1/distance units)
- $\beta$  (1/distance<sup>6</sup> units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global cutoff  $R_c$  is used.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.10.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, coefficients must be specified. No default mixing rules are used.

This pair style does not support the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

## 19.10.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.10.6 Related commands

*pair\_coeff*

## 19.10.7 Default

none

---

(Beck) Beck, Molecular Physics, 14, 311 (1968).

## 19.11 pair\_style body/nparticle command

### 19.11.1 Syntax

```
pair_style body/nparticle cutoff
```

cutoff = global cutoff for interactions (distance units)

### 19.11.2 Examples

```
pair_style body/nparticle 3.0
pair_coeff * * 1.0 1.0
pair_coeff 1 1 1.0 1.5 2.5
```

### 19.11.3 Description

Style *body/nparticle* is for use with body particles and calculates pairwise body/body interactions as well as interactions between body and point-particles. See the *Howto body* doc page for more details on using body particles.

This pair style is designed for use with the “nparticle” body style, which is specified as an argument to the “atom-style body” command. See the *Howto body* page for more details about the body styles LAMMPS supports. The “nparticle” style treats a body particle as a rigid body composed of N sub-particles.

The coordinates of a body particle are its center-of-mass (COM). If the COMs of a pair of body particles are within the cutoff (global or type-specific, as specified above), then all interactions between pairs of sub-particles in the two body particles are computed. E.g. if the first body particle has 3 sub-particles, and the second has 10, then 30 interactions are computed and summed to yield the total force and torque on each body particle.

---

**Note:** In the example just described, all 30 interactions are computed even if the distance between a particular pair of sub-particles is greater than the cutoff. Likewise, no interaction between two body particles is computed if the two

COMs are further apart than the cutoff, even if the distance between some pairs of their sub-particles is within the cutoff. Thus care should be used in defining the cutoff distances for body particles, depending on their shape and size.

---

Similar rules apply for a body particle interacting with a point particle. The distance between the two particles is calculated using the COM of the body particle and the position of the point particle. If the distance is within the cutoff and the body particle has  $N$  sub-particles, then  $N$  interactions with the point particle are computed and summed. If the distance is not within the cutoff, no interactions between the body and point particle are computed.

The interaction between two sub-particles, or a sub-particle and point particle, or between two point particles is computed as a Lennard-Jones interaction, using the standard formula

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < R_c$$

where  $R_c$  is the cutoff. As explained above, an interaction involving one or two body sub-particles may be computed even for  $r > R_c$ .

For style *body*, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global cutoff is used.

---

#### 19.11.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs  $I, J$  and  $I \neq J$ , the epsilon and sigma coefficients and cutoff distance for all of this pair style can be mixed. The default mix value is *geometric*. See the *pair\_modify* command for details.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.11.5 Restrictions

This style is part of the BODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Defining particles to be bodies so they participate in body/body or body/particle interactions requires the use of the *atom\_style body* command.

## 19.11.6 Related commands

*pair\_coeff*, *fix rigid*

## 19.11.7 Default

none

# 19.12 pair\_style body/rounded/polygon command

## 19.12.1 Syntax

```
pair_style body/rounded/polygon c_n c_t mu delta_ua cutoff
```

```
c_n = normal damping coefficient
c_t = tangential damping coefficient
mu = normal friction coefficient during gross sliding
delta_ua = multiple contact scaling factor
cutoff = global separation cutoff for interactions (distance units), see below for definition
```

## 19.12.2 Examples

```
pair_style body/rounded/polygon 20.0 5.0 0.0 1.0 0.5
pair_coeff * * 100.0 1.0
pair_coeff 1 1 100.0 1.0
```

## 19.12.3 Description

Style *body/rounded/polygon* is for use with 2d models of body particles of style *rounded/polygon*. It calculates pairwise body/body interactions which can include body particles modeled as 1-vertex circular disks with a specified diameter. See the [Howto body](#) page for more details on using body rounded/polygon particles.

This pairwise interaction between rounded polygons is described in *Fraige*, where a polygon does not have sharp corners, but is rounded at its vertices by circles centered on each vertex with a specified diameter. The edges of the polygon are defined between pairs of adjacent vertices. The circle diameter for each polygon is specified in the data file read by the *read data* command. This is a 2d discrete element model (DEM) which allows for multiple contact points.

Note that when two particles interact, the effective surface of each polygon particle is displaced outward from each of its vertices and edges by half its circle diameter (as in the diagram below of a gray and yellow square particle). The interaction forces and energies between two particles are defined with respect to the separation of their respective rounded surfaces, not by the separation of the vertices and edges themselves.

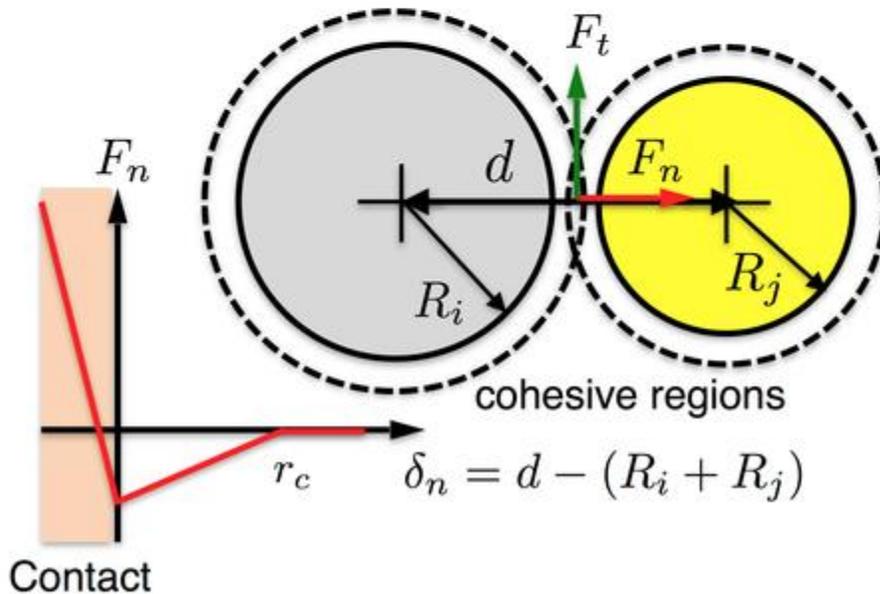
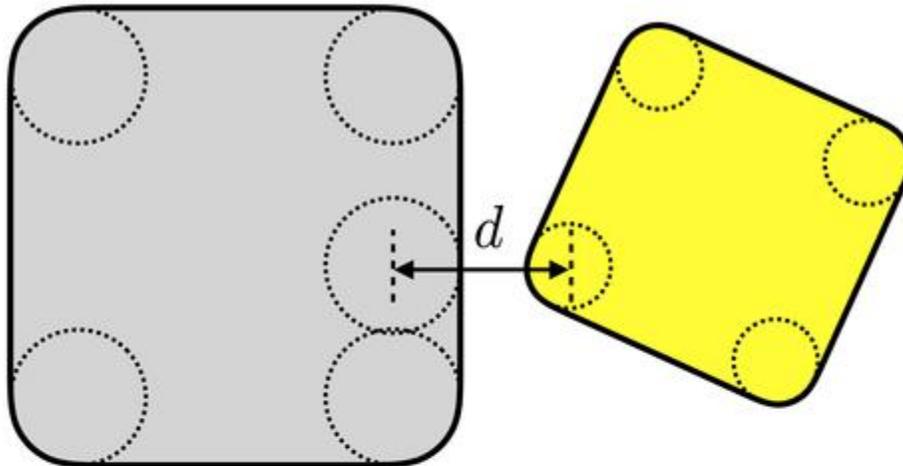
This means that the specified cutoff in the *pair\_style* command is the cutoff distance,  $r_c$ , for the surface separation,  $\delta_n$  (see figure below). This is the distance at which two particles no longer interact. If  $r_c$  is specified as 0.0, then it is a contact-only interaction. I.e. the two particles must overlap in order to exert a repulsive force on each other. If  $r_c > 0.0$ , then the force between two particles will be attractive for surface separations from 0 to  $r_c$ , and repulsive once the particles overlap.

Note that unlike for other pair styles, the specified cutoff is not the distance between the centers of two particles at which they stop interacting. This center-to-center distance depends on the shape and size of the two particles and their relative orientation. LAMMPS takes that into account when computing the surface separation distance and applying the  $r_c$  cutoff.

The forces between vertex-vertex, vertex-edge, and edge-edge overlaps are given by:

$$F_n = \begin{cases} k_n \delta_n - c_n v_n & \delta_n \leq 0 \\ -k_{na} \delta_n - c_n v_n & 0 < \delta_n \leq r_c \\ 0 & \delta_n > r_c \end{cases}$$

$$F_t = \begin{cases} \mu k_n \delta_n - c_t v_t & \delta_n \leq 0 \\ 0 & \delta_n > 0 \end{cases}$$



Note that  $F_n$  and  $F_t$  are functions of the surface separation  $\delta_n = d - (R_i + R_j)$ . In this model, when  $(R_i + R_j) <$



$d < (R_i + R_j) + r_c$ , that is,  $0 < \delta_n < r_c$ , the cohesive region of the two surfaces overlap and the two surfaces are attractive to each other.

In *Fraige*, the tangential friction force between two particles that are in contact is modeled differently prior to gross sliding (i.e. static friction) and during gross-sliding (kinetic friction). The latter takes place when the tangential deformation exceeds the Coulomb frictional limit. In the current implementation, however, we do not take into account frictional history, i.e. we do not keep track of how many time steps the two particles have been in contact nor calculate the tangential deformation. Instead, we assume that gross sliding takes place as soon as two particles are in contact.

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file read by the *read\_data* command:

- $k_n$  (energy/distance<sup>2</sup> units)
- $k_{na}$  (energy/distance<sup>2</sup> units)

Effectively,  $k_n$  and  $k_{na}$  are the slopes of the red lines in the plot above for force versus surface separation, for  $\delta_n < 0$  and  $0 < \delta_n < r_c$  respectively.

### 19.12.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* mix, shift, table, and tail options.

This pair style does not write its information to *binary restart files*. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.12.5 Restrictions

These pair styles are part of the BODY package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

### 19.12.6 Related commands

*pair\_coeff*

### 19.12.7 Default

none

**(Fraige)** F. Y. Fraige, P. A. Langston, A. J. Matchett, J. Dodds, *Particuology*, 6, 455 (2008).

## 19.13 pair\_style body/rounded/polyhedron command

### 19.13.1 Syntax

```
pair_style body/rounded/polyhedron c_n c_t mu delta_ua cutoff
```

```
c_n = normal damping coefficient
c_t = tangential damping coefficient
mu = normal friction coefficient during gross sliding
delta_ua = multiple contact scaling factor
cutoff = global separation cutoff for interactions (distance units), see below for definition
```

### 19.13.2 Examples

```
pair_style body/rounded/polyhedron 20.0 5.0 0.0 1.0 0.5
pair_coeff * * 100.0 1.0
pair_coeff 1 1 100.0 1.0
```

### 19.13.3 Description

Style *body/rounded/polygon* is for use with 3d models of body particles of style *rounded/polyhedron*. It calculates pairwise body/body interactions which can include body particles modeled as 1-vertex spheres with a specified diameter. See the [Howto body](#) page for more details on using body rounded/polyhedron particles.

This pairwise interaction between the rounded polyhedra is described in [Wang](#), where a polyhedron does not have sharp corners and edges, but is rounded at its vertices and edges by spheres centered on each vertex with a specified diameter. The edges of the polyhedron are defined between pairs of adjacent vertices. Its faces are defined by a loop of edges. The sphere diameter for each polygon is specified in the data file read by the [read data](#) command. This is a discrete element model (DEM) which allows for multiple contact points.

Note that when two particles interact, the effective surface of each polyhedron particle is displaced outward from each of its vertices, edges, and faces by half its sphere diameter. The interaction forces and energies between two particles are defined with respect to the separation of their respective rounded surfaces, not by the separation of the vertices, edges, and faces themselves.

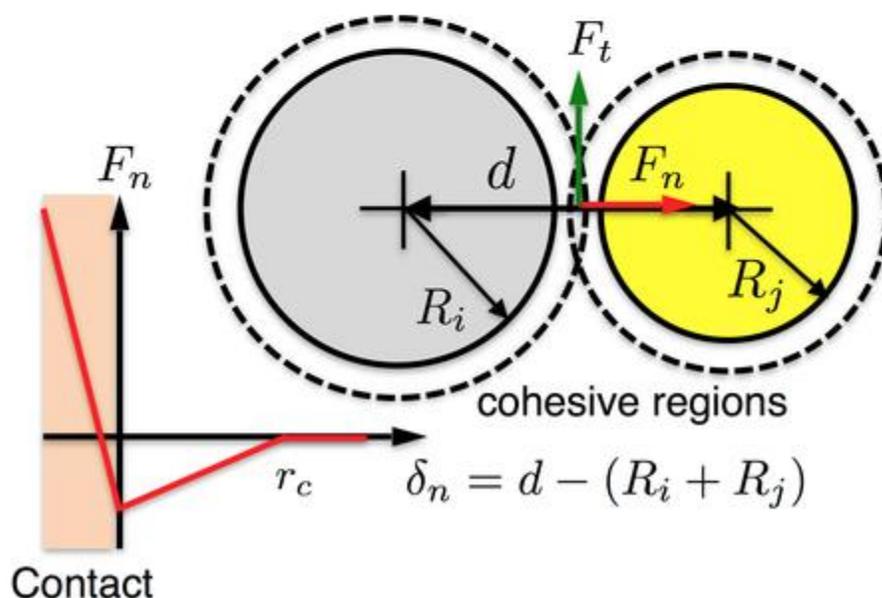
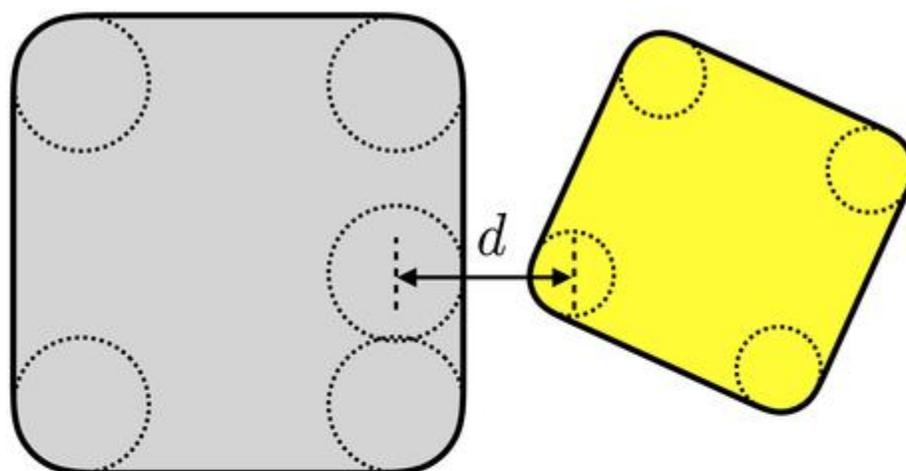
This means that the specified cutoff in the `pair_style` command is the cutoff distance,  $r_c$ , for the surface separation,  $\delta_n$  (see figure below). This is the distance at which two particles no longer interact. If  $r_c$  is specified as 0.0, then it is a contact-only interaction. I.e. the two particles must overlap in order to exert a repulsive force on each other. If  $r_c > 0.0$ , then the force between two particles will be attractive for surface separations from 0 to  $r_c$ , and repulsive once the particles overlap.

Note that unlike for other pair styles, the specified cutoff is not the distance between the centers of two particles at which they stop interacting. This center-to-center distance depends on the shape and size of the two particles and their relative orientation. LAMMPS takes that into account when computing the surface separation distance and applying the  $r_c$  cutoff.

The forces between vertex-vertex, vertex-edge, vertex-face, edge-edge, and edge-face overlaps are given by:

$$F_n = \begin{cases} k_n \delta_n - c_n v_n, & \delta_n \leq 0 \\ -k_{na} \delta_n - c_n v_n & 0 < \delta_n \leq r_c \\ 0 & \delta_n > r_c \end{cases}$$

$$F_t = \begin{cases} \mu k_n \delta_n - c_t v_t & \delta_n \leq 0 \\ 0 & \delta_n > 0 \end{cases}$$



In *Wang*, the tangential friction force between two particles that are in contact is modeled differently prior to gross sliding (i.e. static friction) and during gross-sliding (kinetic friction). The latter takes place when the tangential deformation exceeds the Coulomb frictional limit. In the current implementation, however, we do not take into account frictional history, i.e. we do not keep track of how many time steps the two particles have been in contact nor calculate the tangential deformation. Instead, we assume that gross sliding takes place as soon as two particles are in contact.

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file read by the *read\_data* command:

- $k_n$  (energy/distance<sup>2</sup> units)
- $k_{na}$  (energy/distance<sup>2</sup> units)

Effectively,  $k_n$  and  $k_{na}$  are the slopes of the red lines in the plot above for force versus surface separation, for  $\delta_n < 0$  and  $0 < \delta_n < r_c$  respectively.

### 19.13.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* mix, shift, table, and tail options.

This pair style does not write its information to *binary restart files*. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.13.5 Restrictions

These pair styles are part of the BODY package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

### 19.13.6 Related commands

*pair\_coeff*

### 19.13.7 Default

none

(Wang) J. Wang, H. S. Yu, P. A. Langston, F. Y. Fraige, *Granular Matter*, 13, 1 (2011).

## 19.14 *pair\_style* bop command

### 19.14.1 Syntax

```
pair_style bop keyword ...
```

- zero or more keywords may be appended
- keyword = *save*

```
save = pre-compute and save some values
```

## 19.14.2 Examples

```
pair_style bop
pair_coeff * * ../potentials/CdTe_bop Cd Te
pair_style bop save
pair_coeff * * ../potentials/CdTe.bop.table Cd Te Te
comm_modify cutoff 14.70
```

## 19.14.3 Description

The *bop* pair style computes Bond-Order Potentials (BOP) based on quantum mechanical theory incorporating both  $\sigma$  and  $\pi$  bonding. By analytically deriving the BOP from quantum mechanical theory its transferability to different phases can approach that of quantum mechanical methods. This potential is similar to the original BOP developed by Pettifor (*Pettifor\_1*, *Pettifor\_2*, *Pettifor\_3*) and later updated by Murdick, Zhou, and Ward (*Murdick*, *Ward*). Currently, BOP potential files for these systems are provided with LAMMPS: AlCu, CCu, CdTe, CdTeSe, CdZnTe, CuH, GaAs. A system with only a subset of these elements, including a single element (e.g. C or Cu or Al or Ga or Zn or CdZn), can also be modeled by using the appropriate alloy file and assigning all atom types to the single element or subset of elements via the `pair_coeff` command, as discussed below.

The BOP potential consists of three terms:

$$E = \frac{1}{2} \sum_{i=1}^N \sum_{j=i_1}^{i_N} \phi_{ij}(r_{ij}) - \sum_{i=1}^N \sum_{j=i_1}^{i_N} \beta_{\sigma,ij}(r_{ij}) \cdot \Theta_{\sigma,ij} - \sum_{i=1}^N \sum_{j=i_1}^{i_N} \beta_{\pi,ij}(r_{ij}) \cdot \Theta_{\pi,ij} + U_{prom}$$

where  $\phi_{ij}(r_{ij})$  is a short-range two-body function representing the repulsion between a pair of ion cores,  $\beta_{\sigma,ij}(r_{ij})$  and  $\beta_{\pi,ij}(r_{ij})$  are respectively sigma and  $\pi$  bond integrals,  $\Theta_{\sigma,ij}$  and  $\Theta_{\pi,ij}$  are  $\sigma$  and  $\pi$  bond-orders, and  $U_{prom}$  is the promotion energy for sp-valent systems.

The detailed formulas for this potential are given in Ward (*Ward*); here we provide only a brief description.

The repulsive energy  $\phi_{ij}(r_{ij})$  and the bond integrals  $\beta_{\sigma,ij}(r_{ij})$  and  $\beta_{\pi,ij}(r_{ij})$  are functions of the interatomic distance  $r_{ij}$  between atom  $i$  and  $j$ . Each of these potentials has a smooth cutoff at a radius of  $r_{cut,ij}$ . These smooth cutoffs ensure stable behavior at situations with high sampling near the cutoff such as melts and surfaces.

The bond-orders can be viewed as environment-dependent local variables that are ij bond specific. The maximum value of the  $\sigma$  bond-order ( $\Theta_{\sigma}$  is 1, while that of the  $\pi$  bond-order ( $\Theta_{\pi}$ ) is 2, attributing to a maximum value of the total bond-order ( $\Theta_{\sigma} + \Theta_{\pi}$ ) of 3. The  $\sigma$  and  $\pi$  bond-orders reflect the ubiquitous single-, double-, and triple- bond behavior of chemistry. Their analytical expressions can be derived from tight-binding theory by recursively expanding an inter-site Green's function as a continued fraction. To accurately represent the bonding with a computationally efficient potential formulation suitable for MD simulations, the derived BOP only takes (and retains) the first two levels of the recursive representations for both the  $\sigma$  and the  $\pi$  bond-orders. Bond-order terms can be understood in terms of molecular orbital hopping paths based upon the Cyrot-Lackmann theorem (*Pettifor\_1*). The  $\sigma$  bond-order with a half-full valence shell is used to interpolate the bond-order expression that incorporated explicit valence band filling. This  $\pi$  bond-order expression also contains a three-member ring term that allows implementation of an asymmetric density of states, which helps to either stabilize or destabilize close-packed structures. The  $\pi$  bond-order includes hopping paths of length 4. This enables the incorporation of dihedral angles effects.

**Note:** Note that unlike for other potentials, cutoffs for BOP potentials are not set in the `pair_style` or `pair_coeff` command; they are specified in the BOP potential files themselves. Likewise, the BOP potential files list atomic masses; thus you do not need to use the `mass` command to specify them. Note that for BOP potentials with hydrogen, you will likely want to set the mass of H atoms to be 10x or 20x larger to avoid having to use a tiny timestep. You can do this by using the `mass` command after using the `pair_coeff` command to read the BOP potential file.

One option can be specified as a keyword with the `pair_style` command.

The `save` keyword gives you the option to calculate in advance and store a set of distances, angles, and derivatives of angles. The default is to not do this, but to calculate them on-the-fly each time they are needed. The former may be faster, but takes more memory. The latter requires less memory, but may be slower. It is best to test this option to optimize the speed of BOP for your particular system configuration.

---

Only a single `pair_coeff` command is used with the `bop` style which specifies a BOP potential file, with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of BOP elements to atom types

As an example, imagine the `CdTe.bop` file has BOP values for Cd and Te. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Cd, and the fourth to be Te, you would use the following `pair_coeff` command:

```
pair_coeff * * CdTe Cd Cd Cd Te
```

The first 2 arguments must be `**` so as to span all LAMMPS atom types. The first three Cd arguments map LAMMPS atom types 1,2,3 to the Cd element in the BOP file. The final Te argument maps LAMMPS atom type 4 to the Te element in the BOP file.

BOP files in the `potentials` directory of the LAMMPS distribution have a “.bop” suffix. The potentials are in tabulated form containing pre-tabulated pair functions for  $\phi_{ij}(r_{ij})$ ,  $\beta_{(\sigma,ij)}(r_{ij})$ , and  $\beta_{\pi,ij}(r_{ij})$ .

The parameters/coefficients format for the different kinds of BOP files are given below with variables matching the formulation of Ward (*Ward*) and Zhou (*Zhou*). Each header line containing a “:” is preceded by a blank line.

---

### No angular table file format:

The parameters/coefficients format for the BOP potentials input file containing pre-tabulated functions of  $g$  is given below with variables matching the formulation of Ward (*Ward*). This format also assumes the angular functions have the formulation of (*Ward*).

- Line 1: # elements N

The first line is followed by N lines containing the atomic number, mass, and element symbol of each element.

Following the definition of the elements several global variables for the tabulated functions are given.

- Line 1: nr, nBOt (nr is the number of divisions the radius is broken into for function tables and MUST be a factor of 5; nBOt is the number of divisions for the tabulated values of  $\text{THETA}_{(S,ij)}$ )
- Line 2: delta\_1-delta\_7 (if all are not used in the particular formulation, set unused values to 0.0)

Following this N lines for  $e_1$ - $e_N$  containing  $p_{\pi}$ .

- Line 3:  $p_{\pi}$  (for  $e_1$ )
- Line 4:  $p_{\pi}$  (for  $e_2$  and continues to  $e_N$ )

The next section contains several pair constants for the number of interaction types  $e_i$ - $e_j$ , with  $i=1 \rightarrow N$ ,  $j=i \rightarrow N$

- Line 1:  $r_{\text{cut}}$  (for  $e_1$ - $e_1$  interactions)
- Line 2:  $c_{\sigma}$ ,  $a_{\sigma}$ ,  $c_{\pi}$ ,  $a_{\pi}$
- Line 3:  $\delta_{\sigma}$ ,  $\delta_{\pi}$

- Line 4:  $f\_sigma$ ,  $k\_sigma$ ,  $delta\_3$  (This  $delta\_3$  is similar to that of the previous section but is interaction type dependent)

The next section contains a line for each three body interaction type  $e\_j-e\_i-e\_k$  with  $i=0 \rightarrow N$ ,  $j=0 \rightarrow N$ ,  $k=j \rightarrow N$

- Line 1:  $g\_(\sigma_0)$ ,  $g\_(\sigma_1)$ ,  $g\_(\sigma_2)$  (These are coefficients for  $g\_(\sigma,ijk)(\theta_{ijk})$  for  $e\_1-e\_1-e\_1$  interaction. *Ward* contains the full expressions for the constants as functions of  $b\_(\sigma,ijk)$ ,  $p\_(\sigma,ijk)$ ,  $u\_(\sigma,ijk)$ )
- Line 2:  $g\_(\sigma_0)$ ,  $g\_(\sigma_1)$ ,  $g\_(\sigma_2)$  (for  $e\_1-e\_1-e\_2$ )

The next section contains a block for each interaction type for the  $\phi_{ij}(r_{ij})$ . Each block has  $n_r$  entries with 5 entries per line.

- Line 1:  $\phi(r_1)$ ,  $\phi(r_2)$ ,  $\phi(r_3)$ ,  $\phi(r_4)$ ,  $\phi(r_5)$  (for the  $e\_1-e\_1$  interaction type)
- Line 2:  $\phi(r_6)$ ,  $\phi(r_7)$ ,  $\phi(r_8)$ ,  $\phi(r_9)$ ,  $\phi(r_{10})$  (this continues until  $n_r$ )
- ...
- Line  $n_r/5+1$ :  $\phi(r_1)$ ,  $\phi(r_2)$ ,  $\phi(r_3)$ ,  $\phi(r_4)$ ,  $\phi(r_5)$ , (for the  $e\_1-e\_1$  interaction type)

The next section contains a block for each interaction type for the  $\beta_{ij}(\sigma,ij)(r_{ij})$ . Each block has  $n_r$  entries with 5 entries per line.

- Line 1:  $\beta_{ij}(\sigma,ij)(r_1)$ ,  $\beta_{ij}(\sigma,ij)(r_2)$ ,  $\beta_{ij}(\sigma,ij)(r_3)$ ,  $\beta_{ij}(\sigma,ij)(r_4)$ ,  $\beta_{ij}(\sigma,ij)(r_5)$  (for the  $e\_1-e\_1$  interaction type)
- Line 2:  $\beta_{ij}(\sigma,ij)(r_6)$ ,  $\beta_{ij}(\sigma,ij)(r_7)$ ,  $\beta_{ij}(\sigma,ij)(r_8)$ ,  $\beta_{ij}(\sigma,ij)(r_9)$ ,  $\beta_{ij}(\sigma,ij)(r_{10})$  (this continues until  $n_r$ )
- ...
- Line  $n_r/5+1$ :  $\beta_{ij}(\sigma,ij)(r_1)$ ,  $\beta_{ij}(\sigma,ij)(r_2)$ ,  $\beta_{ij}(\sigma,ij)(r_3)$ ,  $\beta_{ij}(\sigma,ij)(r_4)$ ,  $\beta_{ij}(\sigma,ij)(r_5)$  (for the  $e\_1-e\_2$  interaction type)

The next section contains a block for each interaction type for  $\beta_{ij}(\pi,ij)(r_{ij})$ . Each block has  $n_r$  entries with 5 entries per line.

- Line 1:  $\beta_{ij}(\pi,ij)(r_1)$ ,  $\beta_{ij}(\pi,ij)(r_2)$ ,  $\beta_{ij}(\pi,ij)(r_3)$ ,  $\beta_{ij}(\pi,ij)(r_4)$ ,  $\beta_{ij}(\pi,ij)(r_5)$  (for the  $e\_1-e\_1$  interaction type)
- Line 2:  $\beta_{ij}(\pi,ij)(r_6)$ ,  $\beta_{ij}(\pi,ij)(r_7)$ ,  $\beta_{ij}(\pi,ij)(r_8)$ ,  $\beta_{ij}(\pi,ij)(r_9)$ ,  $\beta_{ij}(\pi,ij)(r_{10})$  (this continues until  $n_r$ )
- ...
- Line  $n_r/5+1$ :  $\beta_{ij}(\pi,ij)(r_1)$ ,  $\beta_{ij}(\pi,ij)(r_2)$ ,  $\beta_{ij}(\pi,ij)(r_3)$ ,  $\beta_{ij}(\pi,ij)(r_4)$ ,  $\beta_{ij}(\pi,ij)(r_5)$  (for the  $e\_1-e\_2$  interaction type)

The next section contains a block for each interaction type for the  $\theta_{ij}(\sigma,ij)((\theta_{ij}(\sigma,ij))^{1/2}, f_{ij}(\sigma,ij))$ . Each block has  $n_{BOt}$  entries with 5 entries per line.

- Line 1:  $\theta_{ij}(\sigma,ij)(r_1)$ ,  $\theta_{ij}(\sigma,ij)(r_2)$ ,  $\theta_{ij}(\sigma,ij)(r_3)$ ,  $\theta_{ij}(\sigma,ij)(r_4)$ ,  $\theta_{ij}(\sigma,ij)(r_5)$  (for the  $e\_1-e\_2$  interaction type)
- Line 2:  $\theta_{ij}(\sigma,ij)(r_6)$ ,  $\theta_{ij}(\sigma,ij)(r_7)$ ,  $\theta_{ij}(\sigma,ij)(r_8)$ ,  $\theta_{ij}(\sigma,ij)(r_9)$ ,  $\theta_{ij}(\sigma,ij)(r_{10})$  (this continues until  $n_{BOt}$ )
- ...
- Line  $n_{BOt}/5+1$ :  $\theta_{ij}(\sigma,ij)(r_1)$ ,  $\theta_{ij}(\sigma,ij)(r_2)$ ,  $\theta_{ij}(\sigma,ij)(r_3)$ ,  $\theta_{ij}(\sigma,ij)(r_4)$ ,  $\theta_{ij}(\sigma,ij)(r_5)$  (for the  $e\_1-e\_2$  interaction type)

The next section contains a block of  $N$  lines for  $e\_1-e\_N$

- Line 1:  $\delta^\mu$  (for  $e\_1$ )
- Line 2:  $\delta^\mu$  (for  $e\_2$  and repeats to  $e\_N$ )

The last section contains more constants for  $e_i$ - $e_j$  interactions with  $i=0 \rightarrow N$ ,  $j=i \rightarrow N$

- Line 1:  $(A_{ij})^{(\mu * \nu)}$  (for  $e_1$ - $e_1$ )
  - Line 2:  $(A_{ij})^{(\mu * \nu)}$  (for  $e_1$ - $e_2$  and repeats as above)
- 

**Angular spline table file format:**

The parameters/coefficients format for the BOP potentials input file containing pre-tabulated functions of  $g$  is given below with variables matching the formulation of Ward (*Ward*). This format also assumes the angular functions have the formulation of (*Zhou*).

- Line 1: # elements  $N$

The first line is followed by  $N$  lines containing the atomic number, mass, and element symbol of each element.

Following the definition of the elements several global variables for the tabulated functions are given.

- Line 1:  $nr$ ,  $ntheta$ ,  $nBOt$  ( $nr$  is the number of divisions the radius is broken into for function tables and **MUST** be a factor of 5;  $ntheta$  is the power of the spline used to fit the angular function;  $nBOt$  is the number of divisions for the tabulated values of  $THETA_{(S,ij)}$ )
- Line 2:  $\delta_1$ - $\delta_7$  (if all are not used in the particular formulation, set unused values to 0.0)

Following this  $N$  lines for  $e_1$ - $e_N$  containing  $p_{\pi}$ .

- Line 3:  $p_{\pi}$  (for  $e_1$ )
- Line 4:  $p_{\pi}$  (for  $e_2$  and continues to  $e_N$ )

The next section contains several pair constants for the number of interaction types  $e_i$ - $e_j$ , with  $i=1 \rightarrow N$ ,  $j=i \rightarrow N$

- Line 1:  $r_{cut}$  (for  $e_1$ - $e_1$  interactions)
- Line 2:  $c_{\sigma}$ ,  $a_{\sigma}$ ,  $c_{\pi}$ ,  $a_{\pi}$
- Line 3:  $\delta_{\sigma}$ ,  $\delta_{\pi}$
- Line 4:  $f_{\sigma}$ ,  $k_{\sigma}$ ,  $\delta_3$  (This  $\delta_3$  is similar to that of the previous section but is interaction type dependent)

The next section contains a line for each three body interaction type  $e_j$ - $e_i$ - $e_k$  with  $i=0 \rightarrow N$ ,  $j=0 \rightarrow N$ ,  $k=j \rightarrow N$

- Line 1:  $g_0$ ,  $g_1$ ,  $g_2 \dots$  (These are coefficients for the angular spline of the  $g_{(\sigma,ijk)}(THETA_{ijk})$  for  $e_1$ - $e_1$ - $e_1$  interaction. The function can contain up to 10 term thus 10 constants. The first line can contain up to five constants. If the spline has more than five terms the second line will contain the remaining constants The following lines will then contain the constants for the remaining  $g_0$ ,  $g_1$ ,  $g_2 \dots$  (for  $e_1$ - $e_1$ - $e_2$ ) and the other three body interactions)

The rest of the table has the same structure as the previous section (see above).

---

**Angular no-spline table file format:**

The parameters/coefficients format for the BOP potentials input file containing pre-tabulated functions of  $g$  is given below with variables matching the formulation of Ward (*Ward*). This format also assumes the angular functions have the formulation of (*Zhou*).

- Line 1: # elements  $N$

The first two lines are followed by  $N$  lines containing the atomic number, mass, and element symbol of each element.

Following the definition of the elements several global variables for the tabulated functions are given.

---



- Line 1: nr, ntheta, nBOt (nr is the number of divisions the radius is broken into for function tables and MUST be a factor of 5; ntheta is the number of divisions for the tabulated values of the g angular function; nBOt is the number of divisions for the tabulated values of THETA\_(S,ij))
- Line 2: delta\_1-delta\_7 (if all are not used in the particular formulation, set unused values to 0.0)

Following this N lines for e\_1-e\_N containing p\_pi.

- Line 3: p\_pi (for e\_1)
- Line 4: p\_pi (for e\_2 and continues to e\_N)

The next section contains several pair constants for the number of interaction types e\_i-e\_j, with i=1->N, j=i->N

- Line 1: r\_cut (for e\_1-e\_1 interactions)
- Line 2: c\_sigma, a\_sigma, c\_pi, a\_pi
- Line 3: delta\_sigma, delta\_pi
- Line 4: f\_sigma, k\_sigma, delta\_3 (This delta\_3 is similar to that of the previous section but is interaction type dependent)

The next section contains a line for each three body interaction type e\_j-e\_i-e\_k with i=0->N, j=0->N, k=j->N

- Line 1: g(theta1), g(theta2), g(theta3), g(theta4), g(theta5) (for the e\_1-e\_1-e\_1 interaction type)
- Line 2: g(theta6), g(theta7), g(theta8), g(theta9), g(theta10) (this continues until ntheta)
- ...
- Line ntheta/5+1: g(theta1), g(theta2), g(theta3), g(theta4), g(theta5), (for the e\_1-e\_1-e\_2 interaction type)

The rest of the table has the same structure as the previous section (see above).

#### 19.14.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* mix, shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the pair\_style and pair\_coeff commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

#### 19.14.5 Restrictions

These pair styles are part of the MANYBODY package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These pair potentials require the *newton* setting to be “on” for pair interactions.

Pair style bop is not compatible with being used as a sub-style with doc:*hybrid pair styles <pair\_hybrid>*. Pair style bop is also not compatible with *multi-cutoff neighbor lists* or *multi-cutoff communication*.

The .bop.table potential files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use the BOP potential with any LAMMPS units, but you would need to create your own BOP potential file with coefficients listed in the appropriate units if your simulation does not use “metal” units.

### 19.14.6 Related commands

*pair\_coeff*

### 19.14.7 Default

non-tabulated potential file, *a\_0* is non-zero.

---

**(Pettifor\_1)** D.G. Pettifor and I.I. Oleinik, Phys. Rev. B, 59, 8487 (1999).

**(Pettifor\_2)** D.G. Pettifor and I.I. Oleinik, Phys. Rev. Lett., 84, 4124 (2000).

**(Pettifor\_3)** D.G. Pettifor and I.I. Oleinik, Phys. Rev. B, 65, 172103 (2002).

**(Murdick)** D.A. Murdick, X.W. Zhou, H.N.G. Wadley, D. Nguyen-Manh, R. Drautz, and D.G. Pettifor, Phys. Rev. B, 73, 45206 (2006).

**(Ward)** D.K. Ward, X.W. Zhou, B.M. Wong, F.P. Doty, and J.A. Zimmerman, Phys. Rev. B, 85, 115206 (2012).

**(Zhou)** X.W. Zhou, D.K. Ward, M. Foster (TBP).

## 19.15 *pair\_style born* command

Accelerator Variants: *born/omp*, *born/gpu*

## 19.16 *pair\_style born/coul/long* command

Accelerator Variants: *born/coul/long/gpu*, *born/coul/long/omp*

## 19.17 *pair\_style born/coul/msm* command

Accelerator Variants: *born/coul/msm/omp*

## 19.18 *pair\_style born/coul/wolf* command

Accelerator Variants: *born/coul/wolf/gpu*, *born/coul/wolf/omp*

## 19.19 *pair\_style born/coul/dsf* command

### 19.19.1 Syntax

<code>pair_style</code> style args
------------------------------------

- style = *born* or *born/coul/long* or *born/coul/msm* or *born/coul/wolf*
- args = list of arguments for a particular style

*born* args = cutoff  
 cutoff = global cutoff for non-Coulombic interactions (distance units)  
*born/coul/long* args = cutoff (cutoff2)  
 cutoff = global cutoff for non-Coulombic (and Coulombic if only 1 arg) (distance units)  
 cutoff2 = global cutoff for Coulombic (optional) (distance units)  
*born/coul/msm* args = cutoff (cutoff2)  
 cutoff = global cutoff for non-Coulombic (and Coulombic if only 1 arg) (distance units)  
 cutoff2 = global cutoff for Coulombic (optional) (distance units)  
*born/coul/wolf* args = alpha cutoff (cutoff2)  
 alpha = damping parameter (inverse distance units)  
 cutoff = global cutoff for non-Coulombic (and Coulombic if only 1 arg) (distance units)  
 cutoff2 = global cutoff for Coulombic (optional) (distance units)  
*born/coul/dsf* args = alpha cutoff (cutoff2)  
 alpha = damping parameter (inverse distance units)  
 cutoff = global cutoff for non-Coulombic (and Coulombic if only 1 arg) (distance units)  
 cutoff2 = global cutoff for Coulombic (distance units)

## 19.19.2 Examples

```

pair_style born 10.0
pair_coeff * * 6.08 0.317 2.340 24.18 11.51
pair_coeff 1 1 6.08 0.317 2.340 24.18 11.51

pair_style born/coul/long 10.0
pair_style born/coul/long 10.0 8.
pair_coeff * * 6.08 0.317 2.340 24.18 11.51
pair_coeff 1 1 6.08 0.317 2.340 24.18 11.51

pair_style born/coul/msm 10.0
pair_style born/coul/msm 10.0 8.0
pair_coeff * * 6.08 0.317 2.340 24.18 11.51
pair_coeff 1 1 6.08 0.317 2.340 24.18 11.51

pair_style born/coul/wolf 0.25 10.0
pair_style born/coul/wolf 0.25 10.0 9.0
pair_coeff * * 6.08 0.317 2.340 24.18 11.51
pair_coeff 1 1 6.08 0.317 2.340 24.18 11.51

pair_style born/coul/dsf 0.1 10.0 12.0
pair_coeff * * 0.0 1.00 0.00 0.00 0.00
pair_coeff 1 1 480.0 0.25 0.00 1.05 0.50
  
```

### 19.19.3 Description

The *born* style computes the Born-Mayer-Huggins or Tosi/Fumi potential described in (*Fumi and Tosi*), given by

$$E = A \exp\left(\frac{\sigma - r}{\rho}\right) - \frac{C}{r^6} + \frac{D}{r^8} \quad r < r_c$$

where  $\sigma$  is an interaction-dependent length parameter,  $\rho$  is an ionic-pair dependent length parameter, and  $r_c$  is the cutoff.

The styles with *coul/long* or *coul/msm* add a Coulombic term as described for the *lj/cut* pair styles. An additional damping factor is applied to the Coulombic term so it can be used in conjunction with the *kpace\_style* command and its *ewald* or *pppm* of *msm* option. The Coulombic cutoff specified for this style means that pairwise interactions within this distance are computed directly; interactions outside that distance are computed in reciprocal space.

If one cutoff is specified for the *born/coul/long* and *born/coul/msm* style, it is used for both the A,C,D and Coulombic terms. If two cutoffs are specified, the first is used as the cutoff for the A,C,D terms, and the second is the cutoff for the Coulombic term.

The *born/coul/wolf* style adds a Coulombic term as described for the Wolf potential in the *coul/wolf* pair style.

The *born/coul/dsf* style computes the Coulomb contribution with the damped shifted force model as in the *coul/dsf* style.

Note that these potentials are related to the *Buckingham potential*.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- A (energy units)
- $\rho$  (distance units)
- $\sigma$  (distance units)
- C (energy units \* distance units<sup>6</sup>)
- D (energy units \* distance units<sup>8</sup>)
- cutoff (distance units)

The second coefficient, rho, must be greater than zero.

The last coefficient is optional. If not specified, the global A,C,D cutoff specified in the *pair\_style* command is used.

For *born/coul/long*, *born/coul/wolf* and *born/coul/dsf* no Coulombic cutoff can be specified for an individual I,J type pair. All type pairs use the same global Coulombic cutoff specified in the *pair\_style* command.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.19.4 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles do not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

These styles support the *pair\_modify* shift option for the energy of the exp(),  $1/r^6$ , and  $1/r^8$  portion of the pair interaction.

The *born/coul/long* pair style supports the *pair\_modify* table option to tabulate the short-range portion of the long-range Coulombic interaction.

These styles support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

These styles write their information to binary *restart* files, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

These styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

---

### 19.19.5 Restrictions

The *born/coul/long* style is part of the KSPACE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The *born/coul/dsf* and *born/coul/wolf* pair styles are part of the EXTRA-PAIR package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.19.6 Related commands

*pair\_coeff*, *pair\_style buck*

### 19.19.7 Default

none

---

Fumi and Tosi, J Phys Chem Solids, 25, 31 (1964), Fumi and Tosi, J Phys Chem Solids, 25, 45 (1964).

## 19.20 pair\_style bpm/spring command

### 19.20.1 Syntax

```
pair_style bpm/spring
```

## 19.20.2 Examples

```
pair_style bpm/spring
pair_coeff * * 1.0 1.0 1.0
pair_coeff 1 1 1.0 1.0 1.0
```

## 19.20.3 Description

Style *bpm/spring* computes pairwise forces with the formula

$$F = k(r - r_c)$$

where  $k$  is a stiffness and  $r_c$  is the cutoff length. An additional damping force is also applied to interacting particles. The force is proportional to the difference in the normal velocity of particles

$$F_D = -\gamma w(\hat{r} \bullet \vec{v})$$

where  $\gamma$  is the damping strength,  $\hat{r}$  is the radial normal vector,  $\vec{v}$  is the velocity difference between the two particles, and  $w$  is a smoothing factor. This smoothing factor is constructed such that damping forces go to zero as particles come out of contact to avoid discontinuities. It is given by

$$w = 1.0 - \left(\frac{r}{r_c}\right)^8.$$

This pair style is designed for use in a spring-based bonded particle model. It mirrors the construction of the *bpm/spring* bond style.

This pair interaction is always applied to pairs of non-bonded particles that are within the interaction distance. For pairs of bonded particles that are within the interaction distance, there is the option to either include this pair interaction and overlay the pair force over the bond force or to exclude this pair interaction such that the two particles only interact via the bond force. See discussion of the *overlay/pair* option for BPM bond styles and the *special\_bonds* command in the *:doc: how to <Howto\_BPM>* page on BPMs for more details.

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $k$  (force/distance units)
- $r_c$  (distance units)
- $\gamma$  (force/velocity units)

---

## 19.20.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs  $I, J$  and  $I \neq J$ , the  $A$  coefficient and cutoff distance for this pair style can be mixed.  $A$  is always mixed via a *geometric* rule. The cutoff is mixed according to the *pair\_modify* mix value. The default mix value is *geometric*. See the “*pair\_modify*” command for details.

This pair style does not support the *pair\_modify* shift option, since the pair interaction goes to 0.0 at the cutoff.

The *pair\_modify* table and tail options are not relevant for this pair style.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

## 19.20.5 Restrictions

none

## 19.20.6 Related commands

*pair\_coeff*, *bond bpm/spring*

## 19.20.7 Default

none

## 19.21 pair\_style brownian command

Accelerator Variants: *brownian/omp*

## 19.22 pair\_style brownian/poly command

Accelerator Variants: *brownian/poly/omp*

### 19.22.1 Syntax

```
pair_style style mu flaglog flagfld cutinner cutoff t_target seed flagHI flagVF
```

- style = *brownian* or *brownian/poly*
- mu = dynamic viscosity (dynamic viscosity units)
- flaglog = 0/1 log terms in the lubrication approximation on/off
- flagfld = 0/1 to include/exclude Fast Lubrication Dynamics effects
- cutinner = inner cutoff distance (distance units)
- cutoff = outer cutoff for interactions (distance units)
- t\_target = target temp of the system (temperature units)
- seed = seed for the random number generator (positive integer)
- flagHI (optional) = 0/1 to include/exclude 1/r hydrodynamic interactions
- flagVF (optional) = 0/1 to include/exclude volume fraction corrections in the long-range isotropic terms

## 19.22.2 Examples

```
pair_style brownian 1.5 1 1 2.01 2.5 2.0 5878567 # (assuming radius = 1)
pair_coeff 1 1 2.05 2.8
pair_coeff * *
```

## 19.22.3 Description

Styles *brownian* and *brownian/poly* compute Brownian forces and torques on finite-size spherical particles. The former requires monodisperse spherical particles; the latter allows for polydisperse spherical particles.

These pair styles are designed to be used with either the *pair\_style lubricate* or *pair\_style lubricateU* commands to provide thermostating when dissipative lubrication forces are acting. Thus the parameters *mu*, *flaglog*, *flagfld*, *cutinner*, and *cutoff* should be specified consistent with the settings in the lubrication pair styles. For details, refer to either of the lubrication pair styles.

The *t\_target* setting is used to specify the target temperature of the system. The random number *seed* is used to generate random numbers for the thermostating procedure.

The *flagHI* and *flagVF* settings are optional. Neither should be used, or both must be defined.

---

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- *cutinner* (distance units)
- *cutoff* (distance units)

The two coefficients are optional. If neither is specified, the two cutoffs specified in the *pair\_style* command are used. Otherwise both must be specified.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---



### 19.22.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the two cutoff distances for this pair style can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

This pair style does not support the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.22.5 Restrictions

These styles are part of the COLLOID package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Only spherical monodisperse particles are allowed for pair\_style brownian.

Only spherical particles are allowed for pair\_style brownian/poly.

### 19.22.6 Related commands

*pair\_coeff*, *pair\_style lubricate*, *pair\_style lubricateU*

### 19.22.7 Default

The default settings for the optional args are flagHI = 1 and flagVF = 1.

## 19.23 pair\_style buck command

Accelerator Variants: *buck/gpu*, *buck/intel*, *buck/kk*, *buck/omp*

## 19.24 pair\_style buck/coul/cut command

Accelerator Variants: *buck/coul/cut/gpu*, *buck/coul/cut/intel*, *buck/coul/cut/kk*, *buck/coul/cut/omp*

## 19.25 pair\_style buck/coul/long command

Accelerator Variants: *buck/coul/long/gpu*, *buck/coul/long/intel*, *buck/coul/long/kk*, *buck/coul/long/omp*

## 19.26 pair\_style buck/coul/msm command

Accelerator Variants: *buck/coul/msm/omp*

### 19.26.1 Syntax

```
pair_style style args
```

- style = *buck* or *buck/coul/cut* or *buck/coul/long* or *buck/coul/msm*
- args = list of arguments for a particular style

*buck* args = cutoff

cutoff = global cutoff for Buckingham interactions (distance units)

*buck/coul/cut* args = cutoff (cutoff2)

cutoff = global cutoff for Buckingham (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

*buck/coul/long* args = cutoff (cutoff2)

cutoff = global cutoff for Buckingham (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

*buck/coul/msm* args = cutoff (cutoff2)

cutoff = global cutoff for Buckingham (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

### 19.26.2 Examples

```
pair_style buck 2.5
pair_coeff * * 100.0 1.5 200.0
pair_coeff * * 100.0 1.5 200.0 3.0

pair_style buck/coul/cut 10.0
pair_style buck/coul/cut 10.0 8.0
pair_coeff * * 100.0 1.5 200.0
pair_coeff 1 1 100.0 1.5 200.0 9.0
pair_coeff 1 1 100.0 1.5 200.0 9.0 8.0

pair_style buck/coul/long 10.0
pair_style buck/coul/long 10.0 8.0
pair_coeff * * 100.0 1.5 200.0
pair_coeff 1 1 100.0 1.5 200.0 9.0

pair_style buck/coul/msm 10.0
pair_style buck/coul/msm 10.0 8.0
pair_coeff * * 100.0 1.5 200.0
pair_coeff 1 1 100.0 1.5 200.0 9.0
```

### 19.26.3 Description

The *buck* style computes a Buckingham potential ( $\exp/6$  instead of Lennard-Jones  $12/6$ ) given by

$$E = Ae^{-r/\rho} - \frac{C}{r^6} \quad r < r_c$$

where  $\rho$  is an ionic-pair dependent length parameter, and  $r_c$  is the cutoff on both terms.

The styles with *coul/cut* or *coul/long* or *coul/msm* add a Coulombic term as described for the *lj/cut* pair styles. For *buck/coul/long* and *buc/coul/msm*, an additional damping factor is applied to the Coulombic term so it can be used in conjunction with the *kpace\_style* command and its *ewald* or *pppm* or *msm* option. The Coulombic cutoff specified for this style means that pairwise interactions within this distance are computed directly; interactions outside that distance are computed in reciprocal space.

If one cutoff is specified for the *born/coul/cut* and *born/coul/long* and *born/coul/msm* styles, it is used for both the A,C and Coulombic terms. If two cutoffs are specified, the first is used as the cutoff for the A,C terms, and the second is the cutoff for the Coulombic term.

Note that these potentials are related to the *Born-Mayer-Huggins potential*.

---

**Note:** For all these pair styles, the terms with A and C are always cutoff. The additional Coulombic term can be cutoff or long-range (no cutoff) depending on whether the style name includes *coul/cut* or *coul/long* or *coul/msm*. If you wish the  $C/r^6$  term to be long-range (no cutoff), then see the *pair\_style buck/long/coul/long* command.

---

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- A (energy units)
- $\rho$  (distance units)
- C (energy-distance<sup>6</sup> units)
- cutoff (distance units)
- cutoff2 (distance units)

The second coefficient,  $\rho$ , must be greater than zero. The coefficients A,  $\rho$ , and C can be written as analytical expressions of  $\epsilon$  and  $\sigma$ , in analogy to the Lennard-Jones potential (*Khrapak*).

The latter 2 coefficients are optional. If not specified, the global A,C and Coulombic cutoffs are used. If only one cutoff is specified, it is used as the cutoff for both A,C and Coulombic interactions for this type pair. If both coefficients are specified, they are used as the A,C and Coulombic cutoffs for this type pair. You cannot specify 2 cutoffs for style *buck*, since it has no Coulombic terms. For *buck/coul/long* only the LJ cutoff can be specified since a Coulombic cutoff cannot be specified for an individual I,J type pair. All type pairs use the same global Coulombic cutoff specified in the *pair\_style* command.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.26.4 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles do not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

These styles support the *pair\_modify* shift option for the energy of the  $\exp()$  and  $1/r^6$  portion of the pair interaction.

The *buck/coul/long* pair style supports the *pair\_modify* table option to tabulate the short-range portion of the long-range Coulombic interaction.

These styles support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure for the A,C terms in the pair interaction.

These styles write their information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

These styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

### 19.26.5 Restrictions

The *buck/coul/long* style is part of the KSPACE package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.26.6 Related commands

*pair\_coeff*, *pair\_style born*

### 19.26.7 Default

none

(**Khrapak**) Khrapak, Chaudhuri, and Morfill, J Chem Phys, 134, 054120 (2011).

## 19.27 pair\_style buck6d/coul/gauss/dsf command

## 19.28 pair\_style buck6d/coul/gauss/long command

### 19.28.1 Syntax

```
pair_style style args
```

- style = *buck6d/coul/gauss/dsf* or *buck6d/coul/gauss/long*
- args = list of arguments for a particular style

```

buck6d/coul/gauss/dsf args = smooth cutoff (cutoff2)
  smooth = smoothing onset within Buckingham cutoff (ratio)
  cutoff = global cutoff for Buckingham (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)
buck6d/coul/gauss/long args = smooth smooth2 cutoff (cutoff2)
  smooth = smoothing onset within Buckingham cutoff (ratio)
  smooth2 = smoothing onset within Coulombic cutoff (ratio)
  cutoff = global cutoff for Buckingham (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)

```

## 19.28.2 Examples

```

pair_style buck6d/coul/gauss/dsf 0.9000 12.0000
pair_coeff 1 1 1030. 3.061 457.179 4.521 0.608

pair_style buck6d/coul/gauss/long 0.9000 1.0000 12.0000
pair_coeff 1 1 1030. 3.061 457.179 4.521 0.608

```

## 19.28.3 Description

The *buck6d/coul/gauss* styles evaluate vdW and Coulomb interactions following the MOF-FF force field after (*Schmid*). The vdW term of the *buck6d* styles computes a dispersion damped Buckingham potential:

$$E = Ae^{-\kappa r} - \frac{C}{r^6} \cdot \frac{1}{1 + Dr^{14}} \quad r < r_c$$

where A and C are a force constant,  $\kappa$  is an ionic-pair dependent reciprocal length parameter, D is a dispersion correction parameter, and the cutoff  $r_c$  truncates the interaction distance. The first term in the potential corresponds to the Buckingham repulsion term and the second term to the dispersion attraction with a damping correction analog to the Grimme correction used in DFT. The latter corrects for artifacts occurring at short distances which become an issue for soft vdW potentials.

The *buck6d* styles include a smoothing function which is invoked according to the global smoothing parameter within the specified cutoff. Hereby a parameter of i.e. 0.9 invokes the smoothing within 90% of the cutoff. No smoothing is applied at a value of 1.0. For the *gauss/dsf* style this smoothing is only applicable for the dispersion damped Buckingham potential. For the *gauss/long* styles the smoothing function can also be invoked for the real space coulomb interactions which enforce continuous energies and forces at the cutoff.

Both styles *buck6d/coul/gauss/dsf* and *buck6d/coul/gauss/long* evaluate a Coulomb potential using spherical Gaussian type charge distributions which effectively dampen electrostatic interactions for high charges at close distances. The electrostatic potential is thus evaluated as:

$$E = \frac{C_{q_i q_j}}{\epsilon r_{ij}} \operatorname{erf}(\alpha_{ij} r_{ij}) \quad r < r_c$$

where C is an energy-conversion constant,  $q_i$  and  $q_j$  are the charges on the 2 atoms, epsilon is the dielectric constant which can be set by the *dielectric* command,  $\alpha$  is the ion pair dependent damping parameter and erf() is the error-function. The cutoff  $r_c$  truncates the interaction distance.

The style *buck6d/coul/gauss/dsf* computes the Coulomb interaction via the damped shifted force model described in (*Fennell*) approximating an Ewald sum similar to the *pair coul/dsf* styles. In *buck6d/coul/gauss/long* an additional damping factor is applied to the Coulombic term so it can be used in conjunction with the *kpace\_style* command and its *ewald* or *pppm* options. The Coulombic cutoff in this case separates the real and reciprocal space evaluation of the Ewald sum.

If one cutoff is specified it is used for both the vdW and Coulomb terms. If two cutoffs are specified, the first is used as the cutoff for the vdW terms, and the second is the cutoff for the Coulombic term.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- A (energy units)
- $\rho$  (distance<sup>-1</sup> units)
- C (energy-distance<sup>6</sup> units)
- D (distance<sup>14</sup> units)
- $\alpha$  (distance<sup>-1</sup> units)
- cutoff (distance units)

The second coefficient,  $\rho$ , must be greater than zero. The latter coefficient is optional. If not specified, the global vdW cutoff is used.

---

#### 19.28.4 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles do not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

These styles do not support the *pair\_modify* shift option for the energy. Instead the smoothing function should be applied by setting the global smoothing parameter to a value < 1.0.

These styles write their information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

#### 19.28.5 Restrictions

These styles are part of the MOFFF package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 19.28.6 Related commands

*pair\_coeff*

#### 19.28.7 Default

none

**(Schmid)** S. Bureekaew, S. Amirjalayer, M. Tafipolsky, C. Spickermann, T.K. Roy and R. Schmid, Phys. Status Solidi B, 6, 1128 (2013).

**(Fennell)** C. J. Fennell, J. D. Gezelter, J Chem Phys, 124, 234104 (2006).

## 19.29 pair\_style buck/long/coul/long command

Accelerator Variants: *buck/long/coul/long/omp*

### 19.29.1 Syntax

```
pair_style buck/long/coul/long flag_buck flag_coul cutoff (cutoff2)
```

- `flag_buck` = *long* or *cut*  
*long* = use Kspace long-range summation for the dispersion term  $1/r^6$   
*cut* = use a cutoff
- `flag_coul` = *long* or *off*  
*long* = use Kspace long-range summation for the Coulombic term  $1/r$   
*off* = omit the Coulombic term
- `cutoff` = global cutoff for Buckingham (and Coulombic if only 1 cutoff) (distance units)
- `cutoff2` = global cutoff for Coulombic (optional) (distance units)

### 19.29.2 Examples

```
pair_style buck/long/coul/long cut off 2.5
pair_style buck/long/coul/long cut long 2.5 4.0
pair_style buck/long/coul/long long long 4.0
pair_coeff * * 1 1
pair_coeff 1 1 1 3 4
```

### 19.29.3 Description

The *buck/long/coul/long* style computes a Buckingham potential ( $\exp/6$  instead of Lennard-Jones  $12/6$ ) and Coulombic potential, given by

$$E = Ae^{-r/\rho} - \frac{C}{r^6} \quad r < r_c$$

$$E = \frac{Cq_iq_j}{\epsilon r} \quad r < r_c$$

$r_c$  is the cutoff. If one cutoff is specified in the `pair_style` command, it is used for both the Buckingham and Coulombic terms. If two cutoffs are specified, they are used as cutoffs for the Buckingham and Coulombic terms respectively.

The purpose of this pair style is to capture long-range interactions resulting from both attractive  $1/r^6$  Buckingham and Coulombic  $1/r$  interactions. This is done by use of the `flag_buck` and `flag_coul` settings. The *Ismail* paper has more details on when it is appropriate to include long-range  $1/r^6$  interactions, using this potential.

If `flag_buck` is set to *long*, no cutoff is used on the Buckingham  $1/r^6$  dispersion term. The long-range portion can be calculated by using the `kpace_style ewald/disp` or `pppm/disp` commands. The specified Buckingham cutoff then determines which portion of the Buckingham interactions are computed directly by the pair potential versus which part is computed in reciprocal space via the Kspace style. If `flag_buck` is set to *cut*, the Buckingham interactions are simply cutoff, as with `pair_style buck`.

If `flag_coul` is set to *long*, no cutoff is used on the Coulombic interactions. The long-range portion can be calculated by using any of several `kpace_style` command options such as `pppm` or `ewald`. Note that if `flag_buck` is also set to *long*,

then the *ewald/disp* or *pppm/disp* Kspace style needs to be used to perform the long-range calculations for both the Buckingham and Coulombic interactions. If *flag\_coul* is set to *off*, Coulombic interactions are not computed.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- A (energy units)
- rho (distance units)
- C (energy-distance<sup>6</sup> units)
- cutoff (distance units)
- cutoff2 (distance units)

The second coefficient, rho, must be greater than zero.

The latter 2 coefficients are optional. If not specified, the global Buckingham and Coulombic cutoffs specified in the *pair\_style* command are used. If only one cutoff is specified, it is used as the cutoff for both Buckingham and Coulombic interactions for this type pair. If both coefficients are specified, they are used as the Buckingham and Coulombic cutoffs for this type pair. Note that if you are using *flag\_buck* set to *long*, you cannot specify a Buckingham cutoff for an atom type pair, since only one global Buckingham cutoff is allowed. Similarly, if you are using *flag\_coul* set to *long*, you cannot specify a Coulombic cutoff for an atom type pair, since only one global Coulombic cutoff is allowed.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.29.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This pair style supports the *pair\_modify* shift option for the energy of the  $\exp()$  and  $1/r^6$  portion of the pair interaction, assuming *flag\_buck* is *cut*.

This pair style does not support the *pair\_modify* shift option for the energy of the Buckingham portion of the pair interaction.

This pair style supports the *pair\_modify* table and table/disp options since they can tabulate the short-range portion of the long-range Coulombic and dispersion interactions.

This pair style write its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style supports the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command, meaning the pairwise forces can be partitioned by distance at different levels of the rRESPA hierarchy. See the *run\_style* command for details.

---



### 19.29.5 Restrictions

This style is part of the KSPACE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.29.6 Related commands

*pair\_coeff*

### 19.29.7 Default

none

---

(Ismail) Ismail, Tsige, In 't Veld, Grest, Molecular Physics (accepted) (2007).

## 19.30 pair\_style lj/charmm/coul/charmm command

Accelerator Variants: *lj/charmm/coul/charmm/gpu*, *lj/charmm/coul/charmm/intel*, *lj/charmm/coul/charmm/kk*, *lj/charmm/coul/charmm/omp*

## 19.31 pair\_style lj/charmm/coul/charmm/implicit command

Accelerator Variants: *lj/charmm/coul/charmm/implicit/kk*, *lj/charmm/coul/charmm/implicit/omp*

## 19.32 pair\_style lj/charmm/coul/long command

Accelerator Variants: *lj/charmm/coul/long/gpu*, *lj/charmm/coul/long/intel*, *lj/charmm/coul/long/kk*, *lj/charmm/coul/long/opt*, *lj/charmm/coul/long/omp*

## 19.33 pair\_style lj/charmm/coul/msm command

Accelerator Variants: *lj/charmm/coul/msm/omp*

## 19.34 pair\_style lj/charmmfsw/coul/charmmfsh command

## 19.35 pair\_style lj/charmmfsw/coul/long command

### 19.35.1 Syntax

<code>pair_style</code> style args
------------------------------------

- style = *lj/charmm/coul/charmm* or *lj/charmm/coul/charmm/implicit* or *lj/charmm/coul/long* or *lj/charmm/coul/msm* or *lj/charmmfsw/coul/charmmfsh* or *lj/charmmfsw/coul/long*
- args = list of arguments for a particular style

```
lj/charmm/coul/charmm args = inner outer (inner2) (outer2)
  inner, outer = global switching cutoffs for Lennard Jones (and Coulombic if only 2
↳args)
  inner2, outer2 = global switching cutoffs for Coulombic (optional)
lj/charmm/coul/charmm/implicit args = inner outer (inner2) (outer2)
  inner, outer = global switching cutoffs for LJ (and Coulombic if only 2 args)
  inner2, outer2 = global switching cutoffs for Coulombic (optional)
lj/charmm/coul/long args = inner outer (cutoff)
  inner, outer = global switching cutoffs for LJ (and Coulombic if only 2 args)
  cutoff = global cutoff for Coulombic (optional, outer is Coulombic cutoff if only 2
↳args)
lj/charmm/coul/msm args = inner outer (cutoff)
  inner, outer = global switching cutoffs for LJ (and Coulombic if only 2 args)
  cutoff = global cutoff for Coulombic (optional, outer is Coulombic cutoff if only 2
↳args)
lj/charmmfsw/coul/charmmfsh args = inner outer (cutoff)
  inner, outer = global cutoffs for LJ (and Coulombic if only 2 args)
  cutoff = global cutoff for Coulombic (optional, outer is Coulombic cutoff if only 2
↳args)
lj/charmmfsw/coul/long args = inner outer (cutoff)
  inner, outer = global cutoffs for LJ (and Coulombic if only 2 args)
  cutoff = global cutoff for Coulombic (optional, outer is Coulombic cutoff if only 2
↳args)
```

## 19.35.2 Examples

```
pair_style lj/charmm/coul/charmm 8.0 10.0
pair_style lj/charmm/coul/charmm 8.0 10.0 7.0 9.0
pair_style lj/charmmfsw/coul/charmmfsh 10.0 12.0
pair_style lj/charmmfsw/coul/charmmfsh 10.0 12.0 9.0
pair_coeff * * 100.0 2.0
pair_coeff 1 1 100.0 2.0 150.0 3.5

pair_style lj/charmm/coul/charmm/implicit 8.0 10.0
pair_style lj/charmm/coul/charmm/implicit 8.0 10.0 7.0 9.0
pair_coeff * * 100.0 2.0
pair_coeff 1 1 100.0 2.0 150.0 3.5

pair_style lj/charmm/coul/long 8.0 10.0
pair_style lj/charmm/coul/long 8.0 10.0 9.0
pair_style lj/charmmfsw/coul/long 8.0 10.0
pair_style lj/charmmfsw/coul/long 8.0 10.0 9.0
pair_coeff * * 100.0 2.0
pair_coeff 1 1 100.0 2.0 150.0 3.5

pair_style lj/charmm/coul/msm 8.0 10.0
pair_style lj/charmm/coul/msm 8.0 10.0 9.0
pair_coeff * * 100.0 2.0
```

(continues on next page)

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```
pair_coeff 1 1 100.0 2.0 150.0 3.5
```

### 19.35.3 Description

These pair styles compute Lennard Jones (LJ) and Coulombic interactions with additional switching or shifting functions that ramp the energy and/or force smoothly to zero between an inner and outer cutoff. They are implementations of the widely used CHARMM force field used in the CHARMM MD code (and others). See (*MacKerell*) for a description of the CHARMM force field.

The styles with *charmm* (not *charmmfsw* or *charmmfsh*) in their name are the older, original LAMMPS implementations. They compute the LJ and Coulombic interactions with an energy switching function (esw, shown in the formula below as  $S(r)$ ), which ramps the energy smoothly to zero between the inner and outer cutoff. This can cause irregularities in pairwise forces (due to the discontinuous second derivative of energy at the boundaries of the switching region), which in some cases can result in detectable artifacts in an MD simulation.

The newer styles with *charmmfsw* or *charmmfsh* in their name replace the energy switching with force switching (fsw) and force shifting (fsh) functions, for LJ and Coulombic interactions respectively. These follow the formulas and description given in (*Steinbach*) and (*Brooks*) to minimize these artifacts.

---

**Note:** The newer *charmmfsw* or *charmmfsh* styles were released in March 2017. We recommend they be used instead of the older *charmm* styles. This includes the newer *dihedral\_style charmmfsw* command. Eventually code from the new styles will propagate into the related pair styles (e.g. implicit, accelerator, free energy variants).

---



---

**Note:** The newest CHARMM pair styles reset the Coulombic energy conversion factor used internally in the code, from the LAMMPS value to the CHARMM value, as if it were effectively a parameter of the force field. This is because the CHARMM code uses a slightly different value for the this conversion factor in *real units* (Kcal/mole), namely CHARMM = 332.0716, LAMMPS = 332.06371. This is to enable more precise agreement by LAMMPS with the CHARMM force field energies and forces, when using one of these two CHARMM pair styles.

---

$$\begin{aligned}
 E &= LJ(r) & r < r_{\text{in}} \\
 &= S(r) * LJ(r) & r_{\text{in}} < r < r_{\text{out}} \\
 &= 0 & r > r_{\text{out}} \\
 E &= C(r) & r < r_{\text{in}} \\
 &= S(r) * C(r) & r_{\text{in}} < r < r_{\text{out}} \\
 &= 0 & r > r_{\text{out}}
 \end{aligned}$$

$$LJ(r) = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right]$$

$$C(r) = \frac{Cq_i q_j}{\epsilon r}$$

$$S(r) = \frac{[r_{\text{out}}^2 - r^2]^2 [r_{\text{out}}^2 + 2r^2 - 3r_{\text{in}}^2]}{[r_{\text{out}}^2 - r_{\text{in}}^2]^3}$$

where  $S(r)$  is the energy switching function mentioned above for the *charmm* styles. See the (*Steinbach*) paper for the functional forms of the force switching and force shifting functions used in the *charmmfsw* and *charmmfsh* styles.

When using the *lj/charmm/coul/charmm* styles, both the LJ and Coulombic terms require an inner and outer cutoff. They can be the same for both formulas or different depending on whether 2 or 4 arguments are used in the pair\_style

command. For the *lj/charmmfsw/coul/charmmfsh* style, the LJ term requires both an inner and outer cutoff, while the Coulombic term requires only one cutoff. If the Coulombic cutoff is not specified (2 instead of 3 arguments), the LJ outer cutoff is used for the Coulombic cutoff. In all cases where an inner and outer cutoff are specified, the inner cutoff distance must be less than the outer cutoff. It is typical to make the difference between the inner and outer cutoffs about 2.0 Angstroms.

Style *lj/charmm/coul/charmm/implicit* computes the same formulas as style *lj/charmm/coul/charmm* except that an additional  $1/r$  term is included in the Coulombic formula. The Coulombic energy thus varies as  $1/r^2$ . This is effectively a distance-dependent dielectric term which is a simple model for an implicit solvent with additional screening. It is designed for use in a simulation of an unsolvated biomolecule (no explicit water molecules).

Styles *lj/charmm/coul/long* and *lj/charmm/coul/msm* compute the same formulas as style *lj/charmm/coul/charmm* and style *lj/charmmfsw/coul/long* computes the same formulas as style *lj/charmmfsw/coul/charmmfsh*, except that an additional damping factor is applied to the Coulombic term, so it can be used in conjunction with the *kspc\_style* command and its *ewald* or *pppm* or *msm* option. Only one Coulombic cutoff is specified for these styles; if only 2 arguments are used in the *pair\_style* command, then the outer LJ cutoff is used as the single Coulombic cutoff. The Coulombic cutoff specified for these styles means that pairwise interactions within this distance are computed directly; interactions outside that distance are computed in reciprocal space.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- $\epsilon_{14}$  (energy units)
- $\sigma_{14}$  (distance units)

Note that  $\sigma$  is defined in the LJ formula as the zero-crossing distance for the potential, not as the energy minimum at  $2^{1/6}\sigma$ .

The latter 2 coefficients are optional. If they are specified, they are used in the LJ formula between 2 atoms of these types which are also first and fourth atoms in any dihedral. No cutoffs are specified because the CHARMM force field does not allow varying cutoffs for individual atom pairs; all pairs use the global cutoff(s) specified in the *pair\_style* command.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* [command-line switch](#) when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 19.35.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon, sigma, epsilon\_14, and sigma\_14 coefficients for all of the lj/charmm pair styles can be mixed. The default mix value is *arithmetic* to coincide with the usual settings for the CHARMM force field. See the “pair\_modify” command for details.

None of the *lj/charmm* or *lj/charmmfsw* pair styles support the *pair\_modify* shift option, since the Lennard-Jones portion of the pair interaction is smoothed to 0.0 at the cutoff.

The *lj/charmm/coul/long* and *lj/charmmfsw/coul/long* styles support the *pair\_modify* table option since they can tabulate the short-range portion of the long-range Coulombic interaction.

None of the *lj/charmm* or *lj/charmmfsw* pair styles support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure, since the Lennard-Jones portion of the pair interaction is smoothed to 0.0 at the cutoff.

All of the *lj/charmm* and *lj/charmmfsw* pair styles write their information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

The *lj/charmm/coul/long* and *lj/charmmfsw/coul/long* pair styles support the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command, meaning the pairwise forces can be partitioned by distance at different levels of the rRESPA hierarchy. The other styles only support the *pair* keyword of run\_style respa. See the *run\_style* command for details.

---

### 19.35.5 Restrictions

All the styles with *coul/charmm* or *coul/charmmfsh* styles are part of the MOLECULE package. All the styles with *coul/long* style are part of the KSPACE package. They are only enabled if LAMMPS was built with those packages. See the *Build package* doc page for more info.

### 19.35.6 Related commands

*pair\_coeff*

### 19.35.7 Default

none

---

**(Brooks)** Brooks, et al, J Comput Chem, 30, 1545 (2009).

**(MacKerell)** MacKerell, Bashford, Bellott, Dunbrack, Evanseck, Field, Fischer, Gao, Guo, Ha, et al, J Phys Chem, 102, 3586 (1998).

**(Steinbach)** Steinbach, Brooks, J Comput Chem, 15, 667 (1994).

## 19.36 pair\_style lj/class2 command

Accelerator Variants: *lj/class2/gpu*, *lj/class2/kk*, *lj/class2/omp*

## 19.37 pair\_style lj/class2/coul/cut command

Accelerator Variants: *lj/class2/coul/cut/kk*, *lj/class2/coul/cut/omp*

## 19.38 pair\_style lj/class2/coul/long command

Accelerator Variants: *lj/class2/coul/long/gpu*, *lj/class2/coul/long/kk*, *lj/class2/coul/long/omp*

### 19.38.1 Syntax

```
pair_style style args
```

- style = *lj/class2* or *lj/class2/coul/cut* or *lj/class2/coul/long*
- args = list of arguments for a particular style

*lj/class2* args = cutoff

cutoff = global cutoff for class 2 interactions (distance units)

*lj/class2/coul/cut* args = cutoff (cutoff2)

cutoff = global cutoff for class 2 (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

*lj/class2/coul/long* args = cutoff (cutoff2)

cutoff = global cutoff for class 2 (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

### 19.38.2 Examples

```
pair_style lj/class2 10.0
pair_coeff * * 100.0 2.5
pair_coeff 1 2* 100.0 2.5 9.0

pair_style lj/class2/coul/cut 10.0
pair_style lj/class2/coul/cut 10.0 8.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0
pair_coeff 1 1 100.0 3.5 9.0 9.0

pair_style lj/class2/coul/long 10.0
pair_style lj/class2/coul/long 10.0 8.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0
```

### 19.38.3 Description

The *lj/class2* styles compute a 6/9 Lennard-Jones potential given by

$$E = \epsilon \left[ 2 \left( \frac{\sigma}{r} \right)^9 - 3 \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_c$$

$r_c$  is the cutoff.

The *lj/class2/coul/cut* and *lj/class2/coul/long* styles add a Coulombic term as described for the *lj/cut* pair styles.

See (*Sun*) for a description of the COMPASS class2 force field.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- cutoff1 (distance units)
- cutoff2 (distance units)

The latter 2 coefficients are optional. If not specified, the global class 2 and Coulombic cutoffs are used. If only one cutoff is specified, it is used as the cutoff for both class 2 and Coulombic interactions for this type pair. If both coefficients are specified, they are used as the class 2 and Coulombic cutoffs for this type pair. You cannot specify 2 cutoffs for style *lj/class2*, since it has no Coulombic terms.

For *lj/class2/coul/long* only the class 2 cutoff can be specified since a Coulombic cutoff cannot be specified for an individual I,J type pair. All type pairs use the same global Coulombic cutoff specified in the *pair\_style* command.

If the *pair\_coeff* command is not used to define coefficients for a particular I != J type pair, the mixing rule for  $\epsilon$  and  $\sigma$  for all class2 potentials is to use the *sixthpower* formulas documented by the *pair\_modify* command. The *pair\_modify mix* setting is thus ignored for class2 potentials for epsilon and sigma. However it is still followed for mixing the cutoff distance.

A version of these styles with a soft core, *lj/cut/soft*, suitable for use in free energy calculations, is part of the FEP package and is documented with the *pair\_style \*/soft* styles. The version with soft core is only available if LAMMPS was built with that package. See the *Build package* page for more info.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

## 19.38.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the `lj/class2` pair styles can be mixed. Epsilon and sigma are always mixed with the value *sixthpower*. The cutoff distance is mixed by whatever option is set by the `pair_modify` command (default = `geometric`). See the “`pair_modify`” command for details.

All of the `lj/class2` pair styles support the *pair\_modify* `shift` option for the energy of the Lennard-Jones portion of the pair interaction.

The `lj/class2/coul/long` pair style does not support the *pair\_modify* `table` option since a tabulation capability has not yet been added to this potential.

All of the `lj/class2` pair styles support the *pair\_modify* `tail` option for adding a long-range tail correction to the energy and pressure of the Lennard-Jones portion of the pair interaction.

All of the `lj/class2` pair styles write their information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

Only the `lj/class2` and `lj/class2/coul/long` pair styles support the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command, meaning the pairwise forces can be partitioned by distance at different levels of the rRESPA hierarchy. The other styles only support the *pair* keyword of *run\_style respa*. See the *run\_style* command for details.

## 19.38.5 Restrictions

These styles are part of the CLASS2 package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.38.6 Related commands

*pair\_coeff*, *pair\_style \*/soft*

## 19.38.7 Default

none

---

(Sun) Sun, J Phys Chem B 102, 7338-7364 (1998).

## 19.39 pair\_style colloid command

Accelerator Variants: *colloid/gpu*, *colloid/omp*

### 19.39.1 Syntax

```
pair_style colloid cutoff
```

- `cutoff` = global cutoff for colloidal interactions (distance units)



## 19.39.2 Examples

```
pair_style colloid 10.0
pair_coeff * * 25 1.0 10.0 10.0
pair_coeff 1 1 144 1.0 0.0 0.0 3.0
pair_coeff 1 2 75.398 1.0 0.0 10.0 9.0
pair_coeff 2 2 39.478 1.0 10.0 10.0 25.0
```

## 19.39.3 Description

Style *colloid* computes pairwise interactions between large colloidal particles and small solvent particles using 3 formulas. A colloidal particle has a size > sigma; a solvent particle is the usual Lennard-Jones particle of size sigma.

The colloid-colloid interaction energy is given by

$$U_A = -\frac{A_{cc}}{6} \left[ \frac{2a_1a_2}{r^2 - (a_1 + a_2)^2} + \frac{2a_1a_2}{r^2 - (a_1 - a_2)^2} + \ln \left( \frac{r^2 - (a_1 + a_2)^2}{r^2 - (a_1 - a_2)^2} \right) \right]$$

$$U_R = \frac{A_{cc}}{37800} \frac{\sigma^6}{r} \left[ \frac{r^2 - 7r(a_1 + a_2) + 6(a_1^2 + 7a_1a_2 + a_2^2)}{(r - a_1 - a_2)^7} + \frac{r^2 + 7r(a_1 + a_2) + 6(a_1^2 + 7a_1a_2 + a_2^2)}{(r + a_1 + a_2)^7} - \frac{r^2 + 7r(a_1 - a_2) + 6(a_1^2 - 7a_1a_2 + a_2^2)}{(r + a_1 - a_2)^7} - \frac{r^2 - 7r(a_1 - a_2) + 6(a_1^2 - 7a_1a_2 + a_2^2)}{(r - a_1 + a_2)^7} \right]$$

$$U = U_A + U_R, \quad r < r_c$$

where  $A_{cc}$  is the Hamaker constant,  $a_1$  and  $a_2$  are the radii of the two colloidal particles, and  $r_c$  is the cutoff. This equation results from describing each colloidal particle as an integrated collection of Lennard-Jones particles of size sigma and is derived in (Everaers).

The colloid-solvent interaction energy is given by

$$U = \frac{2 a^3 \sigma^3 A_{cs}}{9(a^2 - r^2)^3} \left[ 1 - \frac{(5 a^6 + 45 a^4 r^2 + 63 a^2 r^4 + 15 r^6) \sigma^6}{15 (a - r)^6 (a + r)^6} \right], \quad r < r_c$$

where  $A_{cs}$  is the Hamaker constant,  $a$  is the radius of the colloidal particle, and  $r_c$  is the cutoff. This formula is derived from the colloid-colloid interaction, letting one of the particle sizes go to zero.

The solvent-solvent interaction energy is given by the usual Lennard-Jones formula

$$U = \frac{A_{ss}}{36} \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right], \quad r < r_c$$

with  $A_{ss}$  set appropriately, which results from letting both particle sizes go to zero.

When used in combination with *pair\_style yukawa/colloid*, the two terms become the so-called DLVO potential, which combines electrostatic repulsion and van der Waals attraction.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- A (energy units)
- $\sigma$  (distance units)
- d1 (distance units)
- d2 (distance units)
- cutoff (distance units)

A is the Hamaker energy prefactor and should typically be set as follows:

- $A_{cc} = \text{colloid/colloid} = 4\pi^2 = 39.5$
- $A_{cs} = \text{colloid/solvent} = \sqrt{A_{cc}A_{ss}}$
- $A_{ss} = \text{solvent/solvent} = 144$  (assuming  $\epsilon = 1$ , so that  $144/36 = 4$ )

$\sigma$  is the size of the solvent particle or the constituent particles integrated over in the colloidal particle and should typically be set as follows:

- $\sigma_{cc} = \text{colloid/colloid} = 1.0$
- $\sigma_{cs} = \text{colloid/solvent} = \text{arithmetic mixing between colloid } \sigma \text{ and solvent } \sigma$
- $\sigma_{ss} = \text{solvent/solvent} = 1.0$  or whatever size the solvent particle is

Thus typically  $\sigma_{cs} = 1.0$ , unless the solvent particle's size  $\neq 1.0$ .

D1 and d2 are particle diameters, so that  $d1 = 2*a1$  and  $d2 = 2*a2$  in the formulas above. Both d1 and d2 must be values  $\geq 0$ . If  $d1 > 0$  and  $d2 > 0$ , then the pair interacts via the colloid-colloid formula above. If  $d1 = 0$  and  $d2 = 0$ , then the pair interacts via the solvent-solvent formula. I.e. a d value of 0 is a Lennard-Jones particle of size  $\sigma$ . If either  $d1 = 0$  or  $d2 = 0$  and the other is larger, then the pair interacts via the colloid-solvent formula.

Note that the diameter of a particular particle type may appear in multiple `pair_coeff` commands, as it interacts with other particle types. You should insure the particle diameter is specified consistently each time it appears.

The last coefficient is optional. If not specified, the global cutoff specified in the `pair_style` command is used. However, you typically want different cutoffs for interactions between different particle sizes. E.g. if colloidal particles of diameter 10 are used with solvent particles of diameter 1, then a solvent-solvent cutoff of 2.5 would correspond to a colloid-colloid cutoff of 25. A good rule-of-thumb is to use a colloid-solvent cutoff that is half the big diameter + 4 times the small diameter. I.e.  $9 = 5 + 4$  for the colloid-solvent cutoff in this case.

---

**Note:** When using `pair_style colloid` for a mixture with 2 (or more) widely different particles sizes (e.g.  $\sigma=10$  colloids in a background  $\sigma=1$  LJ fluid), you will likely want to use these commands for efficiency: *[neighbor multi](#)* and *[comm\\_modify multi](#)*.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *[Accelerator packages](#)* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *[Build package](#)* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *[-suffix command-line switch](#)* when you invoke LAMMPS, or you can use the *[suffix](#)* command in your input script.

See the *[Accelerator packages](#)* page for more instructions on how to use the accelerated styles effectively.

---

### 19.39.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs  $I, J$  and  $I \neq J$ , the  $A$ ,  $\sigma$ ,  $d1$ , and  $d2$  coefficients and cutoff distance for this pair style can be mixed.  $A$  is an energy value mixed like a LJ epsilon.  $D1$  and  $d2$  are distance values and are mixed like  $\sigma$ . The default mix value is *geometric*. See the “pair\_modify” command for details.

This pair style supports the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.39.5 Restrictions

This style is part of the COLLOID package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Normally, this pair style should be used with finite-size particles which have a diameter, e.g. see the *atom\_style sphere* command. However, this is not a requirement, since the only definition of particle size is via the *pair\_coeff* parameters for each type. In other words, the physical radius of the particle is ignored. Thus you should insure that the  $d1, d2$  parameters you specify are consistent with the physical size of the particles of that type.

Per-particle polydispersity is not yet supported by this pair style; only per-type polydispersity is enabled via the *pair\_coeff* parameters.

### 19.39.6 Related commands

*pair\_coeff*

### 19.39.7 Default

none

---

(Everaers) Everaers, Ejtehadi, Phys Rev E, 67, 041710 (2003).

## 19.40 pair\_style comb command

Accelerator Variants: *comb/omp*

## 19.41 pair\_style comb3 command

### 19.41.1 Syntax

```
pair_style comb
pair_style comb3 keyword
```

keyword = *polar*

*polar* value = *polar\_on* or *polar\_off* = whether or not to include atomic polarization

### 19.41.2 Examples

```
pair_style comb
pair_coeff * * ../potentials/ffield.comb Si
pair_coeff * * ../potentials/ffield.comb Hf Si 0

pair_style comb3 polar_off
pair_coeff * * ../potentials/ffield.comb3 0 Cu N C O
```

### 19.41.3 Description

Style *comb* computes the second generation variable charge COMB (Charge-Optimized Many-Body) potential. Style *comb3* computes the third-generation COMB potential. These COMB potentials are described in (*COMB*) and (*COMB3*). Briefly, the total energy  $E_T$  of a system of atoms is given by

$$E_T = \sum_i [E_i^{self}(q_i)] + \sum_{j>i} [E_{ij}^{short}(r_{ij}, q_i, q_j) + E_{ij}^{Coul}(r_{ij}, q_i, q_j)] + E^{polar}(q_i, r_{ij}) + E^{vdW}(r_{ij}) + E^{barr}(q_i) + E^{corr}(r_{ij}, \theta_{jik})$$

where  $E_i^{self}$  is the self-energy of atom  $i$  (including atomic ionization energies and electron affinities),  $E_{ij}^{short}$  is the bond-order potential between atoms  $i$  and  $j$ ,  $E_{ij}^{Coul}$  is the Coulomb interactions,  $E^{polar}$  is the polarization term for organic systems (style *comb3* only),  $E^{vdW}$  is the van der Waals energy (style *comb3* only),  $E^{barr}$  is a charge barrier function, and  $E^{corr}$  are angular correction terms.

The COMB potentials (styles *comb* and *comb3*) are variable charge potentials. The equilibrium charge on each atom is calculated by the electronegativity equalization (QEq) method. See [Rick](#) for further details. This is implemented by the *fix qeq/comb* command, which should normally be specified in the input script when running a model with the COMB potential. The *fix qeq/comb* command has options that determine how often charge equilibration is performed, its convergence criterion, and which atoms are included in the calculation.

Only a single *pair\_coeff* command is used with the *comb* and *comb3* styles which specifies the COMB potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the potential file in the *pair\_coeff* command, where N is the number of LAMMPS atom types.

For example, if your LAMMPS simulation of a Si/SiO<sub>2</sub>/HfO<sub>2</sub> interface has 4 atom types, and you want the first and last to be Si, the second to be Hf, and the third to be O, and you would use the following *pair\_coeff* command:

```
pair_coeff * * ../potentials/ffield.comb Si Hf 0 Si
```

The first two arguments must be \* \* so as to span all LAMMPS atom types. The first and last Si arguments map LAMMPS atom types 1 and 4 to the Si element in the *ffield.comb* file. The second Hf argument maps LAMMPS atom type 2 to the Hf element, and the third O argument maps LAMMPS atom type 3 to the O element in the potential file.

If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *comb* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

For style *comb*, the provided potential file *ffield.comb* contains all currently-available second generation COMB parameterizations: for Si, Cu, Hf, Ti, O, their oxides and Zr, Zn and U metals. For style *comb3*, the potential file *ffield.comb3* contains all currently-available third generation COMB parameterizations: O, Cu, N, C, H, Ti, Zn and Zr. The status of the optimization of the compounds, for example Cu<sub>2</sub>O, TiN and hydrocarbons, are given in the following table:

	O	Cu	N	C	H	Ti	Zn	Zr
O	F	F	F	F	F	F	F	F
Cu	F	F	P	F	F	P	F	P
N	F	P	F	M	F	P	P	P
C	F	F	M	F	F	M	M	M
H	F	F	F	F	F	M	M	F
Ti	F	P	P	M	M	F	P	P
Zn	F	F	P	M	M	P	F	P
Zr	F	P	P	M	F	P	P	F

- F = Fully optimized
- M = Only optimized for dimer molecule
- P = in progress, but have it from mixing rule

For style *comb3*, in addition to *ffield.comb3*, a special parameter file, *lib.comb3*, that is exclusively used for C/O/H systems, will be automatically loaded if carbon atom is detected in LAMMPS input structure. This file must be in your working directory or in the directories listed in the environment variable LAMMPS\_POTENTIALS, as described on the *pair\_coeff* command doc page.

The keyword *polar* indicates whether the force field includes the atomic polarization. Since the equilibration of the polarization has not yet been implemented, it can only set *polar\_off* at present.

---

**Note:** You can not use potential file *ffield.comb* with style *comb3*, nor file *ffield.comb3* with style *comb*.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.41.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , where types I and J correspond to two different element types, mixing is performed by LAMMPS as described above from values in the potential file.

These pair styles does not support the *pair\_modify* shift, table, and tail options.

These pair styles do not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style*, *pair\_coeff*, and *fix qeq/comb* commands in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.41.5 Restrictions

These pair styles are part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These pair styles requires the *newton* setting to be “on” for pair interactions.

The COMB potentials in the *ffield.comb* and *ffield.comb3* files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use the COMB potential with any LAMMPS units, but you would need to create your own COMB potential file with coefficients listed in the appropriate units if your simulation does not use “metal” units.

### 19.41.6 Related commands

*pair\_style*, *pair\_coeff*, *fix qeq/comb*

### 19.41.7 Default

none

---

**(COMB)** T.-R. Shan, B. D. Devine, T. W. Kemper, S. B. Sinnott, and S. R. Phillpot, Phys. Rev. B 81, 125328 (2010)

**(COMB3)** T. Liang, T.-R. Shan, Y.-T. Cheng, B. D. Devine, M. Noordhoek, Y. Li, Z. Lu, S. R. Phillpot, and S. B. Sinnott, Mat. Sci. & Eng: R 74, 255-279 (2013).

**(Rick)** S. W. Rick, S. J. Stuart, B. J. Berne, J Chem Phys 101, 6141 (1994).

## 19.42 pair\_style cosine/squared command

### 19.42.1 Syntax

```
pair_style cosine/squared cutoff
```

- cutoff = global cutoff for cosine-squared interactions (distance units)

```
pair_coeff i j eps sigma
pair_coeff i j eps sigma cutoff
pair_coeff i j eps sigma wca
pair_coeff i j eps sigma cutoff wca
```

- $i, j$  = a particle type
- $\text{eps}$  = interaction strength, i.e. the depth of the potential minimum (energy units)
- $\text{sigma}$  = distance of the potential minimum from 0
- $\text{cutoff}$  = the cutoff distance for this pair type, if different from global (distance units)
- $\text{wca}$  = if specified a Weeks-Chandler-Andersen potential (with  $\text{eps}$  strength and minimum at  $\text{sigma}$ ) is added, otherwise not

## 19.42.2 Examples

```
pair_style cosine/squared 3.0
pair_coeff * * 1.0 1.3
pair_coeff 1 3 1.0 1.3 2.0
pair_coeff 1 3 1.0 1.3 wca
pair_coeff 1 3 1.0 1.3 2.0 wca
```

## 19.42.3 Description

Style *cosine/squared* computes a potential of the form

$$E = \begin{cases} -\epsilon & r < \sigma \\ -\epsilon \cos\left(\frac{\pi(r-\sigma)}{2(r_c-\sigma)}\right)^2 & \sigma \leq r < r_c \\ 0 & r \geq r_c \end{cases}$$

between two point particles, where  $(\sigma, -\epsilon)$  is the location of the (rightmost) minimum of the potential, as explained in the syntax section above.

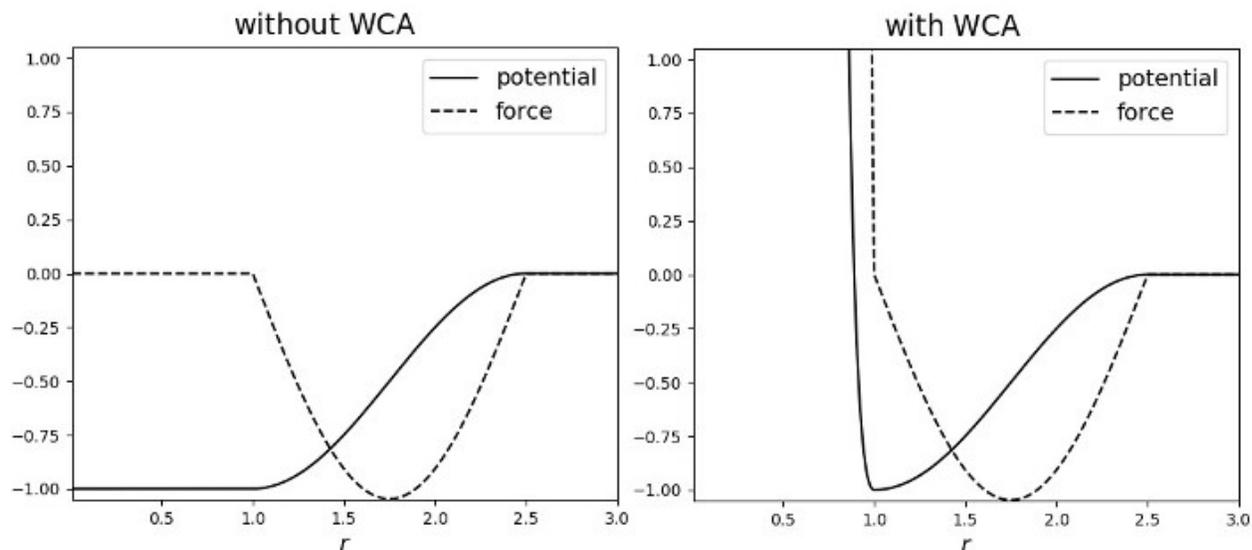
This potential was first used in (*Cooke*) for a coarse-grained lipid membrane model. It is generally very useful as a non-specific interaction potential because it is fully adjustable in depth and width while joining the minimum at  $(\text{sigma}, -\text{epsilon})$  and zero at  $(\text{cutoff}, 0)$  smoothly, requiring no shifting and causing no related artifacts, tail energy calculations etc. This evidently requires *cutoff* to be larger than *sigma*.

If the *wca* option is used then a Weeks-Chandler-Andersen potential (*Weeks*) is added to the above specified cosine-squared potential, specifically the following:

$$E = \epsilon \left[ \left(\frac{\sigma}{r}\right)^{12} - 2 \left(\frac{\sigma}{r}\right)^6 + 1 \right], \quad r < \sigma$$

In this case, and this case only, the  $\sigma$  parameter can be equal to *cutoff* ( $\sigma = \text{cutoff}$ ) which will result in ONLY the WCA potential being used (and print a warning), so the minimum will be attained at  $(\text{sigma}, 0)$ . This is a convenience feature that enables a purely repulsive potential to be used without a need to define an additional pair style and use the hybrid styles.

The energy and force of this pair style for parameters  $\text{epsilon} = 1.0$ ,  $\text{sigma} = 1.0$ ,  $\text{cutoff} = 2.5$ , with and without the WCA potential, are shown in the graphs below:



#### 19.42.4 Mixing, shift, table, tail correction, restart, rRESPA info

Mixing is not supported for this style.

The *shift*, *table* and *tail* options are not relevant for this style.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

#### 19.42.5 Restrictions

The *cosine/squared* style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS is build with that package. See the *Build package* page for more info.

#### 19.42.6 Related commands

*pair\_coeff*, *pair\_style lj/cut*

#### 19.42.7 Default

none

(**Cooke**) “Cooke, Kremer and Deserno, Phys. Rev. E, 72, 011506 (2005)”

(**Weeks**) “Weeks, Chandler and Andersen, J. Chem. Phys., 54, 5237 (1971)”



### 19.43 pair\_style coul/cut command

Accelerator Variants: *coul/cut/gpu*, *coul/cut/kk*, *coul/cut/omp*

### 19.44 pair\_style coul/debye command

Accelerator Variants: *coul/debye/gpu*, *coul/debye/kk*, *coul/debye/omp*

### 19.45 pair\_style coul/dsf command

Accelerator Variants: *coul/dsf/gpu*, *coul/dsf/kk*, *coul/dsf/omp*

### 19.46 pair\_style coul/exclude command

### 19.47 pair\_style coul/cut/global command

Accelerator Variants: *coul/cut/omp*

### 19.48 pair\_style coul/long command

Accelerator Variants: *coul/long/omp*, *coul/long/kk*, *coul/long/gpu*

### 19.49 pair\_style coul/msm command

Accelerator Variants: *coul/msm/omp*

### 19.50 pair\_style coul/streitz command

### 19.51 pair\_style coul/wolf command

Accelerator Variants: *coul/wolf/kk*, *coul/wolf/omp*

## 19.52 pair\_style tip4p/cut command

Accelerator Variants: *tip4p/cut/omp*

## 19.53 pair\_style tip4p/long command

Accelerator Variants: *tip4p/long/omp*

### 19.53.1 Syntax

```
pair_style coul/cut cutoff
pair_style coul/debye kappa cutoff
pair_style coul/dsf alpha cutoff
pair_style coul/exclude cutoff
pair_style coul/cut/global cutoff
pair_style coul/long cutoff
pair_style coul/wolf alpha cutoff
pair_style coul/streitz cutoff keyword alpha
pair_style tip4p/cut otype htype btype atype qdist cutoff
pair_style tip4p/long otype htype btype atype qdist cutoff
```

- cutoff = global cutoff for Coulombic interactions
- kappa = Debye length (inverse distance units)
- alpha = damping parameter (inverse distance units)

### 19.53.2 Examples

```
pair_style coul/cut 2.5
pair_coeff * *
pair_coeff 2 2 3.5

pair_style coul/debye 1.4 3.0
pair_coeff * *
pair_coeff 2 2 3.5

pair_style coul/dsf 0.05 10.0
pair_coeff * *

pair_style hybrid/overlay coul/exclude 10.0 ...
pair_coeff * * coul/exclude

pair_style coul/long 10.0
pair_coeff * *

pair_style coul/msm 10.0
pair_coeff * *
```

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(continued from previous page)

```

pair_style coul/wolf 0.2 9.0
pair_coeff * *

pair_style coul/streitz 12.0 ewald
pair_style coul/streitz 12.0 wolf 0.30
pair_coeff * * Al0.streitz Al 0

pair_style tip4p/cut 1 2 7 8 0.15 12.0
pair_coeff * *

pair_style tip4p/long 1 2 7 8 0.15 10.0
pair_coeff * *

```

### 19.53.3 Description

The *coul/cut* style computes the standard Coulombic interaction potential given by

$$E = \frac{Cq_iq_j}{\epsilon r} \quad r < r_c$$

where C is an energy-conversion constant, Qi and Qj are the charges on the 2 atoms, and  $\epsilon$  is the dielectric constant which can be set by the *dielectric* command. The cutoff  $r_c$  truncates the interaction distance.

Style *coul/debye* adds an additional exp() damping factor to the Coulombic term, given by

$$E = \frac{Cq_iq_j}{\epsilon r} \exp(-\kappa r) \quad r < r_c$$

where  $\kappa$  is the Debye length. This potential is another way to mimic the screening effect of a polar solvent.

Style *coul/dsf* computes Coulombic interactions via the damped shifted force model described in *Fennell*, given by:

$$E = q_iq_j \left[ \frac{\text{erfc}(\alpha r)}{r} - \frac{\text{erfc}(\alpha r_c)}{r_c} + \left( \frac{\text{erfc}(\alpha r_c)}{r_c^2} + \frac{2\alpha}{\sqrt{\pi}} \frac{\exp(-\alpha^2 r_c^2)}{r_c} \right) (r - r_c) \right] \quad r < r_c$$

where  $\alpha$  is the damping parameter and erfc() is the complementary error-function. The potential corrects issues in the Wolf model (described below) to provide consistent forces and energies (the Wolf potential is not differentiable at the cutoff) and smooth decay to zero.

Style *coul/wolf* computes Coulombic interactions via the Wolf summation method, described in *Wolf*, given by:

$$E_i = \frac{1}{2} \sum_{j \neq i} \frac{q_i q_j \text{erfc}(\alpha r_{ij})}{r_{ij}} + \frac{1}{2} \sum_{j \neq i} \frac{q_i q_j \text{erf}(\alpha r_{ij})}{r_{ij}} \quad r < r_c$$

where  $\alpha$  is the damping parameter, and erc() and erfc() are error-function and complementary error-function terms. This potential is essentially a short-range, spherically-truncated, charge-neutralized, shifted, pairwise  $1/r$  summation. With a manipulation of adding and subtracting a self term (for  $i = j$ ) to the first and second term on the right-hand-side, respectively, and a small enough  $\alpha$  damping parameter, the second term shrinks and the potential becomes a rapidly-converging real-space summation. With a long enough cutoff and small enough  $\alpha$  parameter, the energy and forces

calculated by the Wolf summation method approach those of the Ewald sum. So it is a means of getting effective long-range interactions with a short-range potential.

---

Style *coul/streitz* is the Coulomb pair interaction defined as part of the Streitz-Mintmire potential, as described in [this paper](#), in which charge distribution about an atom is modeled as a Slater 1s orbital. More details can be found in the referenced paper. To fully reproduce the published Streitz-Mintmire potential, which is a variable charge potential, style *coul/streitz* must be used with *pair\_style eam/alloy* (or some other short-range potential that has been parameterized appropriately) via the *pair\_style hybrid/overlay* command. Likewise, charge equilibration must be performed via the *fix qeq/slater* command. For example:

```
pair_style hybrid/overlay coul/streitz 12.0 wolf 0.31 eam/alloy
pair_coeff * * coul/streitz AIO.streitz Al O
pair_coeff * * eam/alloy AIO.eam.alloy Al O
fix 1 all qeq/slater 1 12.0 1.0e-6 100 coul/streitz
```

The keyword *wolf* in the *coul/streitz* command denotes computing Coulombic interactions via Wolf summation. An additional damping parameter is required for the Wolf summation, as described for the *coul/wolf* potential above. Alternatively, Coulombic interactions can be computed via an Ewald summation. For example:

```
pair_style hybrid/overlay coul/streitz 12.0 ewald eam/alloy
kspace_style ewald 1e-6
```

Keyword *ewald* does not need a damping parameter, but a *kspace\_style* must be defined, which can be style *ewald* or *pppm*. The Ewald method was used in Streitz and Mintmire's original paper, but a Wolf summation offers a speed-up in some cases.

For the *fix qeq/slater* command, the *qfile* can be a filename that contains QEq parameters as discussed on the [fix qeq](#) command doc page. Alternatively *qfile* can be replaced by "coul/streitz", in which case the *fix* will extract QEq parameters from the *coul/streitz* pair style itself.

See the `examples/streitz` directory for an example input script that uses the Streitz-Mintmire potential. The potentials directory has the `AIO.eam.alloy` and `AIO.streitz` potential files used by the example.

Note that the Streitz-Mintmire potential is generally used for oxides, but there is no conceptual problem with extending it to nitrides and carbides (such as SiC, TiN). Pair *coul/streitz* used by itself or with any other pair style such as EAM, MEAM, Tersoff, or LJ in *hybrid/overlay* mode. To do this, you would need to provide a Streitz-Mintmire parameterization for the material being modeled.

---

Pair style *coul/cut/global* computes the same Coulombic interactions as style *coul/cut* except that it allows only a single global cutoff and thus makes it compatible for use in combination with long-range coulomb styles in *hybrid pair styles*.

Pair style *coul/exclude* computes Coulombic interactions like *coul/cut* but **only** applies them to excluded pairs using a scaling factor of  $\gamma - 1.0$  with  $\gamma$  being the factor assigned to that excluded pair via the *special\_bonds coul* setting. With this it is possible to treat Coulomb interactions for molecular systems with *kspace style scafacos*, which always computes the *full* Coulomb interactions without exclusions. Pair style *coul/exclude* will then *subtract* the excluded interactions accordingly. So to achieve the same forces as with `pair_style lj/cut/coul/long 12.0` with `kspace_style ppm 1.0e-6`, one would use `pair_style hybrid/overlay lj/cut 12.0 coul/exclude 12.0` with `kspace_style scafacos p3m 1.0e-6`.

Styles *coul/long* and *coul/msm* compute the same Coulombic interactions as style *coul/cut* except that an additional damping factor is applied so it can be used in conjunction with the *kspace\_style* command and its *ewald* or *pppm* option. The Coulombic cutoff specified for this style means that pairwise interactions within this distance are computed directly; interactions outside that distance are computed in reciprocal space.

Styles *tip4p/cut* and *tip4p/long* implement the Coulomb part of the TIP4P water model of ([Jorgensen](#)), which introduces a massless site located a short distance away from the oxygen atom along the bisector of the HOH angle. The atomic

types of the oxygen and hydrogen atoms, the bond and angle types for OH and HOH interactions, and the distance to the massless charge site are specified as `pair_style` arguments. Style `tip4p/cut` uses a global cutoff for Coulomb interactions; style `tip4p/long` is for use with a long-range Coulombic solver (Ewald or PPPM).

---

**Note:** For each TIP4P water molecule in your system, the atom IDs for the O and 2 H atoms must be consecutive, with the O atom first. This is to enable LAMMPS to “find” the 2 H atoms associated with each O atom. For example, if the atom ID of an O atom in a TIP4P water molecule is 500, then its 2 H atoms must have IDs 501 and 502.

---

See the [Howto tip4p](#) page for more information on how to use the TIP4P pair styles and lists of parameters to set. Note that the neighbor list cutoff for Coulomb interactions is effectively extended by a distance  $2*qd_{\text{dist}}$  when using the TIP4P pair style, to account for the offset distance of the fictitious charges on O atoms in water molecules. Thus it is typically best in an efficiency sense to use a LJ cutoff  $\geq$  Coulombic cutoff +  $2*qd_{\text{dist}}$ , to shrink the size of the neighbor list. This leads to slightly larger cost for the long-range calculation, so you can test the trade-off for your model.

---

Note that these potentials are designed to be combined with other pair potentials via the `pair_style hybrid/overlay` command. This is because they have no repulsive core. Hence if they are used by themselves, there will be no repulsion to keep two oppositely charged particles from moving arbitrarily close to each other.

The following coefficients must be defined for each pair of atoms types via the `pair_coeff` command as in the examples above, or in the data or restart files read by the `read_data` or `read_restart` commands, or by mixing as described below:

- cutoff (distance units)

For `coul/cut` and `coul/debye` the cutoff coefficient is optional. If it is not used (as in some of the examples above), the default global value specified in the `pair_style` command is used.

For `coul/cut/global`, `coul/long` and `coul/msm` no cutoff can be specified for an individual I,J type pair via the `pair_coeff` command. All type pairs use the same global Coulomb cutoff specified in the `pair_style` command.

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix` *command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 19.53.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , the cutoff distance for the `coul/cut` style can be mixed. The default mix value is *geometric*. See the “`pair_modify`” command for details.

The `pair_modify` shift option is not relevant for these pair styles.

The `coul/long` style supports the `pair_modify` table option for tabulation of the short-range portion of the long-range Coulombic interaction.

These pair styles do not support the `pair_modify` tail option for adding long-range tail corrections to energy and pressure.

---

These pair styles write their information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the `pair` keyword of the `run_style respa` command. They do not support the *inner*, *middle*, *outer* keywords.

---

### 19.53.5 Restrictions

The *coul/cut/global*, *coul/long*, *coul/msm*, *coul/streitz*, and *tip4p/long* styles are part of the KSPACE package. They are only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

### 19.53.6 Related commands

*pair\_coeff*, *pair\_style*, *hybrid/overlay*, *kpace\_style*

### 19.53.7 Default

none

---

**(Wolf)** D. Wolf, P. Keflinski, S. R. Phillpot, J. Eggebrecht, J Chem Phys, 110, 8254 (1999).

**(Fennell)** C. J. Fennell, J. D. Gezelter, J Chem Phys, 124, 234104 (2006).

**(Streitz)** F. H. Streitz, J. W. Mintmire, Phys Rev B, 50, 11996-12003 (1994).

**(Jorgensen)** Jorgensen, Chandrasekhar, Madura, Impey, Klein, J Chem Phys, 79, 926 (1983).

## 19.54 pair\_style coul/diel command

Accelerator Variants: *coul/diel/omp*

### 19.54.1 Syntax

```
pair_style coul/diel cutoff
```

cutoff = global cutoff (distance units)

### 19.54.2 Examples

```
pair_style coul/diel 3.5
pair_coeff 1 4 78. 1.375 0.112
```

### 19.54.3 Description

Style *coul/diel* computes a Coulomb correction for implicit solvent ion interactions in which the dielectric permittivity is distance dependent. The dielectric permittivity  $\epsilon_D(r)$  connects to limiting regimes: One limit is defined by a small dielectric permittivity (close to vacuum) at or close to contact separation between the ions. At larger separations the dielectric permittivity reaches a bulk value used in the regular Coulomb interaction *coul/long* or *coul/cut*. The transition is modeled by a hyperbolic function which is incorporated in the Coulomb correction term for small ion separations as follows

$$E = \frac{Cq_iq_j}{\epsilon r} \left( \frac{\epsilon}{\epsilon_D(r)} - 1 \right) \quad r < r_c$$

$$\epsilon_D(r) = \frac{5.2 + \epsilon}{2} + \frac{\epsilon - 5.2}{2} \tanh \left( \frac{r - r_{me}}{\sigma_e} \right)$$

where  $r_{me}$  is the inflection point of  $\epsilon_D(r)$  and  $\sigma_e$  is a slope defining length scale. C is the same Coulomb conversion factor as in the pair\_styles *coul/cut*, *coul/long*, and *coul/debye*. In this way the Coulomb interaction between ions is corrected at small distances r. The lower limit of  $\epsilon_D(r \rightarrow 0) = 5.2$  due to dielectric saturation (*Stiles*) while the Coulomb interaction reaches its bulk limit by setting  $\epsilon_D(r \rightarrow \infty) = \epsilon$ , the bulk value of the solvent which is 78 for water at 298K.

Examples of the use of this type of Coulomb interaction include implicit solvent simulations of salt ions (*Lenart*) and of ionic surfactants (*Jusuifi*). Note that this potential is only reasonable for implicit solvent simulations and in combination with *coul/cut* or *coul/long*. It is also usually combined with *gauss/cut*, see (*Lenart*) or (*Jusuifi*).

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $\epsilon$  (no units)
- $r_{me}$  (distance units)
- $\sigma_e$  (distance units)

The global cutoff ( $r_c$ ) specified in the *pair\_style* command is used.

---

### 19.54.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support parameter mixing. Coefficients must be given explicitly for each type of particle pairs.

This pair style supports the *pair\_modify* shift option for the energy of the Gauss-potential portion of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.54.5 Restrictions

This style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.54.6 Related commands

*pair\_coeff pair\_style gauss/cut*

### 19.54.7 Default

none

---

**(Stiles)** Stiles , Hubbard, and Kayser, J Chem Phys, 77, 6189 (1982).

**(Lenart)** Lenart , Jusufi, and Panagiotopoulos, J Chem Phys, 126, 044509 (2007).

**(Jusufi)** Jusufi, Hynninen, and Panagiotopoulos, J Phys Chem B, 112, 13783 (2008).

## 19.55 pair\_style coul/shield command

### 19.55.1 Syntax

```
pair_style coul/shield cutoff tap_flag
```

- cutoff = global cutoff (distance units)
- tap\_flag = 0/1 to turn off/on the taper function

### 19.55.2 Examples

```
pair_style coul/shield 16.0 1
pair_coeff 1 2 0.70
```

### 19.55.3 Description

Style *coul/shield* computes a Coulomb interaction for boron and nitrogen atoms located in different layers of hexagonal boron nitride. This potential is designed be used in combination with the pair style *ilp/graphene/hbn*

---

**Note:** This potential is intended for electrostatic interactions between two different layers of hexagonal boron nitride. Therefore, to avoid interaction within the same layers, each layer should have a separate molecule id and is recommended to use the “full” atom style, so that charge and molecule ID information is included.

---



$$E = \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij}$$

$$V_{ij} = \text{Tap}(r_{ij}) \frac{\kappa q_i q_j}{\sqrt[3]{r_{ij}^3 + (1/\lambda_{ij})^3}}$$

$$\text{Tap}(r_{ij}) = 20 \left( \frac{r_{ij}}{R_{cut}} \right)^7 - 70 \left( \frac{r_{ij}}{R_{cut}} \right)^6 + 84 \left( \frac{r_{ij}}{R_{cut}} \right)^5 - 35 \left( \frac{r_{ij}}{R_{cut}} \right)^4 + 1$$

Where  $\text{Tap}(r_{ij})$  is the taper function which provides a continuous cutoff (up to third derivative) for inter-atomic separations larger than  $r_c$  (*Leven1*), (*Leven2*) and (*Maaravi*). Here  $\lambda$  is the shielding parameter that eliminates the short-range singularity of the classical mono-polar electrostatic interaction expression (*Maaravi*).

The shielding parameter  $\lambda$  (1/distance units) must be defined for each pair of atom types via the *pair\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

The global cutoff ( $r_c$ ) specified in the *pair\_style* command is used.

#### 19.55.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support parameter mixing. Coefficients must be given explicitly for each type of particle pairs.

The *pair\_modify table* option is not relevant for this pair style.

This pair style does not support the *pair\_modify tail* option for adding long-range tail corrections to energy and pressure.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

#### 19.55.5 Restrictions

This pair style is part of the INTERLAYER package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 19.55.6 Related commands

*pair\_coeff pair\_style ilp/graphene/hbn*

#### 19.55.7 Default

*tap\_flag* = 1

**(Leven1)** I. Leven, I. Azuri, L. Kronik and O. Hod, J. Chem. Phys. 140, 104106 (2014).

**(Leven2)** I. Leven et al, J. Chem.Theory Comput. 12, 2896-905 (2016).

**(Maaravi)** T. Maaravi et al, J. Phys. Chem. C 121, 22826-22835 (2017).

## 19.56 pair\_style coul/slater command

## 19.57 pair\_style coul/slater/cut command

## 19.58 pair\_style coul/slater/long command

### 19.58.1 Syntax

```
pair_style coul/slater/cut lambda cutoff
pair_style coul/slater/long lambda cutoff
```

lambda = decay length of the charge (distance units) cutoff = cutoff (distance units)

### 19.58.2 Examples

```
pair_style coul/slater/cut 1.0 3.5
pair_coeff * *
pair_coeff 2 2 2.5

pair_style coul/slater/long 1.0 12.0
pair_coeff * *
pair_coeff 1 1 5.0
```

### 19.58.3 Description

Styles *coul/slater* compute electrostatic interactions in mesoscopic models which employ potentials without explicit excluded-volume interactions. The goal is to prevent artificial ionic pair formation by including a charge distribution in the Coulomb potential, following the formulation of (*Melchor*):

$$E = \frac{Cq_iq_j}{\epsilon r} \left( 1 - \left( 1 + \frac{r_{ij}}{\lambda} \exp(-2r_{ij}/\lambda) \right) \right) \quad r < r_c$$

where  $r_c$  is the cutoff distance and  $\lambda$  is the decay length of the charge.  $C$  is the same Coulomb conversion factor as in the pair\_styles *coul/cut* and *coul/long*. In this way the Coulomb interaction between ions is corrected at small distances  $r$ . For the *coul/slater/cut* style, the potential energy for distances larger than the cutoff is zero, while for the *coul/slater/long*, the long-range interactions are computed either by the Ewald or the PPPM technique.

Phenomena that can be captured at a mesoscopic level using this type of electrostatic interactions include the formation of polyelectrolyte-surfactant aggregates, charge stabilization of colloidal suspensions, and the formation of complexes driven by charged species in biological systems. (*Vaiwala*).

The cutoff distance is optional. If it is not used, the default global value specified in the *pair\_style* command is used. For each pair of atom types, a specific cutoff distance can be defined via the *pair\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $r_c$  (distance units)

The global decay length of the charge ( $\lambda$ ) specified in the *pair\_style* command is used for all pairs.

## 19.58.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , the cutoff distance for the *coul/slater* styles can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

The *pair\_modify* shift and table options are not relevant for these pair styles.

These pair styles do not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

These pair styles write their information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

## 19.58.5 Restrictions

The *coul/slater/long* style requires the long-range solvers included in the KSPACE package.

These styles are part of the EXTRA-PAIR package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.58.6 Related commands

*pair\_coeff*, *pair\_style*, *hybrid/overlay*, *kpace\_style*

## 19.58.7 Default

none

---

(Melchor) Gonzalez-Melchor, Mayoral, Velazquez, and Alejandre, J Chem Phys, 125, 224107 (2006).

(Vaiwala) Vaiwala, Jadhav, and Thakkar, J Chem Phys, 146, 124904 (2017).

## 19.59 pair\_style coul/tt command

### 19.59.1 Syntax

```
pair_style style args
```

- style = *coul/tt*
- args = list of arguments for a particular style

*coul/tt* args = n cutoff

n = degree of polynomial

cutoff = global cutoff (distance units)

## 19.59.2 Examples

```
pair_style hybrid/overlay ... coul/tt 4 12.0
pair_coeff 1 2 coul/tt 4.5 1.0
pair_coeff 1 2 coul/tt 4.0 1.0 4 12.0
pair_coeff 1 3* coul/tt 4.5 1.0 4
```

Example input scripts available: examples/PACKAGES/drude

## 19.59.3 Description

The *coul/tt* pair style is meant to be used with force fields that include explicit polarization through Drude dipoles.

The *coul/tt* pair style should be used as a sub-style within in the *pair\_style hybrid/overlay* command, in conjunction with a main pair style including Coulomb interactions and *thole* pair style, or with *lj/cut/thole/long* pair style that is equivalent to the combination of preceding two.

The *coul/tt* pair styles compute the charge-dipole Coulomb interaction damped at short distances by a function

$$f_{n,ij}(r) = 1 - c_{ij} \cdot e^{-b_{ij}r} \sum_{k=0}^n \frac{(b_{ij}r)^k}{k!}$$

This function results from an adaptation to the Coulomb interaction (*Salanne*) of the damping function originally proposed by *Tang Toennies* for van der Waals interactions.

The polynomial takes the degree 4 for damping the Coulomb interaction. The parameters  $b_{ij}$  and  $c_{ij}$  could be determined from first-principle calculations for small, mainly mono-atomic, ions (*Salanne*), or else treated as empirical for large molecules.

In pair styles with Drude induced dipoles, this damping function is typically applied to the interactions between a Drude charge (either  $q_{D,i}$  on a Drude particle or  $-q_{D,i}$  on the respective Drude core) and a charge on a non-polarizable atom,  $q_j$ .

The Tang-Toennies function could also be used to damp electrostatic interactions between the (non-polarizable part of the) charge of a core,  $q_i - q_{D,i}$ , and the Drude charge of another,  $-q_{D,j}$ . The  $b_{ij}$  and  $c_{ij}$  are equal to  $b_{ji}$  and  $c_{ji}$  in the case of core-core interactions.

For pair\_style *coul/tt*, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the example above.

- $b_{ij}$
- $c_{ij}$
- degree of polynomial (positive integer)
- cutoff (distance units)

The last two coefficients are optional. If not specified the global degree of the polynomial or the global cutoff specified in the pair\_style command are used. In order to specify a cutoff (forth argument), the degree of the polynomial (third argument) must also be specified.

### 19.59.4 Mixing, shift, table, tail correction, restart, rRESPA info

The *coul/tt* pair style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

### 19.59.5 Restrictions

These pair styles are part of the DRUDE package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair\_style should currently not be used with the *charmm dihedral style* if the latter has non-zero 1-4 weighting factors. This is because the *coul/tt* pair style does not know which pairs are 1-4 partners of which dihedrals.

### 19.59.6 Related commands

*fix drude*, *fix langevin/drude*, *fix drude/transform*, *compute temp/drude*, *pair\_style thole*

### 19.59.7 Default

none

---

(**Thole**) Chem Phys, 59, 341 (1981).

(**Salanne**) Salanne, Rotenberg, Jahn, Vuilleumier, Simon, Christian and Madden, Theor Chem Acc, 131, 1143 (2012).

(**Tang and Toennies**) J Chem Phys, 80, 3726 (1984).

## 19.60 pair\_style born/coul/dsf/cs command

## 19.61 pair\_style born/coul/long/cs command

Accelerator Variants: *born/coul/long/cs/gpu*

## 19.62 pair\_style born/coul/wolf/cs command

Accelerator Variants: *born/coul/wolf/cs/gpu*

## 19.63 pair\_style buck/coul/long/cs command

## 19.64 pair\_style coul/long/cs command

Accelerator Variants: *coul/long/cs/gpu*

## 19.65 pair\_style coul/wolf/cs command

## 19.66 pair\_style lj/cut/coul/long/cs command

## 19.67 pair\_style lj/class2/coul/long/cs command

### 19.67.1 Syntax

```
pair_style style args
```

- style = *born/coul/dsf/cs* or *born/coul/long/cs* or *born/coul/wolf/cs* or *buck/coul/long/cs* or *coul/long/cs* or *coul/wolf/cs* or *lj/cut/coul/long/cs* or *lj/class2/coul/long/cs*
- args = list of arguments for a particular style

*born/coul/dsf/cs* args = alpha cutoff (cutoff2)

alpha = damping parameter (inverse distance units)

cutoff = global cutoff for non-Coulombic (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (distance units)

*born/coul/long/cs* args = cutoff (cutoff2)

cutoff = global cutoff for non-Coulombic (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

*born/coul/wolf/cs* args = alpha cutoff (cutoff2)

alpha = damping parameter (inverse distance units)

cutoff = global cutoff for Buckingham (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

*buck/coul/long/cs* args = cutoff (cutoff2)

cutoff = global cutoff for Buckingham (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

*coul/long* args = cutoff

cutoff = global cutoff for Coulombic (distance units)

*coul/wolf* args = alpha cutoff

alpha = damping parameter (inverse distance units)

cutoff = global cutoff for Coulombic (distance units)

*lj/cut/coul/long/cs* args = cutoff (cutoff2)

cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

*lj/class2/coul/long/cs* args = cutoff (cutoff2)

cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

### 19.67.2 Examples

```
pair_style born/coul/dsf/cs 0.1 10.0 12.0
pair_coeff * * 0.0 1.00 0.00 0.00 0.00
pair_coeff 1 1 480.0 0.25 0.00 1.05 0.50

pair_style born/coul/long/cs 10.0 8.0
pair_coeff 1 1 6.08 0.317 2.340 24.18 11.51
```

(continues on next page)

(continued from previous page)

```

pair_style born/coul/wolf/cs 0.25 10.0 12.0
pair_coeff * * 0.0 1.00 0.00 0.00 0.00
pair_coeff 1 1 480.0 0.25 0.00 1.05 0.50

pair_style buck/coul/long/cs 10.0
pair_style buck/coul/long/cs 10.0 8.0
pair_coeff * * 100.0 1.5 200.0
pair_coeff 1 1 100.0 1.5 200.0 9.0

pair_style coul/long/cs 10.0
pair_coeff * *

pair_style coul/wolf/cs 0.2 9.0
pair_coeff * *

pair_style lj/cut/coul/long/cs 10.0
pair_style lj/cut/coul/long/cs 10.0 8.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0

```

### 19.67.3 Description

These pair styles are designed to be used with the adiabatic core/shell model of (*Mitchell and Fincham*). See the *Howto coreshell* page for an overview of the model as implemented in LAMMPS.

All the styles are identical to the corresponding pair style without the “/cs” in the name:

- *pair\_style born/coul/dsf*
- *pair\_style born/coul/long*
- *pair\_style born/coul/wolf*
- *pair\_style buck/coul/long*
- *pair\_style coul/long*
- *pair\_style coul/wolf*
- *pair\_style lj/cut/coul/long*
- *pair\_style lj/class2/coul/long*

except that they correctly treat the special case where the distance between two charged core and shell atoms in the same core/shell pair approach  $r = 0.0$ .

Styles with a “/long” in the name are used with a long-range solver for Coulombic interactions via the *kpace\_style* command. They require special treatment of the short-range Coulombic interactions within the cor/shell model.

Specifically, the short-range Coulomb interaction between a core and its shell should be turned off using the *special\_bonds* command by setting the 1-2 weight to 0.0, which works because the core and shell atoms are bonded to each other. This induces a long-range correction approximation which fails at small distances ( $\sim < 10e-8$ ). Therefore, the Coulomb term which is used to calculate the correction factor is extended by a minimal distance ( $r_{min} = 1.0-6$ ) when the interaction between a core/shell pair is treated, as follows

$$E = \frac{Cq_iq_j}{\epsilon(r + r_{min})} \quad r \rightarrow 0$$

where  $C$  is an energy-conversion constant,  $q_i$  and  $q_j$  are the charges on the core and shell,  $\epsilon$  is the dielectric constant and  $r_{min}$  is the minimal distance.

For styles that are not used with a long-range solver, i.e. those with “/dsf” or “/wolf” in the name, the only correction is the addition of a minimal distance to avoid the possible  $r = 0.0$  case for a core/shell pair.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.67.4 Mixing, shift, table, tail correction, restart, rRESPA info

See the corresponding doc pages for pair styles without the “cs” suffix to see how mixing, shifting, tabulation, tail correction, restarting, and rRESPA are handled by these pair styles.

---

#### 19.67.5 Restrictions

These pair styles are part of the CORESHELL package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

#### 19.67.6 Related commands

*pair\_coeff*, *pair\_style born*, *pair\_style buck*

#### 19.67.7 Default

none

---

(**Mitchell and Fincham**) Mitchell, Fincham, J Phys Condensed Matter, 5, 1031-1038 (1993).



## 19.68 `pair_style coul/cut/dielectric` command

## 19.69 `pair_style coul/long/dielectric` command

## 19.70 `pair_style lj/cut/coul/cut/dielectric` command

Accelerator Variants: *lj/cut/coul/cut/dielectric/omp*

## 19.71 `pair_style lj/cut/coul/debye/dielectric` command

Accelerator Variants: *lj/cut/coul/debye/dielectric/omp*

## 19.72 `pair_style lj/cut/coul/long/dielectric` command

Accelerator Variants: *lj/cut/coul/long/dielectric/omp*

## 19.73 `pair_style lj/cut/coul/msm/dielectric` command

## 19.74 `pair_style lj/long/coul/long/dielectric` command

### 19.74.1 Syntax

```
pair_style style args
```

- `style` = *lj/cut/coul/cut/dielectric* or *lj/cut/coul/long/dielectric* or *lj/cut/coul/msm/dielectric* or *lj/long/coul/msm/dielectric*
- `args` = list of arguments for a particular style

### 19.74.2 Examples

```

pair_style coul/cut/dielectric 10.0
pair_coeff * *
pair_coeff 1 1 9.0

pair_style lj/cut/coul/cut/dielectric 10.0
pair_style lj/cut/coul/cut/dielectric 10.0 8.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0

pair_style lj/cut/coul/long/dielectric 10.0
pair_style lj/cut/coul/long/dielectric 10.0 8.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0

```

Used in input scripts:

```
examples/PACKAGES/dielectric/in.confined
examples/PACKAGES/dielectric/in.nopbc
```

### 19.74.3 Description

All these pair styles are derived from the corresponding pair styles without the *dielectric* suffix. In addition to computing atom forces and energies, these pair styles compute the electrical field vector at each atom, which are to be used in the *fix polarize* commands.

These pair styles should be used with *atom\_style dielectric*, which uses atom charges rescaled by their local dielectric constant.

The styles *lj/cut/coul/long/dielectric*, *lj/cut/coul/msm/dielectric*, and *lj/long/coul/long/dielectric* should be used with their *k*space style counterparts, namely, *pppm/dielectric*, *pppm/disp/dielectric*, and *msm/dielectric*, respectively.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.74.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs *I,J* and *I != J*, the epsilon and sigma coefficients and cutoff distances for this pair style can be mixed. The default mix value is *geometric*. See the “*pair\_modify*” command for details.

The *pair\_modify* table option is not relevant for this pair style.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.74.5 Restrictions

These styles are part of the DIELECTRIC package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.74.6 Related commands

*pair\_coeff*, *fix polarize*, *read\_data*

## 19.74.7 Default

none

## 19.75 pair\_style lj/cut/dipole/cut command

Accelerator Variants: *lj/cut/dipole/cut/gpu*, *lj/cut/dipole/cut/omp*

## 19.76 pair\_style lj/sf/dipole/sf command

Accelerator Variants: *lj/sf/dipole/sf/gpu*, *lj/sf/dipole/sf/omp*

## 19.77 pair\_style lj/cut/dipole/long command

Accelerator Variants: *lj/cut/dipole/long/gpu*

## 19.78 pair\_style lj/long/dipole/long command

### 19.78.1 Syntax

```
pair_style lj/cut/dipole/cut cutoff (cutoff2)
pair_style lj/sf/dipole/sf cutoff (cutoff2)
pair_style lj/cut/dipole/long cutoff (cutoff2)
pair_style lj/long/dipole/long flag_lj flag_coul cutoff (cutoff2)
```

- cutoff = global cutoff LJ (and Coulombic if only 1 arg) (distance units)
- cutoff2 = global cutoff for Coulombic and dipole (optional) (distance units)
- flag\_lj = *long* or *cut* or *off*
  - long* = use long-range damping on dispersion  $1/r^6$  term
  - cut* = use a cutoff on dispersion  $1/r^6$  term
  - off* = omit dispersion  $1/r^6$  term entirely
- flag\_coul = *long* or *off*
  - long* = use long-range damping on Coulombic  $1/r$  and point-dipole terms
  - off* = omit Coulombic and point-dipole terms entirely

## 19.78.2 Examples

```

pair_style lj/cut/dipole/cut 10.0
pair_coeff * * 1.0 1.0
pair_coeff 2 3 1.0 1.0 2.5 4.0

pair_style lj/sf/dipole/sf 9.0
pair_coeff * * 1.0 1.0
pair_coeff 2 3 1.0 1.0 2.5 4.0 scale 0.5
pair_coeff 2 3 1.0 1.0 2.5 4.0

pair_style lj/cut/dipole/long 10.0
pair_coeff * * 1.0 1.0
pair_coeff 2 3 1.0 1.0 2.5 4.0

pair_style lj/long/dipole/long long long 3.5 10.0
pair_coeff * * 1.0 1.0
pair_coeff 2 3 1.0 1.0 2.5 4.0

```

## 19.78.3 Description

Style *lj/cut/dipole/cut* computes interactions between pairs of particles that each have a charge and/or a point dipole moment. In addition to the usual Lennard-Jones interaction between the particles (Elj) the charge-charge (Eqq), charge-dipole (Eqp), and dipole-dipole (Epp) interactions are computed by these formulas for the energy (E), force (F), and torque (T) between particles I and J.

$$\begin{aligned}
 E_{LJ} &= 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \\
 E_{qq} &= \frac{q_i q_j}{r} \\
 E_{qp} &= \frac{q}{r^3} (\vec{p} \cdot \vec{r}) \\
 E_{pp} &= \frac{1}{r^3} (\vec{p}_i \cdot \vec{p}_j) - \frac{3}{r^5} (\vec{p}_i \cdot \vec{r})(\vec{p}_j \cdot \vec{r}) \\
 \\ 
 F_{qq} &= \frac{q_i q_j}{r^3} \vec{r} \\
 F_{qp} &= -\frac{q}{r^3} \vec{p} + \frac{3q}{r^5} (\vec{p} \cdot \vec{r}) \vec{r} \\
 F_{pp} &= \frac{3}{r^5} (\vec{p}_i \cdot \vec{p}_j) \vec{r} - \frac{15}{r^7} (\vec{p}_i \cdot \vec{r})(\vec{p}_j \cdot \vec{r}) \vec{r} + \frac{3}{r^5} [(\vec{p}_j \cdot \vec{r}) \vec{p}_i + (\vec{p}_i \cdot \vec{r}) \vec{p}_j] \\
 \\ 
 T_{pq} &= T_{ij} = \frac{q_j}{r^3} (\vec{p}_i \times \vec{r}) \\
 T_{qp} &= T_{ji} = -\frac{q_i}{r^3} (\vec{p}_j \times \vec{r}) \\
 T_{pp} &= T_{ij} = -\frac{1}{r^3} (\vec{p}_i \times \vec{p}_j) + \frac{3}{r^5} (\vec{p}_j \cdot \vec{r})(\vec{p}_i \times \vec{r}) \\
 T_{pp} &= T_{ji} = -\frac{1}{r^3} (\vec{p}_j \times \vec{p}_i) + \frac{3}{r^5} (\vec{p}_i \cdot \vec{r})(\vec{p}_j \times \vec{r})
 \end{aligned}$$

where  $q_i$  and  $q_j$  are the charges on the two particles,  $\vec{p}_i$  and  $\vec{p}_j$  are the dipole moment vectors of the two particles,  $r$  is their separation distance, and the vector  $\vec{r} = \vec{R}_i - \vec{R}_j$  is the separation vector between the two particles. Note that Eqq

and  $F_{qq}$  are simply Coulombic energy and force,  $F_{ij} = -F_{ji}$  as symmetric forces, and  $T_{ij} != -T_{ji}$  since the torques do not act symmetrically. These formulas are discussed in (Allen) and in (Toukmaji).

Also note, that in the code, all of these terms (except  $E_{lj}$ ) have a  $C/\epsilon$  prefactor, the same as the Coulombic term in the LJ + Coulombic pair styles discussed [here](#).  $C$  is an energy-conversion constant and  $\epsilon$  is the dielectric constant which can be set by the *dielectric* command. The same is true of the equations that follow for other dipole pair styles.

Style *lj/sf/dipole/sf* computes “shifted-force” interactions between pairs of particles that each have a charge and/or a point dipole moment. In general, a shifted-force potential is a (slightly) modified potential containing extra terms that make both the energy and its derivative go to zero at the cutoff distance; this removes (cutoff-related) problems in energy conservation and any numerical instability in the equations of motion (Allen). Shifted-force interactions for the Lennard-Jones ( $E_{LJ}$ ), charge-charge ( $E_{qq}$ ), charge-dipole ( $E_{pq}$ ), dipole-charge ( $E_{qp}$ ) and dipole-dipole ( $E_{pp}$ ) potentials are computed by these formulas for the energy ( $E$ ), force ( $F$ ), and torque ( $T$ ) between particles  $I$  and  $J$ :

$$E_{LJ} = 4\epsilon \left\{ \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] + \left[ 6 \left( \frac{\sigma}{r_c} \right)^{12} - 3 \left( \frac{\sigma}{r_c} \right)^6 \right] \left( \frac{r}{r_c} \right)^2 - 7 \left( \frac{\sigma}{r_c} \right)^{12} + 4 \left( \frac{\sigma}{r_c} \right)^6 \right\}$$

$$E_{qq} = \frac{q_i q_j}{r} \left( 1 - \frac{r}{r_c} \right)^2$$

$$E_{pq} = E_{ji} = -\frac{q}{r^3} \left[ 1 - 3 \left( \frac{r}{r_c} \right)^2 + 2 \left( \frac{r}{r_c} \right)^3 \right] (\vec{p} \bullet \vec{r})$$

$$E_{qp} = E_{ij} = \frac{q}{r^3} \left[ 1 - 3 \left( \frac{r}{r_c} \right)^2 + 2 \left( \frac{r}{r_c} \right)^3 \right] (\vec{p} \bullet \vec{r})$$

$$E_{pp} = \left[ 1 - 4 \left( \frac{r}{r_c} \right)^3 + 3 \left( \frac{r}{r_c} \right)^4 \right] \left[ \frac{1}{r^3} (\vec{p}_i \bullet \vec{p}_j) - \frac{3}{r^5} (\vec{p}_i \bullet \vec{r})(\vec{p}_j \bullet \vec{r}) \right]$$

$$F_{LJ} = \left\{ \left[ 48\epsilon \left( \frac{\sigma}{r} \right)^{12} - 24\epsilon \left( \frac{\sigma}{r} \right)^6 \right] \frac{1}{r^2} - \left[ 48\epsilon \left( \frac{\sigma}{r_c} \right)^{12} - 24\epsilon \left( \frac{\sigma}{r_c} \right)^6 \right] \frac{1}{r_c^2} \right\} \vec{r}$$

$$F_{qq} = \frac{q_i q_j}{r} \left( \frac{1}{r^2} - \frac{1}{r_c^2} \right) \vec{r}$$

$$F_{pq} = F_{ij} = -\frac{3q}{r^5} \left[ 1 - \left( \frac{r}{r_c} \right)^2 \right] (\vec{p} \bullet \vec{r}) \vec{r} + \frac{q}{r^3} \left[ 1 - 3 \left( \frac{r}{r_c} \right)^2 + 2 \left( \frac{r}{r_c} \right)^3 \right] \vec{p}$$

$$F_{qp} = F_{ij} = \frac{3q}{r^5} \left[ 1 - \left( \frac{r}{r_c} \right)^2 \right] (\vec{p} \bullet \vec{r}) \vec{r} - \frac{q}{r^3} \left[ 1 - 3 \left( \frac{r}{r_c} \right)^2 + 2 \left( \frac{r}{r_c} \right)^3 \right] \vec{p}$$

$$F_{pp} = \frac{3}{r^5} \left\{ \left[ 1 - \left( \frac{r}{r_c} \right)^4 \right] \left[ (\vec{p}_i \bullet \vec{p}_j) - \frac{3}{r^2} (\vec{p}_i \bullet \vec{r})(\vec{p}_j \bullet \vec{r}) \right] \vec{r} + \left[ 1 - 4 \left( \frac{r}{r_c} \right)^3 + 3 \left( \frac{r}{r_c} \right)^4 \right] \left[ (\vec{p}_j \bullet \vec{r}) \vec{p}_i + (\vec{p}_i \bullet \vec{r}) \vec{p}_j - \frac{2}{r^2} (\vec{p}_i \bullet \vec{r})(\vec{p}_j \bullet \vec{r}) \vec{r} \right] \right\}$$

$$\begin{aligned}
T_{pq} = T_{ij} &= \frac{q_j}{r^3} \left[ 1 - 3 \left( \frac{r}{r_c} \right)^2 + 2 \left( \frac{r}{r_c} \right)^3 \right] (\vec{p}_i \times \vec{r}) \\
T_{qp} = T_{ji} &= -\frac{q_i}{r^3} \left[ 1 - 3 \left( \frac{r}{r_c} \right)^2 + 2 \left( \frac{r}{r_c} \right)^3 \right] (\vec{p}_j \times \vec{r}) \\
T_{pp} = T_{ij} &= -\frac{1}{r^3} \left[ 1 - 4 \left( \frac{r}{r_c} \right)^3 + e3 \left( \frac{r}{r_c} \right)^4 \right] (\vec{p}_i \times \vec{p}_j) + \\
&\quad \frac{3}{r^5} \left[ 1 - 4 \left( \frac{r}{r_c} \right)^3 + 3 \left( \frac{r}{r_c} \right)^4 \right] (\vec{p}_j \bullet \vec{r})(\vec{p}_i \times \vec{r}) \\
T_{pp} = T_{ji} &= -\frac{1}{r^3} \left[ 1 - 4 \left( \frac{r}{r_c} \right)^3 + 3 \left( \frac{r}{r_c} \right)^4 \right] (\vec{p}_j \times \vec{p}_i) + \\
&\quad \frac{3}{r^5} \left[ 1 - 4 \left( \frac{r}{r_c} \right)^3 + 3 \left( \frac{r}{r_c} \right)^4 \right] (\vec{p}_i \bullet \vec{r})(\vec{p}_j \times \vec{r})
\end{aligned}$$

where  $\epsilon$  and  $\sigma$  are the standard LJ parameters,  $r_c$  is the cutoff,  $q_i$  and  $q_j$  are the charges on the two particles,  $\vec{p}_i$  and  $\vec{p}_j$  are the dipole moment vectors of the two particles,  $r$  is their separation distance, and the vector  $\mathbf{r} = \mathbf{R}_i - \mathbf{R}_j$  is the separation vector between the two particles. Note that  $E_{qq}$  and  $F_{qq}$  are simply Coulombic energy and force,  $F_{ij} = -F_{ji}$  as symmetric forces, and  $T_{ij} \neq -T_{ji}$  since the torques do not act symmetrically. The shifted-force formula for the Lennard-Jones potential is reported in (*Stoddard*). The original (non-shifted) formulas for the electrostatic potentials, forces and torques can be found in (*Price*). The shifted-force electrostatic potentials have been obtained by applying equation 5.13 of (*Allen*). The formulas for the corresponding forces and torques have been obtained by applying the ‘chain rule’ as in appendix C.3 of (*Allen*).

If one cutoff is specified in the `pair_style` command, it is used for both the LJ and Coulombic (q,p) terms. If two cutoffs are specified, they are used as cutoffs for the LJ and Coulombic (q,p) terms respectively. This pair style also supports an optional `scale` keyword as part of a `pair_coeff` statement, where the interactions can be scaled according to this factor. This scale factor is also made available for use with `fix adapt`.

Style `lj/cut/dipole/long` computes the short-range portion of point-dipole interactions as discussed in (*Toukmaji*). Dipole-dipole, dipole-charge, and charge-charge interactions are all supported, along with the standard 12/6 Lennard-Jones interactions, which are computed with a cutoff. A `kspace_style` must be defined to use this pair style. If only dipoles (not point charges) are included in the model, the `kspace` style can be one of these 3 options, all of which compute the long-range portion of dipole-dipole interactions. If the model includes point charges (in addition to dipoles), then only the first of these `kspace` styles can be used:

- `kspace_style ewald/disp`
- `kspace_style ewald/dipole`
- `kspace_style pppm/dipole`

Style `lj/long/dipole/long` has the same functionality as style `lj/cut/dipole/long`, except it also has an option to compute 12/6 Lennard-Jones interactions for use with a long-range dispersion `kspace` style. This is done by setting its `flag_lj` argument to `long`. For long-range LJ interactions, the `kspace_style ewald/disp` command must be used.

The following coefficients must be defined for each pair of atoms types via the `pair_coeff` command as in the examples above, or in the data file or restart files read by the `read_data` or `read_restart` commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- `cutoff1` (distance units)

- `cutoff2` (distance units)

The latter 2 coefficients are optional. If not specified, the global LJ and Coulombic cutoffs specified in the `pair_style` command are used. If only one cutoff is specified, it is used as the cutoff for both LJ and Coulombic interactions for this type pair. If both coefficients are specified, they are used as the LJ and Coulombic cutoffs for this type pair.

---

Note that for systems using these pair styles, typically particles should be able to exert torque on each other via their dipole moments so that the particle and its dipole moment can rotate. This requires they not be point particles, but finite-size spheres. Thus you should use a command like `atom_style hybrid sphere dipole` to use particles with both attributes.

The magnitude and orientation of the dipole moment for each particle can be defined by the `set` command or in the “Atoms” section of the data file read in by the `read_data` command.

Rotating finite-size particles have 6 degrees of freedom (DOFs), translation and rotational. You can use the `compute temp/sphere` command to monitor a temperature which includes all these DOFs.

Finite-size particles with dipole moments should be integrated using one of these options:

- `fix nve/sphere update dipole`
- `fix nve/sphere update dipole` plus `fix langevin omega yes`
- `fix nvt/sphere update dipole`
- `fix npt/sphere update dipole`

In all cases the “update dipole” setting insures the dipole moments are also rotated when the finite-size spheres rotate. The 2nd and 3rd bullets perform thermostating; in the case of a Langevin thermostat the “omega yes” option also thermostats the rotational degrees of freedom (if desired). The 4th bullet performs thermostating and barostatting.

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix` *command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 19.78.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distances for this pair style can be mixed. The default mix value is *geometric*. See the “`pair_modify`” command for details.

For atom type pairs I,J and I != J, the A, sigma, d1, and d2 coefficients and cutoff distance for this pair style can be mixed. A is an energy value mixed like a LJ epsilon. D1 and d2 are distance values and are mixed like sigma. The default mix value is *geometric*. See the “`pair_modify`” command for details.

This pair style does not support the `pair_modify` shift option for the energy of the Lennard-Jones portion of the pair interaction; such energy goes to zero at the cutoff by construction.

The `pair_modify` table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.78.5 Restrictions

The *lj/cut/dipole/cut*, *lj/cut/dipole/long*, *lj/long/dipole/long*, and *lj/sf/dipole/sf\** styles are part of the DIPOLE package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Using dipole pair styles with *electron units* is not currently supported.

### 19.78.6 Related commands

*pair\_coeff*, *set*, *read\_data*, *fix nve/sphere*, *fix nvt/sphere*

### 19.78.7 Default

none

---

(Allen) Allen and Tildesley, Computer Simulation of Liquids, Clarendon Press, Oxford, 1987.

(Toukmaji) Toukmaji, Sagui, Board, and Darden, J Chem Phys, 113, 10913 (2000).

(Stoddard) Stoddard and Ford, Phys Rev A, 8, 1504 (1973).

(Price) Price, Stone and Alderton, Mol Phys, 52, 987 (1984).

## 19.79 pair\_style dpd command

Accelerator Variants: *dpd/gpu*, *dpd/intel*, *dpd/kk*, *dpd/omp*

## 19.80 pair\_style dpd/tstat command

Accelerator Variants: *dpd/tstat/gpu*, *dpd/tstat/kk*, *dpd/tstat/omp*

### 19.80.1 Syntax

```
pair_style dpd T cutoff seed
pair_style dpd/tstat Tstart Tstop cutoff seed
```

- T = temperature (temperature units)
- Tstart,Tstop = desired temperature at start/end of run (temperature units)
- cutoff = global cutoff for DPD interactions (distance units)
- seed = random # seed (positive integer)



## 19.80.2 Examples

```
pair_style dpd 1.0 2.5 34387
pair_coeff * * 3.0 1.0
pair_coeff 1 1 3.0 1.0 1.0

pair_style dpd/tstat 1.0 1.0 2.5 34387
pair_coeff * * 1.0
pair_coeff 1 1 1.0 1.0
```

## 19.80.3 Description

Style *dpd* computes a force field for dissipative particle dynamics (DPD) following the exposition in (*Groot*).

Style *dpd/tstat* invokes a DPD thermostat on pairwise interactions, which is equivalent to the non-conservative portion of the DPD force field. This pairwise thermostat can be used in conjunction with any *pair style*, and in lieu of per-particle thermostats like *fix langevin* or ensemble thermostats like Nose Hoover as implemented by *fix nvt*. To use *dpd/tstat* as a thermostat for another pair style, use the *pair\_style hybrid/overlay* command to compute both the desired pair interaction and the thermostat for each pair of particles.

For style *dpd*, the force on atom I due to atom J is given as a sum of 3 terms

$$\begin{aligned}\vec{f} &= (F^C + F^D + F^R)r_{ij}^{\hat{}} & r < r_c \\ F^C &= Aw(r) \\ F^D &= -\gamma w^2(r)(r_{ij}^{\hat{}} \bullet \vec{v}_{ij}) \\ F^R &= \sigma w(r)\alpha(\Delta t)^{-1/2} \\ w(r) &= 1 - \frac{r}{r_c}\end{aligned}$$

where  $F^C$  is a conservative force,  $F^D$  is a dissipative force, and  $F^R$  is a random force.  $r_{ij}^{\hat{}}$  is a unit vector in the direction  $r_i - r_j$ ,  $\vec{v}_{ij}$  is the vector difference in velocities of the two atoms  $\vec{v}_i - \vec{v}_j$ ,  $\alpha$  is a Gaussian random number with zero mean and unit variance,  $dt$  is the timestep size, and  $w(r)$  is a weighting factor that varies between 0 and 1.  $r_c$  is the pairwise cutoff.  $\sigma$  is set equal to  $\sqrt{2k_B T \gamma}$ , where  $k_B$  is the Boltzmann constant and  $T$  is the temperature parameter in the *pair\_style* command.

For style *dpd/tstat*, the force on atom I due to atom J is the same as the above equation, except that the conservative  $F^C$  term is dropped. Also, during the run,  $T$  is set each timestep to a ramped value from  $Tstart$  to  $Tstop$ .

For style *dpd*, the pairwise energy associated with style *dpd* is only due to the conservative force term  $F^C$ , and is shifted to be zero at the cutoff distance  $r_c$ . The pairwise virial is calculated using all 3 terms. For style *dpd/tstat* there is no pairwise energy, but the last two terms of the formula make a contribution to the virial.

For style *dpd*, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- A (force units)
- $\gamma$  (force/velocity units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global DPD cutoff is used. Note that sigma is set equal to  $\sqrt{2 T \gamma}$ , where T is the temperature set by the *pair\_style* command so it does not need to be specified.

For style *dpd/tstat*, the coefficients defined for each pair of atoms types via the *pair\_coeff* command is the same, except that A is not included.

The GPU-accelerated versions of these styles are implemented based on the work of (*Afshar*) and (*Phillips*).

---

**Note:** If you are modeling DPD polymer chains, you may want to use the *pair\_style srp* command in conjunction with these pair styles. It is a soft segmental repulsive potential (SRP) that can prevent DPD polymer chains from crossing each other.

---

---

**Note:** The virial calculation for pressure when using these pair styles includes all the components of force listed above, including the random force. Since the random force depends on random numbers, everything that changes the order of atoms in the neighbor list (e.g. different number of MPI ranks or a different neighbor list skin distance) will also change the sequence in which the random numbers are applied and thus the individual forces and therefore also the virial/pressure.

---

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 19.80.4 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles do not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

These pair styles do not support the *pair\_modify* shift option for the energy of the pair interaction. Note that as discussed above, the energy due to the conservative  $F^C$  term is already shifted to be 0.0 at the cutoff distance  $r_c$ .

The *pair\_modify* table option is not relevant for these pair styles.

These pair styles do not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

These pair styles write their information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file. Note that the user-specified random number seed is stored in the restart file, so when a simulation is restarted, each processor will re-initialize its random number generator the same way it did initially. This means the random forces will be random, but will not be the same as they would have been if the original simulation had continued past the restart time.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

The *dpd/tstat* style can ramp its target temperature over multiple runs, using the *start* and *stop* keywords of the *run* command. See the *run* command for details of how to do this.

---

## 19.80.5 Restrictions

These styles are part of the DPD-BASIC package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The default frequency for rebuilding neighbor lists is every 10 steps (see the *neigh\_modify* command). This may be too infrequent for style *dpd* simulations since particles move rapidly and can overlap by large amounts. If this setting yields a non-zero number of “dangerous” reneighborings (printed at the end of a simulation), you should experiment with forcing reneighborings more often and see if system energies/trajectories change.

These pair styles requires you to use the *comm\_modify vel yes* command so that velocities are stored by ghost atoms.

These pair styles will not restart exactly when using the *read\_restart* command, though they should provide statistically similar results. This is because the forces they compute depend on atom velocities. See the *read\_restart* command for more details.

## 19.80.6 Related commands

*pair\_coeff*, *fix nvt*, *fix langevin*, *pair\_style srp*

## 19.80.7 Default

none

---

**(Groot)** Groot and Warren, J Chem Phys, 107, 4423-35 (1997).

**(Afshar)** Afshar, F. Schmid, A. Pischevar, S. Worley, Comput Phys Comm, 184, 1119-1128 (2013).

**(Phillips)** C. L. Phillips, J. A. Anderson, S. C. Glotzer, Comput Phys Comm, 230, 7191-7201 (2011).

## 19.81 pair\_style dpd/ext command

Accelerator Variants: dpd/ext/kk

## 19.82 pair\_style dpd/ext/tstat command

Accelerator Variants: dpd/ext/tstat/kk

### 19.82.1 Syntax

```
pair_style dpd/ext T cutoff seed
pair_style dpd/ext/tstat Tstart Tstop cutoff seed
```

- T = temperature (temperature units)
- Tstart, Tstop = desired temperature at start/end of run (temperature units)
- cutoff = global cutoff for DPD interactions (distance units)
- seed = random # seed (positive integer)

## 19.82.2 Examples

```

pair_style dpd/ext 1.0 2.5 34387
pair_coeff 1 1 25.0 4.5 4.5 0.5 0.5 1.2
pair_coeff 1 2 40.0 4.5 4.5 0.5 0.5 1.2

pair_style dpd/ext/tstat 1.0 1.0 2.5 34387
pair_coeff 1 1 4.5 4.5 0.5 0.5 1.2
pair_coeff 1 2 4.5 4.5 0.5 0.5 1.2

```

## 19.82.3 Description

The style *dpd/ext* computes an extended force field for dissipative particle dynamics (DPD) following the exposition in (Groot), (Junghans).

Style *dpd/ext/tstat* invokes an extended DPD thermostat on pairwise interactions, equivalent to the non-conservative portion of the extended DPD force field. To use *dpd/ext/tstat* as a thermostat for another pair style, use the *pair\_style hybrid/overlay* command to compute both the desired pair interaction and the thermostat for each pair of particles.

For the style *dpd/ext*, the force on atom I due to atom J is given as a sum of 3 terms

$$\begin{aligned}
 \mathbf{f} &= f^C + f^D + f^R & r < r_c \\
 f^C &= A_{ij} w(r) \hat{\mathbf{r}}_{ij} \\
 f^D &= -\gamma_{\parallel} w_{\parallel}^2(r) (\hat{\mathbf{r}}_{ij} \cdot \mathbf{v}_{ij}) \hat{\mathbf{r}}_{ij} - \gamma_{\perp} w_{\perp}^2(r) (\mathbf{I} - \hat{\mathbf{r}}_{ij} \hat{\mathbf{r}}_{ij}^T) \mathbf{v}_{ij} \\
 f^R &= \sigma_{\parallel} w_{\parallel}(r) \frac{\alpha}{\sqrt{\Delta t}} \hat{\mathbf{r}}_{ij} + \sigma_{\perp} w_{\perp}(r) (\mathbf{I} - \hat{\mathbf{r}}_{ij} \hat{\mathbf{r}}_{ij}^T) \frac{\xi_{ij}}{\sqrt{\Delta t}} \\
 w(r) &= 1 - r/r_c
 \end{aligned}$$

where  $\mathbf{f}^C$  is a conservative force,  $\mathbf{f}^D$  is a dissipative force, and  $\mathbf{f}^R$  is a random force.  $A_{ij}$  is the maximum repulsion between the two atoms,  $\hat{\mathbf{r}}_{ij}$  is a unit vector in the direction  $\mathbf{r}_i - \mathbf{r}_j$ ,  $\mathbf{v}_{ij} = \mathbf{v}_i - \mathbf{v}_j$  is the vector difference in velocities of the two atoms,  $\alpha$  and  $\xi_{ij}$  are Gaussian random numbers with zero mean and unit variance,  $\Delta t$  is the timestep,  $w(r) = 1 - r/r_c$  is a weight function for the conservative interactions that varies between 0 and 1,  $r_c$  is the corresponding cutoff,  $w_{\alpha}(r) = (1 - r/\bar{r}_c)^{s_{\alpha}}$ ,  $\alpha \equiv (\parallel, \perp)$ , are weight functions with coefficients  $s_{\alpha}$  that vary between 0 and 1,  $\bar{r}_c$  is the corresponding cutoff,  $\mathbf{I}$  is the unit matrix,  $\sigma_{\alpha} = \sqrt{2k_B T \gamma_{\alpha}}$ , where  $k_B$  is the Boltzmann constant and  $T$  is the temperature in the *pair\_style* command.

For the style *dpd/ext/tstat*, the force on atom I due to atom J is the same as the above equation, except that the conservative  $\mathbf{f}^C$  term is dropped. Also, during the run, T is set each timestep to a ramped value from Tstart to Tstop.

For the style *dpd/ext*, the pairwise energy associated with style *dpd/ext* is only due to the conservative force term  $\mathbf{f}^C$ , and is shifted to be zero at the cutoff distance  $r_c$ . The pairwise virial is calculated using all three terms. There is no pairwise energy for style *dpd/ext/tstat*, but the last two terms of the formula contribute the virial.

For the style *dpd/ext/tstat*, the force on atom I due to atom J is the same as the above equation, except that the conservative  $\mathbf{f}^C$  term is dropped. Also, during the run, T is set each timestep to a ramped value from Tstart to Tstop.

For the style *dpd/ext*, the pairwise energy associated with style *dpd/ext* is only due to the conservative force term  $\mathbf{f}^C$ , and is shifted to be zero at the cutoff distance  $r_c$ . The pairwise virial is calculated using all three terms. There is no pairwise energy for style *dpd/ext/tstat*, but the last two terms of the formula contribute the virial.

For the style *dpd/ext*, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above:

- A (force units)
- $\gamma_{\perp}$  (force/velocity units)

- $\gamma_{\parallel}$  (force/velocity units)
- $s_{\perp}$  (unitless)
- $s_{\parallel}$  (unitless)
- $r_c$  (distance units)

The last coefficient is optional. If not specified, the global DPD cutoff is used. Note that  $\sigma$ 's are set equal to  $\sqrt{2k_B T \gamma}$ , where  $T$  is the temperature set by the *pair\_style* command so it does not need to be specified.

For the style *dpd/ext/tstat*, the coefficients defined for each pair of atoms types via the *pair\_coeff* command is the same, except that A is not included.

---

**Note:** If you are modeling DPD polymer chains, you may want to use the *pair\_style srp* command in conjunction with these pair styles. It is a soft segmental repulsive potential (SRP) that can prevent DPD polymer chains from crossing each other.

---

---

**Note:** The virial calculation for pressure when using this pair style includes all the components of force listed above, including the random force.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### Mixing, shift, table, tail correction, restart, rRESPA info:

The style *dpd/ext* does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

The pair styles do not support the *pair\_modify* shift option for the energy of the pair interaction. Note that as discussed above, the energy due to the conservative  $f^C$  term is already shifted to be zero at the cutoff distance  $r_c$ .

The *pair\_modify* table option is not relevant for the style *dpd/ext*.

The style *dpd/ext* does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

The pair styles can only be used via the pair keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, and *outer* keywords.

The style *dpd/ext/tstat* can ramp its target temperature over multiple runs, using the start and stop keywords of the *run* command. See the *run* command for details of how to do this.

---

## 19.82.4 Restrictions

These styles are part of the DPD-BASIC package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The default frequency for rebuilding neighbor lists is every 10 steps (see the *neigh\_modify* command). This may be too infrequent for style *dpd/ext* simulations since particles move rapidly and can overlap by large amounts. If this setting yields a non-zero number of say { dangerous } reneighborings (printed at the end of a simulation), you should experiment with forcing reneighborings more often and see if system energies/trajectories change.

The pair styles require to use the *comm\_modify vel yes* command so that velocities are stored by ghost atoms.

The pair styles will not restart exactly when using the *read\_restart* command, though they should provide statistically similar results. This is because the forces they compute depend on atom velocities. See the *read\_restart* command for more details.

## 19.82.5 Related commands

*pair\_style dpd*, *pair\_coeff*, *fix nvt*, *fix langevin*, *pair\_style srp*

**Default:** none

---

**(Groot)** Groot and Warren, J Chem Phys, 107, 4423-35 (1997).

**(Junghans)** Junghans, Praprotnik and Kremer, Soft Matter 4, 156, 1119-1128 (2008).

## 19.83 pair\_style dpd/fdt command

## 19.84 pair\_style dpd/fdt/energy command

Accelerator Variants: *dpd/fdt/energy/kk*

### 19.84.1 Syntax

```
pair_style style args
```

- style = *dpd/fdt* or *dpd/fdt/energy*
- args = list of arguments for a particular style

*dpd/fdt* args = T cutoff seed

T = temperature (temperature units)

cutoff = global cutoff for DPD interactions (distance units)

seed = random # seed (positive integer)

*dpd/fdt/energy* args = cutoff seed

cutoff = global cutoff for DPD interactions (distance units)

seed = random # seed (positive integer)

## 19.84.2 Examples

```
pair_style dpd/fdt 300.0 2.5 34387
pair_coeff * * 3.0 1.0 2.5

pair_style dpd/fdt/energy 2.5 34387
pair_coeff * * 3.0 1.0 0.1 2.5
```

## 19.84.3 Description

Styles *dpd/fdt* and *dpd/fdt/energy* compute the force for dissipative particle dynamics (DPD) simulations. The *dpd/fdt* style is used to perform DPD simulations under isothermal and isobaric conditions, while the *dpd/fdt/energy* style is used to perform DPD simulations under isoenergetic and isoenthalpic conditions (see *(Lisal)*). For DPD simulations in general, the force on atom I due to atom J is given as a sum of 3 terms

$$\begin{aligned}\vec{f} &= (F^C + F^D + F^R)r_{ij}^{\hat{}} & r < r_c \\ F^C &= Aw(r) \\ F^D &= -\gamma w^2(r)(r_{ij}^{\hat{}} \bullet \vec{v}_{ij}) \\ F^R &= \sigma w(r)\alpha(\Delta t)^{-1/2} \\ w(r) &= 1 - \frac{r}{r_c}\end{aligned}$$

where  $F^C$  is a conservative force,  $F^D$  is a dissipative force, and  $F^R$  is a random force.  $r_{ij}^{\hat{}}$  is a unit vector in the direction  $r_i - r_j$ ,  $\vec{v}_{ij}$  is the vector difference in velocities of the two atoms,  $\vec{v}_i - \vec{v}_j$ ,  $\alpha$  is a Gaussian random number with zero mean and unit variance,  $dt$  is the timestep size, and  $w(r)$  is a weighting factor that varies between 0 and 1,  $r_c$  is the pairwise cutoff. Note that alternative definitions of the weighting function exist, but would have to be implemented as a separate pair style command.

For style *dpd/fdt*, the fluctuation-dissipation theorem defines  $\gamma$  to be set equal to  $\sigma^2/(2T)$ , where  $T$  is the set point temperature specified as a pair style parameter in the above examples. The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- A (force units)
- $\sigma$  (force\*time<sup>1/2</sup>) units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global DPD cutoff is used.

Style *dpd/fdt/energy* is used to perform DPD simulations under isoenergetic and isoenthalpic conditions. The fluctuation-dissipation theorem defines  $\gamma$  to be set equal to  $\sigma^2/(2\theta)$ , where  $\theta$  is the average internal temperature for the pair. The particle internal temperature is related to the particle internal energy through a mesoparticle equation of state (see *fix eos*). The differential internal conductive and mechanical energies are computed within style *dpd/fdt/energy* as:

$$\begin{aligned}du_i^{cond} &= \kappa_{ij} \left( \frac{1}{\theta_i} - \frac{1}{\theta_j} \right) \omega_{ij}^2 + \alpha_{ij} \omega_{ij} \zeta_{ij}^q (\Delta t)^{-1/2} \\ du_i^{mech} &= -\frac{1}{2} \gamma_{ij} \omega_{ij}^2 \left( \frac{\vec{r}_{ij}}{r_{ij}} \bullet \vec{v}_{ij} \right)^2 - \frac{\sigma_{ij}^2}{4} \left( \frac{1}{m_i} + \frac{1}{m_j} \right) \omega_{ij}^2 - \frac{1}{2} \sigma_{ij} \omega_{ij} \left( \frac{\vec{r}_{ij}}{r_{ij}} \bullet \vec{v}_{ij} \right) \zeta_{ij} (\Delta t)^{-1/2}\end{aligned}$$

where

$$\begin{aligned}\alpha_{ij}^2 &= 2k_B\kappa_{ij} \\ \sigma_{ij}^2 &= 2\gamma_{ij}k_B\Theta_{ij} \\ \Theta_{ij}^{-1} &= \frac{1}{2}\left(\frac{1}{\theta_i} + \frac{1}{\theta_j}\right)\end{aligned}$$

$\zeta_{i,j}^q$  is a second Gaussian random number with zero mean and unit variance that is used to compute the internal conductive energy. The fluctuation-dissipation theorem defines  $\alpha^2$  to be set equal to  $2k_B\kappa$ , where  $\kappa$  is the mesoparticle thermal conductivity parameter. The following coefficients must be defined for each pair of atoms types via the `pair_coeff` command as in the examples above, or in the data file or restart files read by the `read_data` or `read_restart` commands:

- A (force units)
- $\sigma$  (force\*time<sup>(1/2)</sup> units)
- $\kappa$  (energy\*temperature/time units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global DPD cutoff is used.

The pairwise energy associated with styles `dpd/fdt` and `dpd/fdt/energy` is only due to the conservative force term  $F^C$ , and is shifted to be zero at the cutoff distance  $r_c$ . The pairwise virial is calculated using only the conservative term.

The forces computed through the `dpd/fdt` and `dpd/fdt/energy` styles can be integrated with the velocity-Verlet integration scheme or the Shardlow splitting integration scheme described by (Lisal). In the cases when these pair styles are combined with the `fix shardlow`, these pair styles differ from the other dpd styles in that the dissipative and random forces are split from the force calculation and are not computed within the pair style. Thus, only the conservative force is computed by the pair style, while the stochastic integration of the dissipative and random forces are handled through the Shardlow splitting algorithm approach. The Shardlow splitting algorithm is advantageous, especially when performing DPD under isoenergetic conditions, as it allows significantly larger timesteps to be taken.

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix` *command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 19.84.4 Restrictions

These commands are part of the DPD-REACT package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

Pair styles `dpd/fdt` and `dpd/fdt/energy` require use of the `comm_modify vel yes` option so that velocities are stored by ghost atoms.

Pair style `dpd/fdt/energy` requires `atom_style dpd` to be used in order to properly account for the particle internal energies and temperatures.



## 19.84.5 Related commands

`pair_coeff`, `fix shardlow`

## 19.84.6 Default

none

(Lisal) M. Lisal, J.K. Brennan, J. Bonet Avalos, “Dissipative particle dynamics at isothermal, isobaric, isoenergetic, and isoenthalpic conditions using Shardlow-like splitting algorithms.”, J. Chem. Phys., 135, 204105 (2011).

## 19.85 pair\_style drip command

### 19.85.1 Syntax

```
pair_style hybrid/overlay drip [styles ...]
```

- styles = other styles to be overlaid with drip (optional)

### 19.85.2 Examples

```
pair_style hybrid/overlay drip
pair_coeff * * none
pair_coeff * * drip C.drip C

pair_style hybrid/overlay drip rebo
pair_coeff * * drip C.drip C
pair_coeff * * rebo CH.airebo C

pair_style hybrid/overlay drip rebo
pair_coeff * * drip C.drip C NULL
pair_coeff * * rebo CH.airebo C H
```

### 19.85.3 Description

Style `drip` computes the interlayer interactions of layered materials using the dihedral-angle-corrected registry-dependent (DRIP) potential as described in (Wen), which is based on the (Kolmogorov) potential and provides an improved prediction for forces. The total potential energy of a system is

$$E = \frac{1}{2} \sum_i \sum_{j \notin \text{layer } i} \phi_{ij}$$

$$\phi_{ij} = f_c(x_r) \left[ e^{-\lambda(r_{ij}-z_0)} \left[ C + f(\rho_{ij}) + g(\rho_{ij}, \{\alpha_{ij}^{(m)}\}) \right] - A \left( \frac{z_0}{r_{ij}} \right)^6 \right]$$

where the  $r^{-6}$  term models the attractive London dispersion, the exponential term is designed to capture the registry effect due to overlapping  $pi$  bonds, and  $f_c$  is a cutoff function.

This potential (DRIP) only provides the interlayer interactions between graphene layers. So, to perform a realistic simulation, it should be used in combination with an intralayer potential such as *REBO* and *Tersoff*. To keep the intralayer interactions unaffected, we should avoid applying DRIP to contribute energy to intralayer interactions. This can be achieved by assigning different molecular IDs to atoms in different layers, and DRIP is implemented such that only atoms with different molecular ID can interact with each other. For this purpose, *atom style* “molecular” or “full” has to be used.

On the other way around, *REBO* (*Tersoff* or any other potential used to provide the intralayer interactions) should not interfere with the interlayer interactions described by DRIP. This is typically automatically achieved using the commands provided in the *Examples* section above, since the cutoff distance for carbon-carbon interaction in the intralayer potentials (e.g. 2 Angstrom for *REBO*) is much smaller than the equilibrium layer distance of graphene layers (about 3.4 Angstrom). If you want, you can enforce this by assigning different atom types to atoms in different layers, and apply an intralayer potential to one atom type. See *pair\_hybrid* for details.

---

The *pair\_coeff* command for DRIP takes  $4+N$  arguments, where  $N$  is the number of LAMMPS atom types. The first three arguments must be fixed to be `**drip`, the fourth argument is the path to the DRIP parameter file, and the remaining  $N$  arguments specifying the mapping between element in the parameter file and atom types. For example, if your LAMMPS simulation has 3 atom types and you want all of them to be C, you would use the following *pair\_coeff* command:

```
pair_coeff * * drip C.drip C C C
```

If a mapping value is specified as NULL, the mapping is not performed. This could be useful when DRIP is used to model part of the system where other element exists. Suppose you have a hydrocarbon system, with C of atom type 1 and H of atom type 2, you can use the following command to inform DRIP not to model H atoms:

```
pair_style hybrid/overlay drip rebo
pair_coeff * * drip C.drip C NULL
pair_coeff * * rebo CH.airebo C H
```

---

**Note:** The potential parameters developed in (*Wen*) are provided with LAMMPS (see the “potentials” directory). Besides those in *Wen*, an additional parameter “normal\_cutoff”, specific to the LAMMPS implementation, is used to find the three nearest neighbors of an atom to construct the normal.

---

## 19.85.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* mix, shift, table, and tail options.

This pair style does not write their information to binary restart files, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

## 19.85.5 Restrictions

This pair style is part of the INTERLAYER package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

The *C.drip* parameter file provided with LAMMPS (see the “potentials” directory) is parameterized for metal *units*. You can use the DRIP potential with any LAMMPS units, but you would need to create your own custom parameter file with coefficients listed in the appropriate units, if your simulation does not use “metal” units.

## 19.85.6 Related commands

*pair\_style lebedeva\_z*, *pair\_style kolmogorov/crespi/z*, *pair\_style kolmogorov/crespi/full*, *pair\_style ilp/graphene/hbn*.

(Wen) M. Wen, S. Carr, S. Fang, E. Kaxiras, and E. B. Tadmor, Phys. Rev. B, 98, 235404 (2018)

(Kolmogorov) A. N. Kolmogorov, V. H. Crespi, Phys. Rev. B 71, 235415 (2005)

## 19.86 pair\_style dsmc command

### 19.86.1 Syntax

```
pair_style dsmc max_cell_size seed weighting Tref Nrecompute Nsample
```

- max\_cell\_size = global maximum cell size for DSMC interactions (distance units)
- seed = random # seed (positive integer)
- weighting = macroparticle weighting
- Tref = reference temperature (temperature units)
- Nrecompute = re-compute v\*sigma\_max every this many timesteps (timesteps)
- Nsample = sample this many times in recomputing v\*sigma\_max

### 19.86.2 Examples

```
pair_style dsmc 2.5 34387 10 1.0 100 20
pair_coeff * * 1.0
pair_coeff 1 1 1.0
```

### 19.86.3 Description

Style *dsmc* computes collisions between pairs of particles for a direct simulation Monte Carlo (DSMC) model following the exposition in (*Bird*). Each collision resets the velocities of the two particles involved. The number of pairwise collisions for each pair or particle types and the length scale within which they occur are determined by the parameters of the `pair_style` and `pair_coeff` commands.

Stochastic collisions are performed using the variable hard sphere (VHS) approach, with the user-defined `max_cell_size` value used as the maximum DSMC cell size, and reference cross-sections for collisions given using the `pair_coeff` command.

There is no pairwise energy or virial contributions associated with this pair style.

The following coefficient must be defined for each pair of atoms types via the `pair_coeff` command as in the examples above, or in the data file or restart files read by the `read_data` or `read_restart` commands:

- `sigma` (area units, i.e. distance-squared)

The global DSMC `max_cell_size` determines the maximum cell length used in the DSMC calculation. A structured mesh is overlaid on the simulation box such that an integer number of cells are created in each direction for each processor's sub-domain. Cell lengths are adjusted up to the user-specified maximum cell size.

---

To perform a DSMC simulation with LAMMPS, several additional options should be set in your input script, though LAMMPS does not check for these settings.

Since this pair style does not compute particle forces, you should use the “fix nve/noforce” time integration fix for the DSMC particles, e.g.

```
fix 1 all nve/noforce
```

This pair style assumes that all particles will be communicated to neighboring processors every timestep as they move. This makes it possible to perform all collisions between pairs of particles that are on the same processor. To ensure this occurs, you should use these commands:

```
neighbor 0.0 bin
neigh_modify every 1 delay 0 check no
atom_modify sort 0 0.0
communicate single cutoff 0.0
```

These commands ensure that LAMMPS communicates particles to neighboring processors every timestep and that no ghost atoms are created. The output statistics for a simulation run should indicate there are no ghost particles or neighbors.

In order to get correct DSMC collision statistics, users should specify a Gaussian velocity distribution when populating the simulation domain. Note that the default velocity distribution is uniform, which will not give good DSMC collision rates. Specify “dist gaussian” when using the `velocity` command as in the following:

```
velocity all create 594.6 87287 loop geom dist gaussian
```

---

## 19.86.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This pair style does not support the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file. Note that the user-specified random number seed is stored in the restart file, so when a simulation is restarted, each processor will re-initialize its random number generator the same way it did initially. This means the random forces will be random, but will not be the same as they would have been if the original simulation had continued past the restart time.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

## 19.86.5 Restrictions

This style is part of the MC package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.86.6 Related commands

*pair\_coeff*, *fix nve/noforce*, *neigh\_modify*, *neighbor*, *comm\_modify*

## 19.86.7 Default

none

**(Bird)** G. A. Bird, “Molecular Gas Dynamics and the Direct Simulation of Gas Flows” (1994).

## 19.87 pair\_style e3b command

### 19.87.1 Syntax

```
pair_style e3b Otype
```

- Otype = atom type for oxygen

```
pair_coeff * * keyword
```

- one or more keyword/value pairs must be appended.
- keyword = *preset* or *Ea* or *Eb* or *Ec* or *E2* or *K3* or *K2* or *Rs* or *Rc3* or *Rc2* or *bondL* or *neigh*
- If the *preset* keyword is given, no others are needed. Otherwise, all are mandatory except for *neigh*. The *neigh* keyword is always optional.

`preset` arg = 2011 or 2015 = which set of predefined parameters to use  
     2011 = use the potential parameters from (Tainter 2011)  
     2015 = use the potential parameters from (Tainter 2015)  
`Ea` arg = three-body energy for type A hydrogen bonding interactions (energy units)  
`Eb` arg = three-body energy for type B hydrogen bonding interactions (energy units)  
`Ec` arg = three-body energy for type C hydrogen bonding interactions (energy units)  
`E2` arg = two-body energy correction (energy units)  
`K3` arg = three-body exponential constant (inverse distance units)  
`K2` arg = two-body exponential constant (inverse distance units)  
`Rc3` arg = three-body cutoff (distance units)  
`Rc2` arg = two-body cutoff (distance units)  
`Rs` arg = three-body switching function cutoff (distance units)  
`bondL` arg = intramolecular OH bond length (distance units)  
`neigh` arg = approximate integer number of molecules within Rc3 of an oxygen atom

## 19.87.2 Examples

```

pair_style e3b 1
pair_coeff * * Ea 35.85 Eb -240.2 Ec 449.3 E2 108269.9 K3 1.907 K2 4.872 Rc3 5.2 Rc2 5.2
↳Rs 5.0 bondL 0.9572

pair_style hybrid/overlay e3b 1 lj/cut/tip4p/long 1 2 1 1 0.15 8.5
pair_coeff * * e3b preset 2011
  
```

Used in example input script:

```
examples/PACKAGES/e3b/in.e3b-tip4p2005
```

## 19.87.3 Description

The *e3b* style computes an "explicit three-body" (E3B) potential for water (Kumar 2008).

$$E = E_2 \sum_{i,j} e^{-k_2 r_{ij}} + E_A \sum_{\substack{i,j,k,\ell \\ \in \text{type A}}} f(r_{ij})f(r_{k\ell}) + E_B \sum_{\substack{i,j,k,\ell \\ \in \text{type B}}} f(r_{ij})f(r_{k\ell}) + E_C \sum_{\substack{i,j,k,\ell \\ \in \text{type C}}} f(r_{ij})f(r_{k\ell})$$

$$f(r) = e^{-k_3 r} s(r)$$

$$s(r) = \begin{cases} 1 & r < R_s \\ \frac{(R_f - r)^2 (R_f - 3R_s + 2r)}{(R_f - R_s)^3} & R_s \leq r \leq R_f \\ 0 & r > R_f \end{cases}$$

This potential was developed as a water model that includes the three-body cooperativity of hydrogen bonding explicitly. To use it in this way, it must be applied in conjunction with a conventional two-body water model, through pair style *hybrid/overlay*. The three body interactions are split into three types: A, B, and C. Type A corresponds to anti-cooperative double hydrogen bond donor interactions. Type B corresponds to the cooperative interaction of molecules that both donate and accept a hydrogen bond. Type C corresponds to anti-cooperative double hydrogen bond acceptor interactions. The three-body interactions are smoothly cutoff by the switching function  $s(r)$  between  $R_s$  and  $R_c3$ . The two-body interactions are designed to correct for the effective many-body interactions implicitly included in the conventional two-body potential. The two-body interactions are cut off sharply at  $R_c2$ , because  $K3$  is typically significantly smaller than  $K2$ . See (Kumar 2008) for more details.

Only a single *pair\_coeff* command is used with the *e3b* style and the first two arguments must be *\*\**. The oxygen atom type for the pair style is passed as the only argument to the *pair\_style* command, not in the *pair\_coeff* command. The hydrogen atom type is inferred from the ordering of the atoms.

---

**Note:** Every atom of type *Otype* must be part of a water molecule. Each water molecule must have consecutive IDs with the oxygen first. This pair style does not test that this criteria is met.

---

The *pair\_coeff* command must have at least one keyword/value pair, as described above. The *preset* keyword sets the potential parameters to the values used in (*Tainter 2011*) or (*Tainter 2015*). To use the water models defined in those references, the *e3b* style should always be used in conjunction with an *lj/cut/tip4p/long* style through *pair\_style hybrid/overlay*, as demonstrated in the second example above. The *preset 2011* option should be used with the *TIP4P water model*. The *preset 2015* option should be used with the *TIP4P/2005 water model*. If the *preset* keyword is used, no other keyword is needed. Changes to the preset parameters can be made by specifying the *preset* keyword followed by the specific parameter to change, like *Ea*. Note that the other keywords must come after *preset* in the *pair\_style* command. The *e3b* style can also be used to implement any three-body potential of the same form by specifying all the keywords except *neigh*: *Ea*, *Eb*, *Ec*, *E2*, *K3*, *K2*, *Rc3*, *Rc2*, *Rs*, and *bondL*. The keyword *bondL* specifies the intramolecular OH bond length of the water model being used. This is needed to include H atoms that are within the cutoff even when the attached oxygen atom is not.

This pair style allocates arrays sized according to the number of pairwise interactions within *Rc3*. To do this it needs an estimate for the number of water molecules within *Rc3* of an oxygen atom. This estimate defaults to 10 and can be changed using the *neigh* keyword, which takes an integer as an argument. If the *neigh* setting is too small, the simulation will fail with the error “neigh is too small”. If the *neigh* setting is too large, the pair style will use more memory than necessary.

This pair style tallies a breakdown of the total E3B potential energy into sub-categories, which can be accessed via the *compute pair* command as a vector of values of length 4. The 4 values correspond to the terms in the first equation above: the *E2* term, the *Ea* term, the *Eb* term, and the *Ec* term.

See the examples/PACKAGES/e3b directory for a complete example script.

---

## 19.87.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style is incompatible with *respa*.

---

## 19.87.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

This pair style requires a fixed number of atoms in the simulation, so it is incompatible with fixes like *fix deposit*. If the number of atoms changes between runs, this pair style must be re-initialized by calling the *pair\_style* and *pair\_coeffs* commands. This is not a fundamental limitation of the pair style, but the code currently does not support a variable number of atoms.

The *preset* keyword currently only works with real, metal, si, and cgs *units*.

### 19.87.6 Related commands

*pair\_coeff*, *compute pair*

### 19.87.7 Default

The option default for the *neigh* keyword is 10.

---

(Kumar) Kumar and Skinner, J. Phys. Chem. B, 112, 8311 (2008)

(Tainter 2011) Tainter, Pieniazek, Lin, and Skinner, J. Chem. Phys., 134, 184501 (2011)

(Tainter 2015) Tainter, Shi, and Skinner, 11, 2268 (2015)

## 19.88 pair\_style eam command

Accelerator Variants: *eam/gpu*, *eam/intel*, *eam/kk*, *eam/omp*, *eam/opt*

## 19.89 pair\_style eam/alloy command

Accelerator Variants: *eam/alloy/gpu*, *eam/alloy/intel*, *eam/alloy/kk*, *eam/alloy/omp*, *eam/alloy/opt*

## 19.90 pair\_style eam/cd command

## 19.91 pair\_style eam/cd/old command

## 19.92 pair\_style eam/fs command

## 19.93 pair\_style eam/he command

Accelerator Variants: *eam/fs/gpu*, *eam/fs/intel*, *eam/fs/kk*, *eam/fs/omp*, *eam/fs/opt*

### 19.93.1 Syntax

```
pair_style style
```

- style = *eam* or *eam/alloy* or *eam/cd* or *eam/cd/old* or *eam/fs* or *eam/he*



## 19.93.2 Examples

```

pair_style eam
pair_coeff * * cuu3
pair_coeff 1*3 1*3 niu3.eam

pair_style eam/alloy
pair_coeff * * ../potentials/NiAlH_jea.eam.alloy Ni Al Ni Ni

pair_style eam/cd
pair_coeff * * ../potentials/FeCr.cdeam Fe Cr

pair_style eam/fs
pair_coeff * * NiAlH_jea.eam.fs Ni Al Ni Ni

pair_style eam/he
pair_coeff * * PdHHe.eam.he Pd H He

```

## 19.93.3 Description

Style *eam* computes pairwise interactions for metals and metal alloys using embedded-atom method (EAM) potentials (*Daw*). The total energy  $E_i$  of an atom  $I$  is given by

$$E_i = F_\alpha \left( \sum_{j \neq i} \rho_\beta(r_{ij}) \right) + \frac{1}{2} \sum_{j \neq i} \phi_{\alpha\beta}(r_{ij})$$

where  $F$  is the embedding energy which is a function of the atomic electron density  $\rho$ ,  $\phi$  is a pair potential interaction, and  $\alpha$  and  $\beta$  are the element types of atoms  $I$  and  $J$ . The multi-body nature of the EAM potential is a result of the embedding energy term. Both summations in the formula are over all neighbors  $J$  of atom  $I$  within the cutoff distance.

The cutoff distance and the tabulated values of the functionals  $F$ ,  $\rho$ , and  $\phi$  are listed in one or more files which are specified by the *pair\_coeff* command. These are ASCII text files in a DYNAMO-style format which is described below. DYNAMO was the original serial EAM MD code, written by the EAM originators. Several DYNAMO potential files for different metals are included in the “potentials” directory of the LAMMPS distribution. All of these files are parameterized in terms of LAMMPS *metal units*.

---

**Note:** The *eam* style reads single-element EAM potentials in the DYNAMO *funcfl* format. Either single element or alloy systems can be modeled using multiple *funcfl* files and style *eam*. For the alloy case LAMMPS mixes the single-element potentials to produce alloy potentials, the same way that DYNAMO does. Alternatively, a single DYNAMO *setfl* file or Finnis/Sinclair EAM file can be used by LAMMPS to model alloy systems by invoking the *eam/alloy* or *eam/cd* or *eam/fs* or *eam/he* styles as described below. These files require no mixing since they specify alloy interactions explicitly.

---



---

**Note:** Note that unlike for other potentials, cutoffs for EAM potentials are not set in the *pair\_style* or *pair\_coeff* command; they are specified in the EAM potential files themselves. Likewise, the EAM potential files list atomic masses; thus you do not need to use the *mass* command to specify them.

---

There are several WWW sites that distribute and document EAM potentials stored in DYNAMO or other formats:

```
http://www.ctcms.nist.gov/potentials
http://cst-www.nrl.navy.mil/ccm6/ap
http://enpub.fulton.asu.edu/cms/potentials/main/main.htm
https://openkim.org
```

These potentials should be usable with LAMMPS, though the alternate formats would need to be converted to the DYNAMO format used by LAMMPS and described on this page. The NIST site is maintained by Chandler Becker (cbecker at nist.gov) who is good resource for info on interatomic potentials and file formats.

The OpenKIM Project at <https://openkim.org/browse/models/by-type> provides EAM potentials that can be used directly in LAMMPS with the *kim command* interface.

---

For style *eam*, potential values are read from a file that is in the DYNAMO single-element *funcfl* format. If the DYNAMO file was created by a Fortran program, it cannot have “D” values in it for exponents. C only recognizes “e” or “E” for scientific notation.

For style *eam* a potential file must be assigned to each I,I pair of atom types by using one or more *pair\_coeff* commands, each with a single argument:

- filename

Thus the following command

```
pair_coeff *2 1*2 cuu3.eam
```

will read the *cuu3* potential file and use the tabulated Cu values for F, phi, rho that it contains for type pairs 1,1 and 2,2 (type pairs 1,2 and 2,1 are ignored). See the *pair\_coeff* doc page for alternate ways to specify the path for the potential file. In effect, this makes atom types 1 and 2 in LAMMPS be Cu atoms. Different single-element files can be assigned to different atom types to model an alloy system. The mixing to create alloy potentials for type pairs with  $I \neq J$  is done automatically the same way that the serial DYNAMO code originally did it; you do not need to specify coefficients for these type pairs.

*Funcfl* files in the *potentials* directory of the LAMMPS distribution have an “.eam” suffix. A DYNAMO single-element *funcfl* file is formatted as follows:

- line 1: comment (ignored)
- line 2: atomic number, mass, lattice constant, lattice type (e.g. FCC)
- line 3: Nrho, drho, Nr, dr, cutoff

On line 2, all values but the mass are ignored by LAMMPS. The mass is in mass *units*, e.g. mass number or grams/mole for metal units. The cubic lattice constant is in Angstroms. On line 3, Nrho and Nr are the number of tabulated values in the subsequent arrays, drho and dr are the spacing in density and distance space for the values in those arrays, and the specified cutoff becomes the pairwise cutoff used by LAMMPS for the potential. The units of dr are Angstroms; I’m not sure of the units for drho - some measure of electron density.

Following the three header lines are three arrays of tabulated values:

- embedding function  $F(\rho)$  (Nrho values)
- effective charge function  $Z(r)$  (Nr values)
- density function  $\rho(r)$  (Nr values)

The values for each array can be listed as multiple values per line, so long as each array starts on a new line. For example, the individual  $Z(r)$  values are for  $r = 0, dr, 2*dr, \dots (Nr-1)*dr$ .

The units for the embedding function  $F$  are eV. The units for the density function  $\rho$  are the same as for drho (see above, electron density). The units for the effective charge  $Z$  are “atomic charge” or  $\sqrt{\text{Hartree} * \text{Bohr-radii}}$ . For

two interacting atoms  $i,j$  this is used by LAMMPS to compute the pair potential term in the EAM energy expression as  $r \cdot \phi$ , in units of eV-Angstroms, via the formula

$$r \cdot \phi = 27.2 \cdot 0.529 \cdot Z_i \cdot Z_j$$

where 1 Hartree = 27.2 eV and 1 Bohr = 0.529 Angstroms.

Style *eam/alloy* computes pairwise interactions using the same formula as style *eam*. However the associated *pair\_coeff* command reads a DYNAMO *setfl* file instead of a *funcfl* file. *Setfl* files can be used to model a single-element or alloy system. In the alloy case, as explained above, *setfl* files contain explicit tabulated values for alloy interactions. Thus they allow more generality than *funcfl* files for modeling alloys.

For style *eam/alloy*, potential values are read from a file that is in the DYNAMO multi-element *setfl* format, except that element names (Ni, Cu, etc) are added to one of the lines in the file. If the DYNAMO file was created by a Fortran program, it cannot have “D” values in it for exponents. C only recognizes “e” or “E” for scientific notation.

Only a single *pair\_coeff* command is used with the *eam/alloy* style which specifies a DYNAMO *setfl* file, which contains information for M elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the *pair\_coeff* command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of *setfl* elements to atom types

As an example, the potentials/NiAlH\_jea.eam.alloy file is a *setfl* file which has tabulated EAM values for 3 elements and their alloy interactions: Ni, Al, and H. See the *pair\_coeff* doc page for alternate ways to specify the path for the potential file. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Ni, and the fourth to be Al, you would use the following *pair\_coeff* command:

```
pair_coeff * * NiAlH_jea.eam.alloy Ni Ni Ni Al
```

The first 2 arguments must be \* \* so as to span all LAMMPS atom types. The first three Ni arguments map LAMMPS atom types 1,2,3 to the Ni element in the *setfl* file. The final Al argument maps LAMMPS atom type 4 to the Al element in the *setfl* file. Note that there is no requirement that your simulation use all the elements specified by the *setfl* file.

If a mapping value is specified as NULL, the mapping is not performed. This can be used when an *eam/alloy* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

*Setfl* files in the *potentials* directory of the LAMMPS distribution have an “.eam.alloy” suffix. A DYNAMO multi-element *setfl* file is formatted as follows:

- lines 1,2,3 = comments (ignored)
- line 4: Nelements Element1 Element2 ... ElementN
- line 5: Nrho, drho, Nr, dr, cutoff

In a DYNAMO *setfl* file, line 4 only lists Nelements = the # of elements in the *setfl* file. For LAMMPS, the element name (Ni, Cu, etc) of each element must be added to the line, in the order the elements appear in the file.

The meaning and units of the values in line 5 is the same as for the *funcfl* file described above. Note that the cutoff (in Angstroms) is a global value, valid for all pairwise interactions for all element pairings.

Following the 5 header lines are Nelements sections, one for each element, each with the following format:

- line 1 = atomic number, mass, lattice constant, lattice type (e.g. FCC)
- embedding function F(rho) (Nrho values)
- density function rho(r) (Nr values)

As with the *funcfl* files, only the mass (in mass *units*, e.g. mass number or grams/mole for metal units) is used by LAMMPS from the first line. The cubic lattice constant is in Angstroms. The F and rho arrays are unique to a single element and have the same format and units as in a *funcfl* file.

Following the Nelements sections, Nr values for each pair potential phi(r) array are listed for all i,j element pairs in the same format as other arrays. Since these interactions are symmetric (i,j = j,i) only phi arrays with i >= j are listed, in the following order: i,j = (1,1), (2,1), (2,2), (3,1), (3,2), (3,3), (4,1), ..., (Nelements, Nelements). Unlike the effective charge array Z(r) in *funcfl* files, the tabulated values for each phi function are listed in *setfl* files directly as r\*phi (in units of eV-Angstroms), since they are for atom pairs.

---

Style *eam/cd* is similar to the *eam/alloy* style, except that it computes alloy pairwise interactions using the concentration-dependent embedded-atom method (CD-EAM). This model can reproduce the enthalpy of mixing of alloys over the full composition range, as described in (*Stukowski*). Style *eam/cd/old* is an older, slightly different and slower two-site formulation of the model (*Caro*).

The pair\_coeff command is specified the same as for the *eam/alloy* style. However the DYNAMO *setfl* file must have two lines added to it, at the end of the file:

- line 1: Comment line (ignored)
- line 2: N Coefficient0 Coefficient1 ... CoefficientN

The last line begins with the degree N of the polynomial function h(x) that modifies the cross interaction between A and B elements. Then N+1 coefficients for the terms of the polynomial are then listed.

Modified EAM *setfl* files used with the *eam/cd* style must contain exactly two elements, i.e. in the current implementation the *eam/cd* style only supports binary alloys. The first and second elements in the input EAM file are always taken as the A and B species.

CD-EAM files in the *potentials* directory of the LAMMPS distribution have a “.cdeam” suffix.

---

Style *eam/fs* computes pairwise interactions for metals and metal alloys using a generalized form of EAM potentials due to Finnis and Sinclair (*Finnis*). Style *eam/he* is similar to *eam/fs* except that it allows for negative electron density in order to capture the behavior of helium in metals (*Zhou6*).

The total energy  $E_i$  of an atom I is given by

$$E_i = F_\alpha \left( \sum_{j \neq i} \rho_{\alpha\beta}(r_{ij}) \right) + \frac{1}{2} \sum_{j \neq i} \phi_{\alpha\beta}(r_{ij})$$

where  $\rho_{\alpha\beta}$  refers to the density contributed by a neighbor atom J of element  $\beta$  at the site of atom I of element  $\alpha$ . This has the same form as the EAM formula above, except that rho is now a functional specific to the elements of both atoms I and J, so that different elements can contribute differently to the total electron density at an atomic site depending on the identity of the element at that atomic site.

The associated *pair\_coeff* command for style *eam/fs* or *eam/he* reads a DYNAMO *setfl* file that has been extended to include additional  $\rho_{\alpha\beta}$  arrays of tabulated values. A discussion of how FS EAM differs from conventional EAM alloy potentials is given in (*Ackland1*). An example of such a potential is the same author’s Fe-P FS potential (*Ackland2*). Note that while FS potentials always specify the embedding energy with a square root dependence on the total density, the implementation in LAMMPS does not require that; the user can tabulate any functional form desired in the FS potential files.

For style *eam/fs* and *eam/he* the form of the pair\_coeff command is exactly the same as for style *eam/alloy*, e.g.

```
pair_coeff * * NiAlH_jea.eam.fs Ni Ni Ni Al
```

with N additional arguments after the filename, where N is the number of LAMMPS atom types. See the *pair\_coeff* doc page for alternate ways to specify the path for the potential file. The N values determine the mapping of LAMMPS atom types to EAM elements in the file, as described above for style *eam/alloy*. As with *eam/alloy*, if a mapping value is NULL, the mapping is not performed. This can be used when an *eam/fs* or *eam/he* potential is used as part of a *hybrid* pair style. The NULL values are used as placeholders for atom types that will be used with other potentials.

FS EAM and HE EAM files include more information than the DYNAMO *setfl* format files read by *eam/alloy*, in that i,j density functionals for all pairs of elements are included as needed by the Finnis/Sinclair formulation of the EAM.

FS EAM files in the *potentials* directory of the LAMMPS distribution have an “.eam.fs” suffix. They are formatted as follows:

- lines 1,2,3 = comments (ignored)
- line 4: Nelements Element1 Element2 ... ElementN
- line 5: Nrho, drho, Nr, dr, cutoff

The 5-line header section is identical to an EAM *setfl* file.

Following the header are Nelements sections, one for each element  $\beta$ , each with the following format:

- line 1 = atomic number, mass, lattice constant, lattice type (e.g. FCC)
- embedding function F(rho) (Nrho values)
- density function  $\rho_{1\beta}(r)$  for element  $\beta$  at element 1 (Nr values)
- density function  $\rho_{2\beta}(r)$  for element  $\beta$  at element 2
- ...
- density function  $\rho_{N_{elem}\beta}(r)$  for element  $\beta$  at element  $N_{elem}$

The units of these quantities in line 1 are the same as for *setfl* files. Note that the rho(r) arrays in Finnis/Sinclair can be asymmetric ( $\rho_{\alpha\beta}(r) \neq \rho_{\beta\alpha}(r)$ ) so there are Nelements<sup>2</sup> of them listed in the file.

Following the Nelements sections, Nr values for each pair potential phi(r) array are listed in the same manner (r\*phi, units of eV-Angstroms) as in EAM *setfl* files. Note that in Finnis/Sinclair, the phi(r) arrays are still symmetric, so only phi arrays for  $i \geq j$  are listed.

HE EAM files in the *potentials* directory of the LAMMPS distribution have an “.eam.he” suffix. They are formatted as follows:

- lines 1,2,3 = comments (ignored)
- line 4: Nelements Element1 Element2 ... ElementN
- line 5: Nrho, drho, Nr, dr, cutoff, rhomax

The 5-line header section is identical to an FS EAM file except that line 5 lists an additional value, rhomax. Unlike in FS EAM files where embedding energies F(rho) are always defined between  $\rho = 0$  and  $\rho = (Nrho - 1)drho$ , F(rho) in HE EAM files are defined between  $\rho = rhomin$  and  $\rho = rhomax$ . Since  $drho = (rhomax - rhomin)/(Nrho - 1)$ ,  $rhomin = rhomax - (Nrho - 1)drho$ . The embedding energies F(rho) are listed for  $\rho = rhomin, rhomin + drho, rhomin + 2drho, \dots, rhomax$ . This gives users additional flexibility to define a negative rhomin and therefore an embedding energy function that works for both positive and negative electron densities. The format and units of these sections are identical to the FS EAM files (see above).

---

New in version TBD.

The *eam*, *eam/alloy*, *eam/fs*, and *eam/he* pair styles support extraction of two per-atom quantities by the *fix pair* command. This allows the quantities to be output to files by the *dump* or otherwise processed by other LAMMPS commands.

The names of the two quantities are “rho” and “fp” for the density and derivative of the embedding energy for each atom. Neither quantity needs to be triggered by the *fix pair* command in order for these pair styles to calculate it.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.93.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, mixing is performed by LAMMPS as described above with the individual styles. You never need to specify a *pair\_coeff* command with I != J arguments for the eam styles.

This pair style does not support the *pair\_modify* shift, table, and tail options.

The eam pair styles do not write their information to *binary restart files*, since it is stored in tabulated potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

The eam pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

---

### 19.93.5 Restrictions

All of these styles are part of the MANYBODY package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.93.6 Related commands

*pair\_coeff*

### 19.93.7 Default

none

---

**(Ackland1)** Ackland, Condensed Matter (2005).

**(Ackland2)** Ackland, Mendeleev, Srolovitz, Han and Barashev, Journal of Physics: Condensed Matter, 16, S2629 (2004).

**(Daw)** Daw, Baskes, Phys Rev Lett, 50, 1285 (1983). Daw, Baskes, Phys Rev B, 29, 6443 (1984).

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(Finnis) Finnis, Sinclair, Philosophical Magazine A, 50, 45 (1984).

(Zhou6) Zhou, Bartelt, Sills, Physical Review B, 103, 014108 (2021).

(Stukowski) Stukowski, Sadigh, Erhart, Caro; Modeling Simulation Materials Science & Engineering, 7, 075005 (2009).

(Caro) A Caro, DA Crowson, M Caro; Phys Rev Lett, 95, 075702 (2005)

## 19.94 pair\_style edip command

Accelerator Variants: *edip/omp*

## 19.95 pair\_style edip/multi command

### 19.95.1 Syntax

```
pair_style style
```

- style = *edip* or *edip/multi*

### 19.95.2 Examples

```
pair_style edip
pair_coeff * * Si.edip Si
```

### 19.95.3 Description

The *edip* and *edip/multi* styles compute a 3-body *EDIP* potential which is popular for modeling silicon materials where it can have advantages over other models such as the *Stillinger-Weber* or *Tersoff* potentials. The *edip* style has been programmed for single element potentials, while *edip/multi* supports multi-element EDIP runs.

In EDIP, the energy  $E$  of a system of atoms is

$$E = \sum_{j \neq i} \phi_2(R_{ij}, Z_i) + \sum_{j \neq i} \sum_{k \neq i, k > j} \phi_3(R_{ij}, R_{ik}, Z_i)$$

$$\phi_2(r, Z) = A \left[ \left( \frac{B}{r} \right)^\rho - e^{-\beta Z^2} \right] \exp\left( \frac{\sigma}{r-a} \right)$$

$$\phi_3(R_{ij}, R_{ik}, Z_i) = \exp\left( \frac{\gamma}{R_{ij}-a} \right) \exp\left( \frac{\gamma}{R_{ik}-a} \right) h(\cos\theta_{ijk}, Z_i)$$

$$Z_i = \sum_{m \neq i} f(R_{im}) \quad f(r) = \begin{cases} 1 & r < c \\ \exp\left( \frac{\alpha}{1-x^{-3}} \right) & c < r < a \\ 0 & r > a \end{cases}$$

$$h(l, Z) = \lambda[(1 - e^{-Q(Z)(l+\tau(Z))^2}) + \eta Q(Z)(l + \tau(Z))^2]$$

$$Q(Z) = Q_0 e^{-\mu Z} \quad \tau(Z) = u_1 + u_2(u_3 e^{-u_4 Z} - e^{-2u_4 Z})$$

where  $\phi_2$  is a two-body term and  $\phi_3$  is a three-body term. The summations in the formula are over all neighbors J and K of atom I within a cutoff distance = a. Both terms depend on the local environment of atom I through its effective coordination number defined by Z, which is unity for a cutoff distance < c and gently goes to 0 at distance = a.

Only a single `pair_coeff` command is used with the *edip* style which specifies a EDIP potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of EDIP elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the potential file.

As an example, imagine a file `Si.edip` has EDIP values for Si.

EDIP files in the *potentials* directory of the LAMMPS distribution have a “.edip” suffix. Lines that are not blank or comments (starting with #) define parameters for a triplet of elements. The parameters in a single entry correspond to the two-body and three-body coefficients in the formula above:

- element 1 (the center atom in a 3-body interaction)
- element 2
- element 3
- A (energy units)
- B (distance units)
- cutoffA (distance units)
- cutoffC (distance units)
- $\alpha$
- $\beta$
- $\eta$
- $\gamma$  (distance units)
- *lambda* (energy units)
- $\mu$
- $\tau$
- $\sigma$  (distance units)
- Q0
- u1
- u2
- u3
- u4

The A, B, beta, sigma parameters are used only for two-body interactions. The eta, gamma, lambda, mu, Q0 and all u1 to u4 parameters are used only for three-body interactions. The alpha and cutoffC parameters are used for the coordination environment function only.

The EDIP potential file must contain entries for all the elements listed in the `pair_coeff` command. It can also contain entries for additional elements not being used in a particular simulation; LAMMPS ignores those entries.



For a single-element simulation, only a single entry is required (e.g. SiSiSi). For a two-element simulation, the file must contain 8 entries (for SiSiSi, SiSiC, SiCSi, SiCC, CSiSi, CSiC, CCSi, CCC), that specify EDIP parameters for all permutations of the two elements interacting in three-body configurations. Thus for 3 elements, 27 entries would be required, etc.

At the moment, only a single element parameterization is implemented. However, the author is not aware of other multi-element EDIP parameterization. If you know any and you are interest in that, please contact the author of the EDIP package.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.95.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.95.5 Restrictions

This pair style can only be used if LAMMPS was built with the MANYBODY package. See the [Build package](#) doc page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

The EDIP potential files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use the EDIP potential with any LAMMPS units, but you would need to create your own EDIP potential file with coefficients listed in the appropriate units if your simulation does not use “metal” units.

## 19.95.6 Related commands

*pair\_coeff*

## 19.95.7 Default

none

---

(EDIP) J F Justo et al, Phys Rev B 58, 2539 (1998).

## 19.96 pair\_style eff/cut command

### 19.96.1 Syntax

```
pair_style eff/cut cutoff keyword args ...
```

- cutoff = global cutoff for Coulombic interactions
- zero or more keyword/value pairs may be appended

keyword = *limit/eradius* or *pressure/evirials* or *ecp*

*limit/eradius* args = none

*pressure/evirials* args = none

*ecp* args = type element type element ...

type = LAMMPS atom type (1 to Ntypes)

element = element symbol (e.g. H, Si)

### 19.96.2 Examples

```
pair_style eff/cut 39.7
pair_style eff/cut 40.0 limit/eradius
pair_style eff/cut 40.0 limit/eradius pressure/evirials
pair_style eff/cut 40.0 ecp 1 Si 3 C
pair_coeff * *
pair_coeff 2 2 20.0
pair_coeff 1 s 0.320852 2.283269 0.814857
pair_coeff 3 p 22.721015 0.728733 1.103199 17.695345 6.693621
```

### 19.96.3 Description

This pair style contains a LAMMPS implementation of the electron Force Field (eFF) potential currently under development at Caltech, as described in (*Jaramillo-Botero*). The eFF for  $Z < 6$  was first introduced by (*Su*) in 2007. It has been extended to higher  $Z$ s by using effective core potentials (ECPs) that now cover up to second and third row p-block elements of the periodic table.

eFF can be viewed as an approximation to QM wave packet dynamics and Fermionic molecular dynamics, combining the ability of electronic structure methods to describe atomic structure, bonding, and chemistry in materials, and of plasma methods to describe nonequilibrium dynamics of large systems with a large number of highly excited electrons. Yet, eFF relies on a simplification of the electronic wave function in which electrons are described as floating Gaussian

wave packets whose position and size respond to the various dynamic forces between interacting classical nuclear particles and spherical Gaussian electron wave packets. The wave function is taken to be a Hartree product of the wave packets. To compensate for the lack of explicit antisymmetry in the resulting wave function, a spin-dependent Pauli potential is included in the Hamiltonian. Substituting this wave function into the time-dependent Schrodinger equation produces equations of motion that correspond - to second order - to classical Hamiltonian relations between electron position and size, and their conjugate momenta. The N-electron wave function is described as a product of one-electron Gaussian functions, whose size is a dynamical variable and whose position is not constrained to a nuclear center. This form allows for straightforward propagation of the wave function, with time, using a simple formulation from which the equations of motion are then integrated with conventional MD algorithms. In addition to this spin-dependent Pauli repulsion potential term between Gaussians, eFF includes the electron kinetic energy from the Gaussians. These two terms are based on first-principles quantum mechanics. On the other hand, nuclei are described as point charges, which interact with other nuclei and electrons through standard electrostatic potential forms.

The full Hamiltonian (shown below), contains then a standard description for electrostatic interactions between a set of delocalized point and Gaussian charges which include, nuclei-nuclei (NN), electron-electron (ee), and nuclei-electron (Ne). Thus, eFF is a mixed QM-classical mechanics method rather than a conventional force field method (in which electron motions are averaged out into ground state nuclear motions, i.e a single electronic state, and particle interactions are described via empirically parameterized interatomic potential functions). This makes eFF uniquely suited to simulate materials over a wide range of temperatures and pressures where electronically excited and ionized states of matter can occur and coexist. Furthermore, the interactions between particles -nuclei and electrons- reduce to the sum of a set of effective pairwise potentials in the eFF formulation. The *eff/cut* style computes the pairwise Coulomb interactions between nuclei and electrons ( $E_{NN}, E_{Ne}, E_{ee}$ ), and the quantum-derived Pauli ( $E_{PR}$ ) and Kinetic energy interactions potentials between electrons ( $E_{KE}$ ) for a total energy expression given as,

$$U(R, r, s) = E_{NN}(R) + E_{Ne}(R, r, s) + E_{ee}(r, s) + E_{KE}(r, s) + E_{PR}(\uparrow\downarrow, S)$$

The individual terms are defined as follows:

$$E_{KE} = \frac{\hbar^2}{m_e} \sum_i \frac{3}{2s_i^2}$$

$$E_{NN} = \frac{1}{4\pi\epsilon_0} \sum_{i<j} \frac{Z_i Z_j}{R_{ij}}$$

$$E_{Ne} = -\frac{1}{4\pi\epsilon_0} \sum_{i,j} \frac{Z_i}{R_{ij}} \text{Erf} \left( \frac{\sqrt{2}R_{ij}}{s_j} \right)$$

$$E_{ee} = \frac{1}{4\pi\epsilon_0} \sum_{i<j} \frac{1}{r_{ij}} \text{Erf} \left( \frac{\sqrt{2}r_{ij}}{\sqrt{s_i^2 + s_j^2}} \right)$$

$$E_{Pauli} = \sum_{\sigma_i=\sigma_j} E(\uparrow\uparrow)_{ij} + \sum_{\sigma_i \neq \sigma_j} E(\uparrow\downarrow)_{ij}$$

where,  $s_i$  correspond to the electron sizes, the  $\sigma_i$ 's to the fixed spins of the electrons,  $Z_i$  to the charges on the nuclei,  $R_{ij}$  to the distances between the nuclei or the nuclei and electrons, and  $r_{ij}$  to the distances between electrons. For additional details see (*Jaramillo-Botero*).

The overall electrostatics energy is given in Hartree units of energy by default and can be modified by an energy-conversion constant, according to the units chosen (see *electron\_units*). The cutoff  $R_c$ , given in Bohrs (by default), truncates the interaction distance. The recommended cutoff for this pair style should follow the minimum image criterion, i.e. half of the minimum unit cell length.

Style *eff/long* (not yet available) computes the same interactions as style *eff/cut* except that an additional damping factor is applied so it can be used in conjunction with the *kpace\_style* command and its *ewald* or *pppm* option. The Coulombic cutoff specified for this style means that pairwise interactions within this distance are computed directly; interactions outside that distance are computed in reciprocal space.

This potential is designed to be used with *atom\_style electron* definitions, in order to handle the description of systems with interacting nuclei and explicit electrons.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- cutoff (distance units)

For *eff/cut*, the cutoff coefficient is optional. If it is not used (as in some of the examples above), the default global value specified in the *pair\_style* command is used.

For *eff/long* (not yet available) no cutoff will be specified for an individual I,J type pair via the *pair\_coeff* command. All type pairs use the same global cutoff specified in the *pair\_style* command.

---

The *limit/radius* and *pressure/evirials* keywords are optional. Neither or both must be specified. If not specified they are unset.

The *limit/radius* keyword is used to restrain electron size from becoming excessively diffuse at very high temperatures where the Gaussian wave packet representation breaks down, and from expanding as free particles to infinite size. If unset, electron radius is free to increase without bounds. If set, a restraining harmonic potential of the form  $E = 1/2k_{ss}s^2$  for  $s > L_{box}/2$ , where  $k_{ss} = 1$  Hartrees/Bohr<sup>2</sup>, is applied on the electron radius.

The *pressure/evirials* keyword is used to control between two types of pressure computation: if unset, the computed pressure does not include the electronic radial virials contributions to the total pressure (scalar or tensor). If set, the computed pressure will include the electronic radial virial contributions to the total pressure (scalar and tensor).

The *ecp* keyword is used to associate an ECP representation for a particular atom type. The ECP captures the orbital overlap between a core pseudo particle and valence electrons within the Pauli repulsion. A list of type:element-symbol pairs may be provided for all ECP representations, after the “ecp” keyword.

---

**Note:** Default ECP parameters are provided for C, N, O, Al, and Si. Users can modify these using the *pair\_coeff* command as exemplified above. For this, the User must distinguish between two different functional forms supported, one that captures the orbital overlap assuming the s-type core interacts with an s-like valence electron (s-s) and another that assumes the interaction is s-p. For systems that exhibit significant p-character (e.g. C, N, O) the s-p form is recommended. The “s” ECP form requires 3 parameters and the “p” 5 parameters.

---

**Note:** there are two different pressures that can be reported for eFF when defining this *pair\_style*, one (default) that considers electrons do not contribute radial virial components (i.e. electrons treated as incompressible ‘rigid’ spheres) and one that does. The radial electronic contributions to the virials are only tallied if the flexible pressure option is set, and this will affect both global and per-atom quantities. In principle, the true pressure of a system is somewhere in between the rigid and the flexible eFF pressures, but, for most cases, the difference between these two pressures will not be significant over long-term averaged runs (i.e. even though the energy partitioning changes, the total energy remains similar).

---

**Note:** This implementation of eFF gives a reasonably accurate description for systems containing nuclei from  $Z = 1-6$  in “all electron” representations. For systems with increasingly non-spherical electrons, Users should use the ECP representations. ECPs are now supported and validated for most of the second and third row elements of the p-block. Predefined parameters are provided for C, N, O, Al, and Si. The ECP captures the orbital overlap between the core and valence electrons (i.e. Pauli repulsion) with one of the functional forms:

---

$$E_{P_{aulli}(ECP_s)} = p_1 \exp\left(-\frac{p_2 r^2}{p_3 + s^2}\right)$$

$$E_{P_{aulli}(ECP_p)} = p_1 \left(\frac{2}{p_2/s + s/p_2}\right) (r - p_3 s)^2 \exp\left[-\frac{p_4 (r - p_3 s)^2}{p_5 + s^2}\right]$$

Where the first form correspond to core interactions with s-type valence electrons and the second to core interactions with p-type valence electrons.

The current version adds full support for models with fixed-core and ECP definitions. to enable larger timesteps (i.e. by avoiding the high frequency vibrational modes -translational and radial- of the 2 s electrons), and in the ECP case to reduce the increased orbital complexity in higher Z elements (up to  $Z < 18$ ). A fixed-core should be defined with a mass that includes the corresponding nuclear mass plus the 2 s electrons in atomic mass units ( $2 \times 5.4857990943 \times 10^{-4}$ ), and a radius equivalent to that of minimized 1s electrons (see examples under /examples/PACKAGES/eff/fixed-core). An pseudo-core should be described with a mass that includes the corresponding nuclear mass, plus all the core electrons (i.e. no outer shell electrons), and a radius equivalent to that of a corresponding minimized full-electron system. The charge for a pseudo-core atom should be given by the number of outer shell electrons.

In general, eFF excels at computing the properties of materials in extreme conditions and tracing the system dynamics over multi-picosecond timescales; this is particularly relevant where electron excitations can change significantly the nature of bonding in the system. It can capture with surprising accuracy the behavior of such systems because it describes consistently and in an unbiased manner many different kinds of bonds, including covalent, ionic, multicenter, ionic, and plasma, and how they interconvert and/or change when they become excited. eFF also excels in computing the relative thermochemistry of isodemic reactions and conformational changes, where the bonds of the reactants are of the same type as the bonds of the products. eFF assumes that kinetic energy differences dominate the overall exchange energy, which is true when the electrons present are nearly spherical and nodeless and valid for covalent compounds such as dense hydrogen, hydrocarbons, and diamond; alkali metals (e.g. lithium), alkali earth metals (e.g. beryllium) and semimetals such as boron; and various compounds containing ionic and/or multicenter bonds, such as boron dihydride.

---

#### 19.96.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , the cutoff distance for the *eff/cut* style can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

The *pair\_modify* shift option is not relevant for these pair styles.

The *eff/long* (not yet available) style supports the *pair\_modify* table option for tabulation of the short-range portion of the long-range Coulombic interaction.

These pair styles do not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

These pair styles write their information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

---

## 19.96.5 Restrictions

These pair styles will only be enabled if LAMMPS is built with the EFF package. It will only be enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These pair styles require that particles store electron attributes such as radius, radial velocity, and radial force, as defined by the *atom\_style*. The *electron* atom style does all of this.

These pair styles require you to use the *comm\_modify vel yes* command so that velocities are stored by ghost atoms.

## 19.96.6 Related commands

*pair\_coeff*

## 19.96.7 Default

If not specified, *limit\_radius* = 0 and *pressure\_with\_evirials* = 0.

---

**(Su)** Su and Goddard, Excited Electron Dynamics Modeling of Warm Dense Matter, Phys Rev Lett, 99:185003 (2007).

**(Jaramillo-Botero)** Jaramillo-Botero, Su, Qi, Goddard, Large-scale, Long-term Non-adiabatic Electron Molecular Dynamics for Describing Material Properties and Phenomena in Extreme Environments, J Comp Chem, 32, 497-512 (2011).

## 19.97 pair\_style eim command

Accelerator Variants: *eim/omp*

### 19.97.1 Syntax

```
pair_style style
```

- style = *eim*

### 19.97.2 Examples

```
pair_style eim
pair_coeff * * Na Cl ../potentials/ffield.eim Na Cl
pair_coeff * * Na Cl ffield.eim Na Na Na Cl
pair_coeff * * Na Cl ../potentials/ffield.eim Cl NULL Na
```

### 19.97.3 Description

Style *eim* computes pairwise interactions for ionic compounds using embedded-ion method (EIM) potentials (Zhou). The energy of the system  $E$  is given by

$$E = \frac{1}{2} \sum_{i=1}^N \sum_{j=i_1}^{i_N} \phi_{ij}(r_{ij}) + \sum_{i=1}^N E_i(q_i, \sigma_i)$$

The first term is a double pairwise sum over the  $J$  neighbors of all  $I$  atoms, where  $\phi_{ij}$  is a pair potential. The second term sums over the embedding energy  $E_i$  of atom  $I$ , which is a function of its charge  $q_i$  and the electrical potential  $\sigma_i$  at its location.  $E_i$ ,  $q_i$ , and  $\sigma_i$  are calculated as

$$q_i = \sum_{j=i_1}^{i_N} \eta_{ji}(r_{ij})$$

$$\sigma_i = \sum_{j=i_1}^{i_N} q_j \cdot \psi_{ij}(r_{ij})$$

$$E_i(q_i, \sigma_i) = \frac{1}{2} \cdot q_i \cdot \sigma_i$$

where  $\eta_{ji}$  is a pairwise function describing electron flow from atom  $I$  to atom  $J$ , and  $\psi_{ij}$  is another pairwise function. The multi-body nature of the EIM potential is a result of the embedding energy term. A complete list of all the pair functions used in EIM is summarized below

$$\phi_{ij}(r) = \begin{cases} \left[ \frac{E_{b,ij}\beta_{ij}}{\beta_{ij}-\alpha_{ij}} \exp\left(-\alpha_{ij}\frac{r-r_{e,ij}}{r_{e,ij}}\right) - \frac{E_{b,ij}\alpha_{ij}}{\beta_{ij}-\alpha_{ij}} \exp\left(-\beta_{ij}\frac{r-r_{e,ij}}{r_{e,ij}}\right) \right] f_c(r, r_{e,ij}, r_c, \phi, ij), & p_{ij} = 1 \\ \left[ \frac{E_{b,ij}\beta_{ij}}{\beta_{ij}-\alpha_{ij}} \left(\frac{r_{e,ij}}{r}\right)^{\alpha_{ij}} - \frac{E_{b,ij}\alpha_{ij}}{\beta_{ij}-\alpha_{ij}} \left(\frac{r_{e,ij}}{r}\right)^{\beta_{ij}} \right] f_c(r, r_{e,ij}, r_c, \phi, ij), & p_{ij} = 2 \end{cases}$$

$$\eta_{ji} = A_{\eta,ij} (\chi_j - \chi_i) f_c(r, r_s, \eta, ij, r_c, \eta, ij)$$

$$\psi_{ij}(r) = A_{\psi,ij} \exp(-\zeta_{ij}r) f_c(r, r_s, \psi, ij, r_c, \psi, ij)$$

$$f_c(r, r_p, r_c) = 0.510204 \cdot \operatorname{erfc}\left[\frac{1.64498(2r - r_p - r_c)}{r_c - r_p}\right] - 0.010204$$

Here  $E_b, r_e, r(c, \phi), \alpha, \beta, A(\psi), \zeta, r(s, \psi), r(c, \psi), A(\eta), r(s, \eta), r(c, \eta), \chi$ , and pair function type  $p$  are parameters, with subscripts  $ij$  indicating the two species of atoms in the atomic pair.

---

**Note:** Even though the EIM potential is treating atoms as charged ions, you should not use a LAMMPS *atom\_style* that stores a charge on each atom and thus requires you to assign a charge to each atom, e.g. the *charge* or *full* atom styles. This is because the EIM potential infers the charge on an atom from the equation above for  $q_i$ ; you do not assign charges explicitly.

---

All the EIM parameters are listed in a potential file which is specified by the *pair\_coeff* command. This is an ASCII text file in a format described below. The “field.eim” file included in the “potentials” directory of the LAMMPS distribution currently includes nine elements Li, Na, K, Rb, Cs, F, Cl, Br, and I. A system with any combination of these elements can be modeled. This file is parameterized in terms of LAMMPS *metal units*.

Note that unlike other potentials, cutoffs for EIM potentials are not set in the *pair\_style* or *pair\_coeff* command; they are specified in the EIM potential file itself. Likewise, the EIM potential file lists atomic masses; thus you do not need to use the *mass* command to specify them.

Only a single *pair\_coeff* command is used with the *eim* style which specifies an EIM potential file and the element(s) to extract information for. The EIM elements are mapped to LAMMPS atom types by specifying  $N$  additional arguments after the filename in the *pair\_coeff* command, where  $N$  is the number of LAMMPS atom types:

- Elem1, Elem2, ...
- EIM potential file
- N element names = mapping of EIM elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the potential file.

As an example like one of those above, suppose you want to model a system with Na and Cl atoms. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Na, and the fourth to be Cl, you would use the following `pair_coeff` command:

```
pair_coeff * * Na Clffield.eim Na Na Na Cl
```

The first 2 arguments must be `**` so as to span all LAMMPS atom types. The filename is the EIM potential file. The Na and Cl arguments (before the file name) are the two elements for which info will be extracted from the potential file. The first three trailing Na arguments map LAMMPS atom types 1,2,3 to the EIM Na element. The final Cl argument maps LAMMPS atom type 4 to the EIM Cl element.

If a mapping value is specified as NULL, the mapping is not performed. This can be used when an *eim* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

The `ffield.eim` file in the *potentials* directory of the LAMMPS distribution is formatted as follows:

Lines starting with # are comments and are ignored by LAMMPS. Lines starting with “global:” include three global values. The first value divides the cations from anions, i.e., any elements with electronegativity above this value are viewed as anions, and any elements with electronegativity below this value are viewed as cations. The second and third values are related to the cutoff function - i.e. the 0.510204, 1.64498, and 0.010204 shown in the above equation can be derived from these values.

Lines starting with “element:” are formatted as follows: name of element, atomic number, atomic mass, electronic negativity, atomic radius (LAMMPS ignores it), ionic radius (LAMMPS ignores it), cohesive energy (LAMMPS ignores it), and q0 (must be 0).

Lines starting with “pair:” are entered as: element 1, element 2, r\_(c,phi), r\_(c,phi) (redundant for historical reasons), E\_b, r\_e, alpha, beta, r\_(c,eta), A\_(eta), r\_(s,eta), r\_(c,psi), A\_(psi), zeta, r\_(s,psi), and p.

The lines in the file can be in any order; LAMMPS extracts the info it needs.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---



## 19.97.4 Restrictions

This style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package.

## 19.97.5 Related commands

*pair\_coeff*

## 19.97.6 Default

none

(Zhou) Zhou, submitted for publication (2010). Please contact Xiaowang Zhou (Sandia) for details via email at xzhou at sandia.gov.

## 19.98 pair\_style exp6/rx command

Accelerator Variants: *exp6/rx/kk*

### 19.98.1 Syntax

```
pair_style exp6/rx cutoff ...
```

- cutoff = global cutoff for DPD interactions (distance units)
- weighting = fractional or molecular (optional)

### 19.98.2 Examples

```
pair_style exp6/rx 10.0
pair_style exp6/rx 10.0 fractional
pair_style exp6/rx 10.0 molecular
pair_coeff * * exp6.params h2o h2o exponent 1.0 1.0 10.0
pair_coeff * * exp6.params h2o 1fluid exponent 1.0 1.0 10.0
pair_coeff * * exp6.params 1fluid 1fluid exponent 1.0 1.0 10.0
pair_coeff * * exp6.params 1fluid 1fluid none 10.0
pair_coeff * * exp6.params 1fluid 1fluid polynomial filename 10.0
```

### 19.98.3 Description

Style *exp6/rx* is used in reaction DPD simulations, where the coarse-grained (CG) particles are composed of  $m$  species whose reaction rate kinetics are determined from a set of  $n$  reaction rate equations through the *fix rx* command. The species of one CG particle can interact with a species in a neighboring CG particle through a site-site interaction potential model. The *exp6/rx* style computes an exponential-6 potential given by

$$U_{ij}(r) = \frac{\epsilon}{\alpha - 6} \left\{ 6 \exp\left[\alpha\left(1 - \frac{r_{ij}}{R_m}\right)\right] - \alpha\left(\frac{R_m}{r_{ij}}\right)^6 \right\}$$

where the  $\epsilon$  parameter determines the depth of the potential minimum located at  $R_m$ , and  $\alpha$  determines the softness of the repulsion.

The coefficients must be defined for each species in a given particle type via the *pair\_coeff* command as in the examples above, where the first argument is the filename that includes the exponential-6 parameters for each species. The file includes the species tag followed by the  $\alpha$ ,  $\epsilon$  and  $R_m$  parameters. The format of the file is described below.

The second and third arguments specify the site-site interaction potential between two species contained within two different particles. The species tags must either correspond to the species defined in the reaction kinetics files specified with the *fix rx* command or they must correspond to the tag “1fluid”, signifying interaction with a product species mixture determined through a one-fluid approximation. The interaction potential is weighted by the geometric average of either the mole fraction concentrations or the number of molecules associated with the interacting coarse-grained particles (see the *fractional* or *molecular* weighting pair style options). The coarse-grained potential is stored before and after the reaction kinetics solver is applied, where the difference is defined to be the internal chemical energy (uChem).

The fourth argument specifies the type of scaling that will be used to scale the EXP-6 parameters as reactions occur. Currently, there are three scaling options: *exponent*, *polynomial* and *none*.

Exponent scaling requires two additional arguments for scaling the  $R_m$  and  $\epsilon$  parameters, respectively. The scaling factor is computed by  $\text{phi}^{\text{exponent}}$ , where phi is the number of molecules represented by the coarse-grain particle and exponent is specified as a pair coefficient argument for  $R_m$  and  $\epsilon$ , respectively. The  $R_m$  and  $\epsilon$  parameters are multiplied by the scaling factor to give the scaled interaction parameters for the CG particle.

Polynomial scaling requires a filename to be specified as a pair coeff argument. The file contains the coefficients to a fifth order polynomial for the  $\alpha$ ,  $\epsilon$  and  $R_m$  parameters that depend upon phi (the number of molecules represented by the CG particle). The format of a polynomial file is provided below.

The *none* option to the scaling does not have any additional pair coeff arguments. This is equivalent to specifying the *exponent* option with  $R_m$  and  $\epsilon$  exponents of 0.0 and 0.0, respectively.

The final argument specifies the interaction cutoff (optional).

---

The format of a tabulated file is as follows (without the parenthesized comments):

```
# exponential-6 parameters for various species      (one or more comment or blank lines)
h2o  exp6  11.00  0.02  3.50                        (species, exp6, alpha, Rm, epsilon)
no2  exp6  13.60  0.01  3.70
...
co2  exp6  13.00  0.03  3.20
```

The format of the polynomial scaling file as follows (without the parenthesized comments):

```
# POLYNOMIAL FILE          (one or more comment or blank lines)

# General Functional Form:
# A*phi^5 + B*phi^4 + C*phi^3 + D*phi^2 + E*phi + F
```

```
#
# Parameter A      B      C      D      E      F
#                (blank)
alpha      0.0000  0.00000  0.00008  0.04955 -0.73804 13.63201
epsilon    0.0000  0.00478 -0.06283  0.24486 -0.33737  2.60097
rm         0.0001 -0.00118 -0.00253  0.05812 -0.00509  1.50106
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections.

Following a blank line, the next N lines list the species and their corresponding parameters. The first argument is the species tag, the second argument is the exp6 tag, the third argument is the  $\alpha$  parameter (energy units), the fourth argument is the  $\epsilon$  parameter (energy-distance<sup>6</sup> units), and the fifth argument is the  $R_m$  parameter (distance units). If a species tag of “1fluid” is listed as a pair coefficient, a one-fluid approximation is specified where a concentration-dependent combination of the parameters is computed through the following equations:

$$R_m^3 = \sum_a \sum_b x_a x_b R_{m,ab}^3$$

$$\epsilon = \frac{1}{R_m^3} \sum_a \sum_b x_a x_b \epsilon_{ab} R_{m,ab}^3$$

$$\alpha = \frac{1}{\epsilon R_m^3} \sum_a \sum_b x_a x_b \alpha_{ab} \epsilon_{ab} R_{m,ab}^3$$

where

$$\epsilon_{ab} = \sqrt{\epsilon_a \epsilon_b}$$

$$R_{m,ab} = \frac{R_{m,a} + R_{m,b}}{2}$$

$$\alpha_{ab} = \sqrt{\alpha_a \alpha_b}$$

and  $x_a$  and  $x_b$  are the mole fractions of a and b, respectively, which comprise the gas mixture.

## 19.98.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This style does not support the *pair\_modify* shift option for the energy of the exp() and 1/r<sup>6</sup> portion of the pair interaction.

This style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure for the A,C terms in the pair interaction.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

### 19.98.5 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.98.6 Related commands

*pair\_coeff*

### 19.98.7 Default

fractional weighting

## 19.99 pair\_style extep command

### 19.99.1 Syntax

```
pair_style extep
```

### 19.99.2 Examples

```
pair_style extep  
pair_coeff * * BN.extep B N
```

### 19.99.3 Description

Style *extep* computes the Extended Tersoff Potential (ExTeP) interactions as described in (*Los2017*).

---

### 19.99.4 Restrictions

none

### 19.99.5 Related commands

*pair\_tersoff* <*pair tersoff*>

### 19.99.6 Default

none

---

(Los2017) J. H. Los et al. “Extended Tersoff potential for boron nitride: Energetics and elastic properties of pristine and defective h-BN”, Phys. Rev. B 96 (184108), 2017.

## 19.100 pair\_style lj/cut/soft command

Accelerator Variants: *lj/cut/soft/omp*

## 19.101 pair\_style lj/cut/coul/cut/soft command

Accelerator Variants: *lj/cut/coul/cut/soft/omp*

## 19.102 pair\_style lj/cut/coul/long/soft command

Accelerator Variants: *lj/cut/coul/long/soft/omp*

## 19.103 pair\_style lj/cut/tip4p/long/soft command

Accelerator Variants: *lj/cut/tip4p/long/soft/omp*

## 19.104 pair\_style lj/charmm/coul/long/soft command

Accelerator Variants: *lj/charmm/coul/long/soft/omp*

## 19.105 pair\_style lj/class2/soft command

## 19.106 pair\_style lj/class2/coul/cut/soft command

## 19.107 pair\_style lj/class2/coul/long/soft command

## 19.108 pair\_style coul/cut/soft command

Accelerator Variants: *coul/cut/soft/omp*

## 19.109 pair\_style coul/long/soft command

Accelerator Variants: *coul/long/soft/omp*

## 19.110 pair\_style tip4p/long/soft command

Accelerator Variants: *tip4p/long/soft/omp*

## 19.111 pair\_style morse/soft command

### 19.111.1 Syntax

```
pair_style style args
```

- style = *lj/cut/soft* or *lj/cut/coul/cut/soft* or *lj/cut/coul/long/soft* or *lj/cut/tip4p/long/soft* or *lj/charmm/coul/long/soft* or *lj/class2/soft* or *lj/class2/coul/cut/soft* or *lj/class2/coul/long/soft* or *coul/cut/soft* or *coul/long/soft* or *tip4p/long/soft* or *morse/soft*
- args = list of arguments for a particular style

```
lj/cut/soft args = n alpha_lj cutoff
  n, alpha_LJ = parameters of soft-core potential
  cutoff = global cutoff for Lennard-Jones interactions (distance units)
lj/cut/coul/cut/soft args = n alpha_LJ alpha_C cutoff (cutoff2)
  n, alpha_LJ, alpha_C = parameters of soft-core potential
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)
lj/cut/coul/long/soft args = n alpha_LJ alpha_C cutoff
  n, alpha_LJ, alpha_C = parameters of the soft-core potential
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)
lj/cut/tip4p/long/soft args = otype htype btype atype qdist n alpha_LJ alpha_C cutoff_
↳(cutoff2)
  otype,htype = atom types for TIP4P O and H
  btype,atype = bond and angle types for TIP4P waters
  qdist = distance from O atom to massless charge (distance units)
  n, alpha_LJ, alpha_C = parameters of the soft-core potential
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)
lj/charmm/coul/long/soft args = n alpha_LJ alpha_C inner outer (cutoff)
  n, alpha_LJ, alpha_C = parameters of the soft-core potential
  inner, outer = global switching cutoffs for LJ (and Coulombic if only 5 args)
  cutoff = global cutoff for Coulombic (optional, outer is Coulombic cutoff if only 5_
↳args)
lj/class2/soft args = n alpha_lj cutoff
  n, alpha_LJ = parameters of soft-core potential
  cutoff = global cutoff for Lennard-Jones interactions (distance units)
lj/class2/coul/cut/soft args = n alpha_LJ alpha_C cutoff (cutoff2)
  n, alpha_LJ, alpha_C = parameters of soft-core potential
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
```

```

  cutoff2 = global cutoff for Coulombic (optional) (distance units)
lj/class2/coul/long/soft args = n alpha_LJ alpha_C cutoff (cutoff2)
  n, alpha_LJ, alpha_C = parameters of soft-core potential
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)
coul/cut/soft args = n alpha_C cutoff
  n, alpha_C = parameters of the soft-core potential
  cutoff = global cutoff for Coulomb interactions (distance units)
coul/long/soft args = n alpha_C cutoff
  n, alpha_C = parameters of the soft-core potential
  cutoff = global cutoff for Coulomb interactions (distance units)
tip4p/long/soft args = otype htype btype atype qdist n alpha_C cutoff
  otype,htype = atom types for TIP4P O and H
  btype,atype = bond and angle types for TIP4P waters
  qdist = distance from O atom to massless charge (distance units)
  n, alpha_C = parameters of the soft-core potential
  cutoff = global cutoff for Coulomb interactions (distance units)
morse/soft args = n lf cutoff
  n = soft-core parameter
  lf = transformation range is lf < lambda < 1
  cutoff = global cutoff for Morse interactions (distance units)

```

## 19.111.2 Examples

```

pair_style lj/cut/soft 2.0 0.5 9.5
pair_coeff * * 0.28 3.1 1.0
pair_coeff 1 1 0.28 3.1 1.0 9.5

pair_style lj/cut/coul/cut/soft 2.0 0.5 10.0 9.5
pair_style lj/cut/coul/cut/soft 2.0 0.5 10.0 9.5 9.5
pair_coeff * * 0.28 3.1 1.0
pair_coeff 1 1 0.28 3.1 0.5 10.0
pair_coeff 1 1 0.28 3.1 0.5 10.0 9.5

pair_style lj/cut/coul/long/soft 2.0 0.5 10.0 9.5
pair_style lj/cut/coul/long/soft 2.0 0.5 10.0 9.5 9.5
pair_coeff * * 0.28 3.1 1.0
pair_coeff 1 1 0.28 3.1 0.0 10.0
pair_coeff 1 1 0.28 3.1 0.0 10.0 9.5

pair_style lj/cut/tip4p/long/soft 1 2 7 8 0.15 2.0 0.5 10.0 9.8
pair_style lj/cut/tip4p/long/soft 1 2 7 8 0.15 2.0 0.5 10.0 9.8 9.5
pair_coeff * * 0.155 3.1536 1.0
pair_coeff 1 1 0.155 3.1536 1.0 9.5

pair_style lj/charmm/coul/long 2.0 0.5 10.0 8.0 10.0
pair_style lj/charmm/coul/long 2.0 0.5 10.0 8.0 10.0 9.0
pair_coeff * * 0.28 3.1 1.0
pair_coeff 1 1 0.28 3.1 1.0 0.14 3.1

pair_style lj/class2/coul/long/soft 2.0 0.5 10.0 9.5
pair_style lj/class2/coul/long/soft 2.0 0.5 10.0 9.5 9.5

```

(continues on next page)

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```

pair_coeff * * 0.28 3.1 1.0
pair_coeff 1 1 0.28 3.1 0.0 10.0
pair_coeff 1 1 0.28 3.1 0.0 10.0 9.5

pair_style coul/long/soft 1.0 10.0 9.5
pair_coeff * * 1.0
pair_coeff 1 1 1.0

pair_style tip4p/long/soft 1 2 7 8 0.15 2.0 0.5 10.0 9.8
pair_coeff * * 1.0
pair_coeff 1 1 1.0

pair_style morse/soft 4 0.9 10.0
pair_coeff * * 100.0 2.0 1.5 1.0
pair_coeff 1 1 100.0 2.0 1.5 1.0 3.0

```

Example input scripts available: `examples/PACKAGES/fep`

### 19.111.3 Description

These pair styles have a soft repulsive core, tunable by a parameter `lambda`, in order to avoid singularities during free energy calculations when sites are created or annihilated (*Beutler*). When `lambda` tends to 0 the pair interaction vanishes with a soft repulsive core. When `lambda` tends to 1, the pair interaction approaches the normal, non-soft potential. These pair styles are suited for “alchemical” free energy calculations using the *fix adapt/fep* and *compute fep* commands.

The *lj/cut/soft* style and related sub-styles compute the 12-6 Lennard-Jones and Coulomb potentials modified by a soft core, with the functional form

$$E = \lambda^n 4\epsilon \left\{ \frac{1}{\left[ \alpha_{LJ}(1-\lambda)^2 + \left(\frac{r}{\sigma}\right)^6 \right]^2} - \frac{1}{\alpha_{LJ}(1-\lambda)^2 + \left(\frac{r}{\sigma}\right)^6} \right\} \quad r < r_c$$

The *lj/class2/soft* style is a 9-6 potential with the exponent of the denominator of the first term in brackets taking the value 1.5 instead of 2 (other details differ, see the form of the potential in *pair\_style lj/class2*).

Coulomb interactions can also be damped with a soft core at short distance,

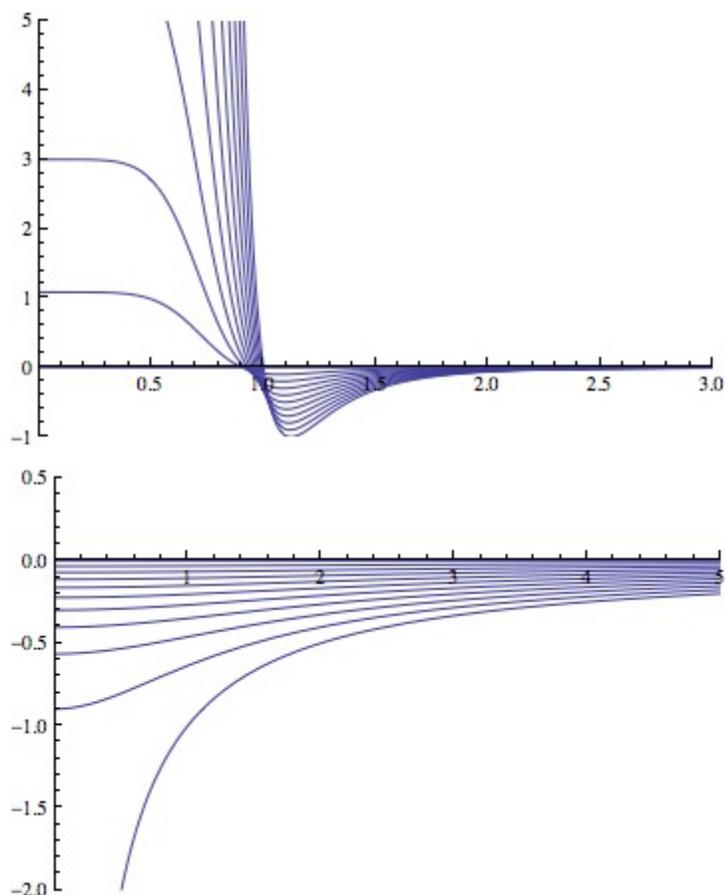
$$E = \lambda^n \frac{Cq_iq_j}{\epsilon [\alpha_C(1-\lambda)^2 + r^2]^{1/2}} \quad r < r_c$$

In the Coulomb part  $C$  is an energy-conversion constant,  $q_i$  and  $q_j$  are the charges on the 2 atoms, and epsilon is the dielectric constant which can be set by the *dielectric* command.

The coefficient `lambda` is an activation parameter. When  $\lambda = 1$  the pair potential is identical to a Lennard-Jones term or a Coulomb term or a combination of both. When  $\lambda = 0$  the interactions are deactivated. The transition between these two extrema is smoothed by a soft repulsive core in order to avoid singularities in potential energy and forces when sites are created or annihilated and can overlap (*Beutler*).

The parameters  $n$ ,  $\alpha_{LJ}$  and  $\alpha_C$  are set in the *pair\_style* command, before the cutoffs. Usual choices for the exponent are  $n = 2$  or  $n = 1$ . For the remaining coefficients  $\alpha_{LJ} = 0.5$  and  $\alpha_C = 10 \text{ \AA}^2$  are appropriate choices. Plots of the 12-6 LJ and Coulomb terms are shown below, for `lambda` ranging from 1 to 0 every 0.1.





For the *lj/cut/coul/cut/soft* or *lj/cut/coul/long/soft* pair styles, as well as for the equivalent *class2* versions, the following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- $\lambda$  (activation parameter, between 0 and 1)
- cutoff1 (distance units)
- cutoff2 (distance units)

The latter two coefficients are optional. If not specified, the global LJ and Coulombic cutoffs specified in the *pair\_style* command are used. If only one cutoff is specified, it is used as the cutoff for both LJ and Coulombic interactions for this type pair. If both coefficients are specified, they are used as the LJ and Coulombic cutoffs for this type pair. You cannot specify 2 cutoffs for style *lj/cut/soft*, since it has no Coulombic terms. For the *coul/cut/soft* and *coul/long/soft* only lambda and the optional cutoff2 are to be specified.

Style *lj/cut/tip4p/long/soft* implements a soft-core version of the TIP4P water model. The usage of the TIP4P pair style is documented in the *pair\_lj* styles. In the soft version the parameters  $n$ ,  $\alpha_{LJ}$  and  $\alpha_C$  are set in the *pair\_style* command, after the specific parameters of the TIP4P water model and before the cutoffs. The activation parameter lambda is supplied as an argument of the *pair\_coeff* command, after epsilon and sigma and before the optional cutoffs.

Style *lj/charmm/coul/long/soft* implements a soft-core version of the modified 12-6 LJ potential used in CHARMM and documented in the *pair\_style lj/charmm/coul/long* style. In the soft version the parameters  $n$ ,  $\alpha_{LJ}$  and  $\alpha_C$  are set in the *pair\_style* command, before the global cutoffs. The activation parameter lambda is introduced as an argument of the *pair\_coeff* command, after  $\epsilon$  and  $\sigma$  and before the optional eps14 and sigma14.

Style *lj/class2/soft* implements a soft-core version of the 9-6 potential in *pair\_style lj/class2*. In the soft version the parameters  $n$ ,  $\alpha_{LJ}$  and  $\alpha_C$  are set in the *pair\_style* command, before the global cutoffs. The activation parameter  $\lambda$  is introduced as an argument of the *pair\_coeff* command, after  $\epsilon$  and  $\sigma$  and before the optional cutoffs.

The *coul/cut/soft*, *coul/long/soft* and *tip4p/long/soft* sub-styles are designed to be combined with other pair potentials via the *pair\_style hybrid/overlay* command. This is because they have no repulsive core. Hence, if used by themselves, there will be no repulsion to keep two oppositely charged particles from overlapping each other. In this case, if  $\lambda = 1$ , a singularity may occur. These sub-styles are suitable to represent charges embedded in the Lennard-Jones radius of another site (for example hydrogen atoms in several water models). The  $\lambda$  must be defined for each pair, and *coul/cut/soft* can accept an optional cutoff as the second coefficient.

---

**Note:** When using the soft-core Coulomb potentials with long-range solvers (*coul/long/soft*, *lj/cut/coul/long/soft*, etc.) in a free energy calculation in which sites holding electrostatic charges are being created or annihilated (using *fix adapt/fep* and *compute fep*) it is important to adapt both the  $\lambda$  activation parameter (from 0 to 1, or the reverse) and the value of the charge (from 0 to its final value, or the reverse). This ensures that long-range electrostatic terms (k-space) are correct. It is not necessary to use soft-core Coulomb potentials if the van der Waals site is present during the free-energy route, thus avoiding overlap of the charges. Examples are provided in the LAMMPS source directory tree, under examples/PACKAGES/fep.

---

---

**Note:** To avoid division by zero do not set  $\sigma = 0$  in the *lj/cut/soft* and related styles; use the  $\lambda$  parameter instead to activate/deactivate interactions, or use  $\epsilon = 0$  and  $\sigma = 1$ . Alternatively, when sites do not interact though the Lennard-Jones term the *coul/long/soft* or similar sub-style can be used via the *pair\_style hybrid/overlay* command.

---

The *morse/soft* variant modifies the *pair\_morse* style at short range to have a soft core. The functional form differs from that of the *lj/soft* styles, and is instead given by:

$$\begin{aligned} s(\lambda) &= (1 - \lambda)/(1 - \lambda_f), & B &= -2De^{-2\alpha r_0}(e^{\alpha r_0} - 1)/3 \\ E &= D_0 \left[ e^{-2\alpha(r-r_0)} - 2e^{-\alpha(r-r_0)} \right] + s(\lambda)Be^{-3\alpha(r-r_0)}, & \lambda &\geq \lambda_f, \quad r < r_c \\ E &= \left( D_0 \left[ e^{-2\alpha(r-r_0)} - 2e^{-\alpha(r-r_0)} \right] + Be^{-3\alpha(r-r_0)} \right) (\lambda/\lambda_f)^n, & \lambda &< \lambda_f, \quad r < r_c \end{aligned}$$

The *morse/soft* style requires the following pair coefficients:

- $D_0$  (energy units)
- $\alpha$  (1/distance units)
- $r_0$  (distance units)
- $\lambda$  (unitless, between 0.0 and 1.0)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global morse cutoff is used.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.111.4 Mixing, shift, table, tail correction, restart, rRESPA info

The different versions of the *lj/cut/soft* pair styles support mixing. For atom type pairs I,J and I != J, the  $\epsilon$  and  $\sigma$  coefficients and cutoff distance for these pair styles can be mixed. The default mix value is *geometric* for 12-6 styles.

The mixing rule for epsilon and sigma for *lj/class2/soft* 9-6 potentials is to use the *sixthpower* formulas. The *pair\_modify mix* setting is thus ignored for class2 potentials for  $\epsilon$  and  $\sigma$ . However it is still followed for mixing the cutoff distance. See the *pair\_modify* command for details.

The *morse/soft* pair style does not support mixing. Thus, coefficients for all LJ pairs must be specified explicitly.

All of the pair styles with soft core support the *pair\_modify* shift option for the energy of the Lennard-Jones portion of the pair interaction.

The different versions of the *lj/cut/soft* pair styles support the *pair\_modify* tail option for adding a long-range tail correction to the energy and pressure for the Lennard-Jones portion of the pair interaction.

---

**Note:** The analytical form of the tail corrections for energy and pressure used in the *lj/cut/soft* potentials are approximate, being identical to that of the corresponding non-soft potentials scaled by a factor  $\lambda^n$ . The errors due to this approximation should be negligible. For example, for a cutoff of  $2.5\sigma$  this approximation leads to maximum relative errors in tail corrections of the order of  $1e-4$  for energy and virial ( $\alpha_{LJ} = 0.5, n = 2$ ). The error vanishes when lambda approaches 0 or 1. Note that these are the errors affecting the long-range tail (itself a correction to the interaction energy) which includes other approximations, namely that the system is homogeneous (local density equal the average density) beyond the cutoff.

---

The *morse/soft* pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

All of these pair styles write information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

---

#### 19.111.5 Restrictions

The pair styles with soft core are only enabled if LAMMPS was built with the FEP package. The *long* versions also require the KSPACE package to be installed. The soft *tip4p* versions also require the MOLECULE package to be installed. These styles are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

### 19.111.6 Related commands

*pair\_coeff*, *fix adapt*, *fix adapt/fep*, *compute fep*

### 19.111.7 Default

none

---

(Beutler) Beutler, Mark, van Schaik, Gerber, van Gunsteren, Chem Phys Lett, 222, 529 (1994).

## 19.112 pair\_style gauss command

Accelerator Variants: *gauss/gpu*, *gauss/omp*

## 19.113 pair\_style gauss/cut command

Accelerator Variants: *gauss/cut/omp*

### 19.113.1 Syntax

```
pair_style gauss cutoff
pair_style gauss/cut cutoff
```

- cutoff = global cutoff for Gauss interactions (distance units)

### 19.113.2 Examples

```
pair_style gauss 12.0
pair_coeff * * 1.0 0.9
pair_coeff 1 4 1.0 0.9 10.0

pair_style gauss/cut 3.5
pair_coeff 1 4 0.2805 1.45 0.112
```

### 19.113.3 Description

Style *gauss* computes a tethering potential of the form

$$E = -A \exp(-Br^2) \quad r < r_c$$

between an atom and its corresponding tether site which will typically be a frozen atom in the simulation.  $r_c$  is the cutoff.

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- A (energy units)
- B (1/distance<sup>2</sup> units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global cutoff is used.

Style *gauss/cut* computes a generalized Gaussian interaction potential between pairs of particles:

$$E = \frac{H}{\sigma_h \sqrt{2\pi}} \exp \left[ -\frac{(r - r_{mh})^2}{2\sigma_h^2} \right]$$

where H determines together with the standard deviation  $\sigma_h$  the peak height of the Gaussian function, and  $r_{mh}$  the peak position. Examples of the use of the Gaussian potentials include implicit solvent simulations of salt ions (*Lenart*) and of surfactants (*Jusufo*). In these instances the Gaussian potential mimics the hydration barrier between a pair of particles. The hydration barrier is located at  $r_{mh}$  and has a width of  $\sigma_h$ . The prefactor determines the height of the potential barrier.

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- H (energy \* distance units)
- $r_{mh}$  (distance units)
- $\sigma_h$  (distance units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global cutoff is used.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 19.113.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the A, B, H, sigma\_h, r\_mh parameters, and the cutoff distance for these pair styles can be mixed: A (energy units) sqrt(1/B) (distance units, see below) H (energy units) sigma\_h (distance units) r\_mh (distance units) cutoff (distance units):ul

The default mix value is *geometric*. Only *arithmetic* and *geometric* mix values are supported. See the “pair\_modify” command for details.

The A and H parameters are mixed using the same rules normally used to mix the “epsilon” parameter in a Lennard Jones interaction. The sigma\_h, r\_mh, and the cutoff distance are mixed using the same rules used to mix the “sigma” parameter in a Lennard Jones interaction. The B parameter is converted to a distance (sigma), before mixing (using sigma=B<sup>-0.5</sup>), and converted back to a coefficient afterwards (using B=sigma<sup>2</sup>). Negative A values are

converted to positive  $A$  values (using  $\text{abs}(A)$ ) before mixing, and converted back after mixing (by multiplying by  $\text{min}(\text{sign}(A_i), \text{sign}(A_j))$ ). This way, if either particle is repulsive (if  $A_i < 0$  or  $A_j < 0$ ), then the default interaction between both particles will be repulsive.

The *gauss* style does not support the *pair\_modify* shift option. There is no effect due to the Gaussian well beyond the cutoff; hence reasonable cutoffs need to be specified.

The *gauss/cut* style supports the *pair\_modify* shift option for the energy of the Gauss-potential portion of the pair interaction.

The *pair\_modify* table and tail options are not relevant for these pair styles.

These pair styles write their information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

The *gauss* pair style tallies an “occupancy” count of how many Gaussian-well sites have an atom within the distance at which the force is a maximum =  $\sqrt{0.5/b}$ . This quantity can be accessed via the *compute pair* command as a vector of values of length 1.

To print this quantity to the log file (with a descriptive column heading) the following commands could be included in an input script:

```
compute gauss all pair gauss
variable occ equal c_gauss[1]
thermo_style custom step temp epair v_occ
```

---

### 19.113.5 Restrictions

The *gauss* and *gauss/cut* styles are part of the EXTRA-PAIR package. They are only enabled if LAMMPS is build with that package. See the *Build package* page for more info.

The *gauss* style does not apply *special\_bonds* factors. When using this pair style on a system that has bonds, the *special\_bonds* factors, if using the default setting of 0.0, may need to be adjusted to some very small number (e.g.  $1.0e-100$ ), so that those special pairs are not completely excluded from the neighbor lists, but won't contribute forces or energies from styles (e.g. when used in combination with a *hybrid pair style*) that do apply those factors.

### 19.113.6 Related commands

*pair\_coeff*, *pair\_style coul/diel*

### 19.113.7 Default

none

**(Lenart)** Lenart, Jusufi, and Panagiotopoulos, J Chem Phys, 126, 044509 (2007).

**(Jusufi)** Jusufi, Hynninen, and Panagiotopoulos, J Phys Chem B, 112, 13783 (2008).

## 19.114 pair\_style gayberne command

Accelerator Variants: *gayberne/gpu*, *gayberne/intel*, *gayberne/omp*

### 19.114.1 Syntax

```
pair_style gayberne gamma epsilon mu cutoff
```

- gamma = shift for potential minimum (typically 1)
- epsilon = exponent for eta orientation-dependent energy function
- mu = exponent for chi orientation-dependent energy function
- cutoff = global cutoff for interactions (distance units)

### 19.114.2 Examples

```
pair_style gayberne 1.0 1.0 1.0 10.0
pair_coeff * * 1.0 1.7 1.7 3.4 3.4 1.0 1.0 1.0
```

### 19.114.3 Description

The *gayberne* styles compute a Gay-Berne anisotropic LJ interaction (*Berardi*) between pairs of ellipsoidal particles or an ellipsoidal and spherical particle via the formulas

$$U(\mathbf{A}_1, \mathbf{A}_2, \mathbf{r}_{12}) = U_r(\mathbf{A}_1, \mathbf{A}_2, \mathbf{r}_{12}, \gamma) \cdot \eta_{12}(\mathbf{A}_1, \mathbf{A}_2, v) \cdot \chi_{12}(\mathbf{A}_1, \mathbf{A}_2, \mathbf{r}_{12}, \mu)$$

$$U_r = 4\epsilon(\varrho^{12} - \varrho^6)$$

$$\varrho = \frac{\sigma}{h_{12} + \gamma\sigma}$$

where  $\mathbf{A}_1$  and  $\mathbf{A}_2$  are the transformation matrices from the simulation box frame to the body frame and  $\mathbf{r}_{12}$  is the center to center vector between the particles.  $U_r$  controls the shifted distance dependent interaction based on the distance of closest approach of the two particles ( $h_{12}$ ) and the user-specified shift parameter gamma. When both particles are spherical, the formula reduces to the usual Lennard-Jones interaction (see details below for when Gay-Berne treats a particle as “spherical”).

For large uniform molecules it has been shown that the energy parameters are approximately representable in terms of local contact curvatures (*Everaers*):

$$\epsilon_a = \sigma \cdot \frac{a}{b \cdot c}; \epsilon_b = \sigma \cdot \frac{b}{a \cdot c}; \epsilon_c = \sigma \cdot \frac{c}{a \cdot b}$$

The variable names utilized as potential parameters are for the most part taken from (*Everaers*) in order to be consistent with the *RE-squared pair potential*. Details on the epsilon and mu parameters are given [here](#).

More details of the Gay-Berne formulation are given in the references listed below and in [this supplementary document](#).

Use of this pair style requires the NVE, NVT, or NPT fixes with the *asphere* extension (e.g. *fix nve/asphere*) in order to integrate particle rotation. Additionally, *atom\_style ellipsoid* should be used since it defines the rotational state and the size and shape of each ellipsoidal particle.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  = well depth (energy units)
- $\sigma$  = minimum effective particle radii (distance units)
- $\epsilon_{i,a}$  = relative well depth of type I for side-to-side interactions
- $\epsilon_{i,b}$  = relative well depth of type I for face-to-face interactions
- $\epsilon_{i,c}$  = relative well depth of type I for end-to-end interactions
- $\epsilon_{j,a}$  = relative well depth of type J for side-to-side interactions
- $\epsilon_{j,b}$  = relative well depth of type J for face-to-face interactions
- $\epsilon_{j,c}$  = relative well depth of type J for end-to-end interactions
- cutoff (distance units)

The last coefficient is optional. If not specified, the global cutoff specified in the `pair_style` command is used.

It is typical with the Gay-Berne potential to define  $\sigma$  as the minimum of the 3 shape diameters of the particles involved in an I,I interaction, though this is not required. Note that this is a different meaning for  $\sigma$  than the *pair\_style resquared* potential uses.

The  $\epsilon_i$  and  $\epsilon_j$  coefficients are actually defined for atom types, not for pairs of atom types. Thus, in a series of `pair_coeff` commands, they only need to be specified once for each atom type.

Specifically, if any of  $\epsilon_{i,a}$ ,  $\epsilon_{i,b}$ ,  $\epsilon_{i,c}$  are non-zero, the three values are assigned to atom type I. If all the  $\epsilon_i$  values are zero, they are ignored. If any of  $\epsilon_{j,a}$ ,  $\epsilon_{j,b}$ ,  $\epsilon_{j,c}$  are non-zero, the three values are assigned to atom type J. If all three `epsilon_j` values are zero, they are ignored. Thus the typical way to define the  $\epsilon_i$  and  $\epsilon_j$  coefficients is to list their values in “`pair_coeff I J`” commands when  $I = J$ , but set them to 0.0 when  $I \neq J$ . If you do list them when  $I \neq J$ , you should insure they are consistent with their values in other `pair_coeff` commands, since only the last setting will be in effect.

Note that if this potential is being used as a sub-style of *pair\_style hybrid*, and there is no “`pair_coeff I I`” setting made for Gay-Berne for a particular type I (because I-I interactions are computed by another hybrid pair potential), then you still need to insure the  $\epsilon_{a,b,c}$  coefficients are assigned to that type. e.g. in a “`pair_coeff I J`” command.

---

**Note:** If the  $\epsilon_{a,b,c}$  for an atom type, and if the shape of the particle itself is spherical, meaning its 3 shape parameters are all the same, then the particle is treated as an LJ sphere by the Gay-Berne potential. This is significant because if two LJ spheres interact, then the simple Lennard-Jones formula is used to compute their interaction energy/force using the specified epsilon and sigma as the standard LJ parameters. This is much cheaper to compute than the full Gay-Berne formula. To treat the particle as a LJ sphere with  $\sigma = D$ , you should normally set  $\epsilon_{a,b,c} = 1.0$ , set the `pair_coeff  $\sigma = D$` , and also set the 3 shape parameters for the particle to  $D$ . The one exception is that if the 3 shape parameters are set to 0.0, which is a valid way in LAMMPS to specify a point particle, then the Gay-Berne potential will treat that as shape parameters of 1.0 (i.e. a LJ particle with  $\sigma = 1$ ), since it requires finite-size particles. In this case you should still set the `pair_coeff  $\sigma = 1.0$`  as well.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.



#### 19.114.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , the epsilon and sigma coefficients and cutoff distance for this pair style can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

This pair style supports the *pair\_modify* shift option for the energy of the Lennard-Jones portion of the pair interaction, but only for sphere-sphere interactions. There is no shifting performed for ellipsoidal interactions due to the anisotropic dependence of the interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.114.5 Restrictions

The *gayberne* style is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These pair styles require that atoms store torque and a quaternion to represent their orientation, as defined by the *atom\_style*. It also require they store a per-type *shape*. The particles cannot store a per-particle diameter.

This pair style requires that atoms be ellipsoids as defined by the *atom\_style ellipsoid* command.

Particles acted on by the potential can be finite-size aspherical or spherical particles, or point particles. Spherical particles have all 3 of their shape parameters equal to each other. Point particles have all 3 of their shape parameters equal to 0.0.

The Gay-Berne potential does not become isotropic as  $r$  increases (*Everaers*). The distance-of-closest-approach approximation used by LAMMPS becomes less accurate when high-aspect ratio ellipsoids are used.

#### 19.114.6 Related commands

*pair\_coeff*, *fix nve/asphere*, *compute temp/asphere*, *pair\_style resquared*

#### 19.114.7 Default

none

---

**(Everaers)** Everaers and Ejtehadi, Phys Rev E, 67, 041710 (2003).

**(Berardi)** Berardi, Fava, Zannoni, Chem Phys Lett, 297, 8-14 (1998). Berardi, Muccioli, Zannoni, J Chem Phys, 128, 024905 (2008).

**(Perram)** Perram and Rasmussen, Phys Rev E, 54, 6565-6572 (1996).

**(Allen)** Allen and Germano, Mol Phys 104, 3225-3235 (2006).

## 19.115 pair\_style gran/hooke command

Accelerator Variants: *gran/hooke/omp*

## 19.116 pair\_style gran/hooke/history command

Accelerator Variants: *gran/hooke/history/omp*, *gran/hooke/history/kk*

## 19.117 pair\_style gran/hertz/history command

Accelerator Variants: *gran/hertz/history/omp*

### 19.117.1 Syntax

```
pair_style style Kn Kt gamma_n gamma_t xmu dampflag keyword
```

- style = *gran/hooke* or *gran/hooke/history* or *gran/hertz/history*
- Kn = elastic constant for normal particle repulsion (force/distance units or pressure units - see discussion below)
- Kt = elastic constant for tangential contact (force/distance units or pressure units - see discussion below)
- gamma\_n = damping coefficient for collisions in normal direction (1/time units or 1/time-distance units - see discussion below)
- gamma\_t = damping coefficient for collisions in tangential direction (1/time units or 1/time-distance units - see discussion below)
- xmu = static yield criterion (unitless value between 0.0 and 1.0e4)
- dampflag = 0 or 1 if tangential damping force is excluded or included
- keyword = *limit\_damping*
  - limit\_damping* value = none
  - limit damping to prevent attractive interaction

---

**Note:** Versions of LAMMPS before 9Jan09 had different style names for granular force fields. This is to emphasize the fact that the Hertzian equation has changed to model polydispersity more accurately. A side effect of the change is that the Kn, Kt, gamma\_n, and gamma\_t coefficients in the pair\_style command must be specified with different values in order to reproduce calculations made with earlier versions of LAMMPS, even for monodisperse systems. See the NOTE below for details.

---

## 19.117.2 Examples

```
pair_style gran/hooke/history 200000.0 NULL 50.0 NULL 0.5 1
pair_style gran/hooke 200000.0 70000.0 50.0 30.0 0.5 0
pair_style gran/hooke 200000.0 70000.0 50.0 30.0 0.5 0 limit_damping
```

## 19.117.3 Description

The *gran* styles use the following formulas for the frictional force between two granular particles, as described in (*Brilliantov*), (*Silbert*), and (*Zhang*), when the distance  $r$  between two particles of radii  $R_i$  and  $R_j$  is less than their contact distance  $d = R_i + R_j$ . There is no force between the particles when  $r > d$ .

The two Hookean styles use this formula:

$$F_{hk} = (k_n \delta \mathbf{n}_{ij} - m_{eff} \gamma_n \mathbf{v}_n) - (k_t \Delta \mathbf{s}_t + m_{eff} \gamma_t \mathbf{v}_t)$$

The Hertzian style uses this formula:

$$F_{hz} = \sqrt{\delta} \sqrt{\frac{R_i R_j}{R_i + R_j}} F_{hk} = \sqrt{\delta} \sqrt{\frac{R_i R_j}{R_i + R_j}} \left[ (k_n \delta \mathbf{n}_{ij} - m_{eff} \gamma_n \mathbf{v}_n) - (k_t \Delta \mathbf{s}_t + m_{eff} \gamma_t \mathbf{v}_t) \right]$$

In both equations the first parenthesized term is the normal force between the two particles and the second parenthesized term is the tangential force. The normal force has 2 terms, a contact force and a damping force. The tangential force also has 2 terms: a shear force and a damping force. The shear force is a “history” effect that accounts for the tangential displacement between the particles for the duration of the time they are in contact. This term is included in pair styles *hooke/history* and *hertz/history*, but is not included in pair style *hooke*. The tangential damping force term is included in all three pair styles if *dampflag* is set to 1; it is not included if *dampflag* is set to 0.

The other quantities in the equations are as follows:

- $\delta = d - r =$  overlap distance of 2 particles
- $K_n =$  elastic constant for normal contact
- $K_t =$  elastic constant for tangential contact
- $\gamma_n =$  viscoelastic damping constant for normal contact
- $\gamma_t =$  viscoelastic damping constant for tangential contact
- $m_{eff} = M_i M_j / (M_i + M_j) =$  effective mass of 2 particles of mass  $M_i$  and  $M_j$
- $\Delta \mathbf{s}_t =$  tangential displacement vector between 2 particles which is truncated to satisfy a frictional yield criterion
- $\mathbf{n}_{ij} =$  unit vector along the line connecting the centers of the 2 particles
- $V_n =$  normal component of the relative velocity of the 2 particles
- $V_t =$  tangential component of the relative velocity of the 2 particles

The  $K_n$ ,  $K_t$ ,  $\gamma_n$ , and  $\gamma_t$  coefficients are specified as parameters to the *pair\_style* command. If a NULL is used for  $K_t$ , then a default value is used where  $K_t = 2/7 K_n$ . If a NULL is used for  $\gamma_t$ , then a default value is used where  $\gamma_t = 1/2 \gamma_n$ .

The interpretation and units for these 4 coefficients are different in the Hookean versus Hertzian equations.

The Hookean model is one where the normal push-back force for two overlapping particles is a linear function of the overlap distance. Thus the specified  $K_n$  is in units of (force/distance). Note that this push-back force is independent of absolute particle size (in the monodisperse case) and of the relative sizes of the two particles (in the polydisperse

case). This model also applies to the other terms in the force equation so that the specified  $\gamma_n$  is in units of (1/time),  $K_t$  is in units of (force/distance), and  $\gamma_t$  is in units of (1/time).

The Hertzian model is one where the normal push-back force for two overlapping particles is proportional to the area of overlap of the two particles, and is thus a non-linear function of overlap distance. Thus  $K_n$  has units of force per area and is thus specified in units of (pressure). The effects of absolute particle size (monodispersity) and relative size (polydispersity) are captured in the radii-dependent prefactors. When these prefactors are carried through to the other terms in the force equation it means that the specified  $\gamma_n$  is in units of (1/(time\*distance)),  $K_t$  is in units of (pressure), and  $\gamma_t$  is in units of (1/(time\*distance)).

Note that in the Hookean case,  $K_n$  can be thought of as a linear spring constant with units of force/distance. In the Hertzian case,  $K_n$  is like a non-linear spring constant with units of force/area or pressure, and as shown in the (Zhang) paper,  $K_n = 4G/(3(1-\nu))$  where  $\nu$  = the Poisson ratio,  $G$  = shear modulus =  $E/(2(1+\nu))$ , and  $E$  = Young's modulus. Similarly,  $K_t = 4G/(2-\nu)$ . (NOTE: in an earlier version of the manual, we incorrectly stated that  $K_t = 8G/(2-\nu)$ .)

Thus in the Hertzian case  $K_n$  and  $K_t$  can be set to values that corresponds to properties of the material being modeled. This is also true in the Hookean case, except that a spring constant must be chosen that is appropriate for the absolute size of particles in the model. Since relative particle sizes are not accounted for, the Hookean styles may not be a suitable model for polydisperse systems.

---

**Note:** In versions of LAMMPS before 9Jan09, the equation for Hertzian interactions did not include the  $\sqrt{r_i r_j / (r_i + r_j)}$  term and thus was not as accurate for polydisperse systems. For monodisperse systems,  $\sqrt{r_i r_j / (r_i + r_j)}$  is a constant factor that effectively scales all 4 coefficients:  $K_n, K_t, \gamma_n, \gamma_t$ . Thus you can set the values of these 4 coefficients appropriately in the current code to reproduce the results of a previous Hertzian monodisperse calculation. For example, for the common case of a monodisperse system with particles of diameter 1, all 4 of these coefficients should now be set 2x larger than they were previously.

---

$xmu$  is also specified in the `pair_style` command and is the upper limit of the tangential force through the Coulomb criterion  $F_t = xmu * F_n$ , where  $F_t$  and  $F_n$  are the total tangential and normal force components in the formulas above. Thus in the Hookean case, the tangential force between 2 particles grows according to a tangential spring and dash-pot model until  $F_t/F_n = xmu$  and is then held at  $F_t = F_n * xmu$  until the particles lose contact. In the Hertzian case, a similar analogy holds, though the spring is no longer linear.

---

**Note:** Normally,  $xmu$  should be specified as a fractional value between 0.0 and 1.0, however LAMMPS allows large values (up to 1.0e4) to allow for modeling of systems which can sustain very large tangential forces.

---

The effective mass  $m_{eff}$  is given by the formula above for two isolated particles. If either particle is part of a rigid body, its mass is replaced by the mass of the rigid body in the formula above. This is determined by searching for a `fix rigid` command (or its variants).

For granular styles there are no additional coefficients to set for each pair of atom types via the `pair_coeff` command. All settings are global and are made via the `pair_style` command. However you must still use the `pair_coeff` for all pairs of granular atom types. For example the command

```
pair_coeff * *
```

should be used if all atoms in the simulation interact via a granular potential (i.e. one of the pair styles above is used). If a granular potential is used as a sub-style of `pair_style hybrid`, then specific atom types can be used in the `pair_coeff` command to determine which atoms interact via a granular potential.

If two particles are moving away from each other while in contact, there is a possibility that the particles could experience an effective attractive force due to damping. If the `limit_damping` keyword is used, this option will zero out the normal component of the force if there is an effective attractive force.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* [command-line switch](#) when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.117.4 Mixing, shift, table, tail correction, restart, rRESPA info

The *pair\_modify* mix, shift, table, and tail options are not relevant for granular pair styles.

These pair styles write their information to *binary restart files*, so a *pair\_style* command does not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

The *single()* function of these pair styles returns 0.0 for the energy of a pairwise interaction, since energy is not conserved in these dissipative potentials. It also returns only the normal component of the pairwise interaction force. However, the *single()* function also calculates 10 extra pairwise quantities. The first 3 are the components of the tangential force between particles I and J, acting on particle I. The fourth is the magnitude of this tangential force. The next 3 (5-7) are the components of the relative velocity in the normal direction (along the line joining the 2 sphere centers). The last 3 (8-10) the components of the relative velocity in the tangential direction.

These extra quantities can be accessed by the *compute pair/local* command, as *p1*, *p2*, ..., *p10*.

---

#### 19.117.5 Restrictions

All the granular pair styles are part of the GRANULAR package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

These pair styles require that atoms store torque and angular velocity (*omega*) as defined by the *atom\_style*. They also require a per-particle radius is stored. The *sphere* atom style does all of this.

This pair style requires you to use the *comm\_modify vel yes* command so that velocities are stored by ghost atoms.

These pair styles will not restart exactly when using the *read\_restart* command, though they should provide statistically similar results. This is because the forces they compute depend on atom velocities. See the *read\_restart* command for more details.

## 19.117.6 Related commands

*pair\_coeff*

## 19.117.7 Default

none

---

**(Brilliantov)** Brilliantov, Spahn, Hertzsch, Poschel, Phys Rev E, 53, p 5382-5392 (1996).

**(Silbert)** Silbert, Ertas, Grest, Halsey, Levine, Plimpton, Phys Rev E, 64, p 051302 (2001).

**(Zhang)** Zhang and Makse, Phys Rev E, 72, p 011301 (2005).

## 19.118 pair\_style granular command

### 19.118.1 Syntax

```
pair_style granular cutoff
```

- cutoff = global cutoff (optional). See discussion below.

### 19.118.2 Examples

```
pair_style granular
pair_coeff * * hooke 1000.0 50.0 tangential linear_nohistory 1.0 0.4 damping mass_
↪velocity

pair_style granular
pair_coeff * * hooke 1000.0 50.0 tangential linear_history 500.0 1.0 0.4 damping mass_
↪velocity

pair_style granular
pair_coeff * * hertz 1000.0 50.0 tangential mindlin 1000.0 1.0 0.4 limit_damping

pair_style granular
pair_coeff * * hertz/material 1e8 0.3 0.3 tangential mindlin_rescale NULL 1.0 0.4_
↪damping tsuji

pair_style granular
pair_coeff 1 * jkr 1000.0 500.0 0.3 10 tangential mindlin 800.0 1.0 0.5 rolling sds 500.
↪0 200.0 0.5 twisting marshall
pair_coeff 2 2 hertz 200.0 100.0 tangential linear_history 300.0 1.0 0.1 rolling sds 200.
↪0 100.0 0.1 twisting marshall

pair_style granular
pair_coeff 1 1 dmt 1000.0 50.0 0.3 0.0 tangential mindlin NULL 0.5 0.5 rolling sds 500.0_
↪200.0 0.5 twisting marshall
pair_coeff 2 2 dmt 1000.0 50.0 0.3 10.0 tangential mindlin NULL 0.5 0.1 rolling sds 500.
↪0 200.0 0.1 twisting marshall
```

### 19.118.3 Description

The *granular* styles support a variety of options for the normal, tangential, rolling and twisting forces resulting from contact between two granular particles. This expands on the options offered by the *pair gran/\** pair styles. The total computed forces and torques are the sum of various models selected for the normal, tangential, rolling and twisting modes of motion.

All model choices and parameters are entered in the *pair\_coeff* command, as described below. Unlike e.g. *pair gran/hooke*, coefficient values are not global, but can be set to different values for different combinations of particle types, as determined by the *pair\_coeff* command. If the contact model choice is the same for two particle types, the mixing for the cross-coefficients can be carried out automatically. This is shown in the last example, where model choices are the same for type 1 - type 1 as for type 2 - type2 interactions, but coefficients are different. In this case, the mixed coefficients for type 1 - type 2 interactions can be determined from mixing rules discussed below. For additional flexibility, coefficients as well as model forms can vary between particle types, as shown in the fourth example: type 1 - type 1 interactions are based on a Johnson-Kendall-Roberts normal contact model and 2-2 interactions are based on a DMT cohesive model (see below). In that example, 1-1 and 2-2 interactions have different model forms, in which case mixing of coefficients cannot be determined, so 1-2 interactions must be explicitly defined via the *pair\_coeff 1 \** command, otherwise an error would result.

The first required keyword for the *pair\_coeff* command is the normal contact model. Currently supported options for normal contact models and their required arguments are:

1. *hooke* :  $k_n, \eta_{n0}$  (or  $e$ )
2. *hertz* :  $k_n, \eta_{n0}$  (or  $e$ )
3. *hertz/material* :  $E, \eta_{n0}$  (or  $e$ ),  $\nu$
4. *dmt* :  $E, \eta_{n0}$  (or  $e$ ),  $\nu, \gamma$
5. *jkr* :  $E, \eta_{n0}$  (or  $e$ ),  $\nu, \gamma$

Here,  $k_n$  is spring stiffness (with units that depend on model choice, see below);  $\eta_{n0}$  is a damping prefactor (or, in its place a coefficient of restitution  $e$ , depending on the choice of damping mode, see below);  $E$  is Young's modulus in units of *forcellength*<sup>2</sup>, i.e. *pressure*;  $\nu$  is Poisson's ratio and  $\gamma$  is a surface energy density, in units of *energy/length*<sup>2</sup>.

For the *hooke* model, the normal, elastic component of force acting on particle  $i$  due to contact with particle  $j$  is given by:

$$\mathbf{F}_{ne,Hooke} = k_n \delta_{ij} \mathbf{n}$$

Where  $\delta_{ij} = R_i + R_j - \|\mathbf{r}_{ij}\|$  is the particle overlap,  $R_i, R_j$  are the particle radii,  $\mathbf{r}_{ij} = \mathbf{r}_i - \mathbf{r}_j$  is the vector separating the two particle centers (note the i-j ordering so that  $F_{ne}$  is positive for repulsion), and  $\mathbf{n} = \frac{\mathbf{r}_{ij}}{\|\mathbf{r}_{ij}\|}$ . Therefore, for *hooke*, the units of the spring constant  $k_n$  are *forcedistance*, or equivalently *mass/time*<sup>2</sup>.

For the *hertz* model, the normal component of force is given by:

$$\mathbf{F}_{ne,Hertz} = k_n R_{eff}^{1/2} \delta_{ij}^{3/2} \mathbf{n}$$

Here,  $R_{eff} = \frac{R_i R_j}{R_i + R_j}$  is the effective radius, denoted for simplicity as  $R$  from here on. For *hertz*, the units of the spring constant  $k_n$  are *forcellength*<sup>2</sup>, or equivalently *pressure*.

For the *hertz/material* model, the force is given by:

$$\mathbf{F}_{ne,Hertz/material} = \frac{4}{3} E_{eff} R_{eff}^{1/2} \delta_{ij}^{3/2} \mathbf{n}$$

Here,  $E_{eff} = E = \left( \frac{1-\nu_i^2}{E_i} + \frac{1-\nu_j^2}{E_j} \right)^{-1}$  is the effective Young's modulus, with  $\nu_i, \nu_j$  the Poisson ratios of the particles of types  $i$  and  $j$ . Note that if the elastic modulus and the shear modulus of the two particles are the same, the *hertz/material* model is equivalent to the *hertz* model with  $k_n = 4/3 E_{eff}$

The *dmt* model corresponds to the (*Derjaguin-Muller-Toporov*) cohesive model, where the force is simply Hertz with an additional attractive cohesion term:

$$\mathbf{F}_{ne,dmt} = \left( \frac{4}{3}ER^{1/2}\delta_{ij}^{3/2} - 4\pi\gamma R \right) \mathbf{n}$$

The *jkr* model is the (*Johnson-Kendall-Roberts*) model, where the force is computed as:

$$\mathbf{F}_{ne,jkr} = \left( \frac{4Ea^3}{3R} - 2\pi a^2 \sqrt{\frac{4\gamma E}{\pi a}} \right) \mathbf{n}$$

Here,  $a$  is the radius of the contact zone, related to the overlap  $\delta$  according to:

$$\delta = a^2/R - 2\sqrt{\pi\gamma a/E}$$

LAMMPS internally inverts the equation above to solve for  $a$  in terms of  $\delta$ , then solves for the force in the previous equation. Additionally, note that the JKR model allows for a tensile force beyond contact (i.e. for  $\delta < 0$ ), up to a maximum of  $3\pi\gamma R$  (also known as the ‘pull-off’ force). Note that this is a hysteretic effect, where particles that are not contacting initially will not experience force until they come into contact  $\delta \geq 0$ ; as they move apart and ( $\delta < 0$ ), they experience a tensile force up to  $3\pi\gamma R$ , at which point they lose contact.

---

In addition, the normal force is augmented by a damping term of the following general form:

$$\mathbf{F}_{n,damp} = -\eta_n \mathbf{v}_{n,rel}$$

Here,  $\mathbf{v}_{n,rel} = (\mathbf{v}_j - \mathbf{v}_i) \cdot \mathbf{n}$  is the component of relative velocity along  $\mathbf{n}$ .

The optional *damping* keyword to the *pair\_coeff* command followed by a keyword determines the model form of the damping factor  $\eta_n$ , and the interpretation of the  $\eta_{n0}$  or  $e$  coefficients specified as part of the normal contact model settings. The *damping* keyword and corresponding model form selection may be appended anywhere in the *pair\_coeff* command. Note that the choice of damping model affects both the normal and tangential damping (and depending on other settings, potentially also the twisting damping). The options for the damping model currently supported are:

1. *velocity*
2. *mass\_velocity*
3. *viscoelastic*
4. *tsuji*

If the *damping* keyword is not specified, the *viscoelastic* model is used by default.

For *damping velocity*, the normal damping is simply equal to the user-specified damping coefficient in the *normal* model:

$$\eta_n = \eta_{n0}$$

Here,  $\eta_{n0}$  is the damping coefficient specified for the normal contact model, in units of *mass/time*.

For *damping mass\_velocity*, the normal damping is given by:

$$\eta_n = \eta_{n0} m_{eff}$$

Here,  $\eta_{n0}$  is the damping coefficient specified for the normal contact model, in units of *1/time* and  $m_{eff} = m_i m_j / (m_i + m_j)$  is the effective mass. Use *damping mass\_velocity* to reproduce the damping behavior of *pair gran/hooke\**.

The *damping viscoelastic* model is based on the viscoelastic treatment of (*Brilliantov et al*), where the normal damping is given by:

$$\eta_n = \eta_{n0} a m_{eff}$$



Here,  $a$  is the contact radius, given by  $a = \sqrt{R\delta}$  for all models except *jkr*, for which it is given implicitly according to  $\delta = a^2/R - 2\sqrt{\pi\gamma a/E}$ . For *damping viscoelastic*,  $\eta_{n0}$  is in units of  $1/(time*distance)$ .

The *tsuji* model is based on the work of (*Tsuji et al*). Here, the damping coefficient specified as part of the normal model is interpreted as a restitution coefficient  $e$ . The damping constant  $\eta_n$  is given by:

$$\eta_n = \alpha(m_{eff}k_n)^{1/2}$$

For normal contact models based on material parameters,  $k_n = 4/3Ea$ . The parameter  $\alpha$  is related to the restitution coefficient  $e$  according to:

$$\alpha = 1.2728 - 4.2783e + 11.087e^2 - 22.348e^3 + 27.467e^4 - 18.022e^5 + 4.8218e^6$$

The dimensionless coefficient of restitution  $e$  specified as part of the normal contact model parameters should be between 0 and 1, but no error check is performed on this.

The total normal force is computed as the sum of the elastic and damping components:

$$\mathbf{F}_n = \mathbf{F}_{ne} + \mathbf{F}_{n,damp}$$

The *pair\_coeff* command also requires specification of the tangential contact model. The required keyword *tangential* is expected, followed by the model choice and associated parameters. Currently supported tangential model choices and their expected parameters are as follows:

1. *linear\_nohistory* :  $x_{\gamma,t}, \mu_s$
2. *linear\_history* :  $k_t, x_{\gamma,t}, \mu_s$
3. *mindlin* :  $k_t$  or NULL,  $x_{\gamma,t}, \mu_s$
4. *mindlin/force* :  $k_t$  or NULL,  $x_{\gamma,t}, \mu_s$
5. *mindlin\_rescale* :  $k_t$  or NULL,  $x_{\gamma,t}, \mu_s$
6. *mindlin\_rescale/force* :  $k_t$  or NULL,  $x_{\gamma,t}, \mu_s$

Here,  $x_{\gamma,t}$  is a dimensionless multiplier for the normal damping  $\eta_n$  that determines the magnitude of the tangential damping,  $\mu_t$  is the tangential (or sliding) friction coefficient, and  $k_t$  is the tangential stiffness coefficient.

For *tangential linear\_nohistory*, a simple velocity-dependent Coulomb friction criterion is used, which mimics the behavior of the *pair gran/hooke* style. The tangential force  $\mathbf{F}_t$  is given by:

$$\mathbf{F}_t = -\min(\mu_t F_{n0}, \|\mathbf{F}_{t,damp}\|)\mathbf{t}$$

The tangential damping force  $\mathbf{F}_{t,damp}$  is given by:

$$\mathbf{F}_{t,damp} = -\eta_t \mathbf{v}_{t,rel}$$

The tangential damping prefactor  $\eta_t$  is calculated by scaling the normal damping  $\eta_n$  (see above):

$$\eta_t = -x_{\gamma,t}\eta_n$$

The normal damping prefactor  $\eta_n$  is determined by the choice of the *damping* keyword, as discussed above. Thus, the *damping* keyword also affects the tangential damping. The parameter  $x_{\gamma,t}$  is a scaling coefficient. Several works in the literature use  $x_{\gamma,t} = 1$  (*Marshall, Tsuji et al, Silbert et al*). The relative tangential velocity at the point of contact is given by  $\mathbf{v}_{t,rel} = \mathbf{v}_t - (R_i\boldsymbol{\Omega}_i + R_j\boldsymbol{\Omega}_j) \times \mathbf{n}$ , where  $\mathbf{v}_t = \mathbf{v}_r - \mathbf{v}_r \cdot \mathbf{n} \mathbf{n}$ ,  $\mathbf{v}_r = \mathbf{v}_j - \mathbf{v}_i$ . The direction of the applied force is  $\mathbf{t} = \mathbf{v}_{t,rel}/\|\mathbf{v}_{t,rel}\|$ .

The normal force value  $F_{n0}$  used to compute the critical force depends on the form of the contact model. For non-cohesive models (*hertz*, *hertz/material*, *hooke*), it is given by the magnitude of the normal force:

$$F_{n0} = \|\mathbf{F}_n\|$$

For cohesive models such as *jkr* and *dmt*, the critical force is adjusted so that the critical tangential force approaches  $\mu_t F_{pulloff}$ , see *Marshall*, equation 43, and *Thornton*. For both models,  $F_{n0}$  takes the form:

$$F_{n0} = \|\mathbf{F}_{ne} + 2F_{pulloff}\|$$

Where  $F_{pulloff} = 3\pi\gamma R$  for *jkr*, and  $F_{pulloff} = 4\pi\gamma R$  for *dmt*.

The remaining tangential options all use accumulated tangential displacement (i.e. contact history), except for the options *mindlin/force* and *mindlin\_rescale/force*, that use accumulated tangential force instead, and are discussed further below. The accumulated tangential displacement is discussed in details below in the context of the *linear\_history* option. The same treatment of the accumulated displacement applies to the other options as well.

For *tangential\_linear\_history*, the tangential force is given by:

$$\mathbf{F}_t = -\min(\mu_t F_{n0}, \| -k_t \xi + \mathbf{F}_{t,damp} \|) \mathbf{t}$$

Here,  $\xi$  is the tangential displacement accumulated during the entire duration of the contact:

$$\xi = \int_{t_0}^t \mathbf{v}_{t,rel}(\tau) d\tau$$

This accumulated tangential displacement must be adjusted to account for changes in the frame of reference of the contacting pair of particles during contact. This occurs due to the overall motion of the contacting particles in a rigid-body-like fashion during the duration of the contact. There are two modes of motion that are relevant: the ‘tumbling’ rotation of the contacting pair, which changes the orientation of the plane in which tangential displacement occurs; and ‘spinning’ rotation of the contacting pair about the vector connecting their centers of mass ( $\mathbf{n}$ ). Corrections due to the former mode of motion are made by rotating the accumulated displacement into the plane that is tangential to the contact vector at each step, or equivalently removing any component of the tangential displacement that lies along  $\mathbf{n}$ , and rescaling to preserve the magnitude. This follows the discussion in *Luding*, see equation 17 and relevant discussion in that work:

$$\xi = (\xi' - (\mathbf{n} \cdot \xi') \mathbf{n}) \frac{\|\xi'\|}{\|\xi' - (\mathbf{n} \cdot \xi') \mathbf{n}\|}$$

Here,  $\xi'$  is the accumulated displacement prior to the current time step and  $\xi$  is the corrected displacement. Corrections to the displacement due to the second mode of motion described above (rotations about  $\mathbf{n}$ ) are not currently implemented, but are expected to be minor for most simulations.

Furthermore, when the tangential force exceeds the critical force, the tangential displacement is re-scaled to match the value for the critical force (see *Luding*, equation 20 and related discussion):

$$\xi = -\frac{1}{k_t} (\mu_t F_{n0} \mathbf{t} - \mathbf{F}_{t,damp})$$

The tangential force is added to the total normal force (elastic plus damping) to produce the total force on the particle. The tangential force also acts at the contact point (defined as the center of the overlap region) to induce a torque on each particle according to:

$$\tau_i = -(R_i - 0.5\delta) \mathbf{n} \times \mathbf{F}_t$$

$$\tau_j = -(R_j - 0.5\delta) \mathbf{n} \times \mathbf{F}_t$$

For *tangential\_mindlin*, the *Mindlin* no-slip solution is used which differs from the *linear\_history* option by an additional factor of  $a$ , the radius of the contact region. The tangential force is given by:

$$\mathbf{F}_t = -\min(\mu_t F_{n0}, \| -k_t a \xi + \mathbf{F}_{t,damp} \|) \mathbf{t}$$

Here,  $a$  is the radius of the contact region, given by  $a = \sqrt{R\delta}$  for all normal contact models, except for *jkr*, where it is given implicitly by  $\delta = a^2/R - 2\sqrt{\pi\gamma a/E}$ , see discussion above. To match the Mindlin solution, one should set  $k_t = 8G_{eff}$ , where  $G_{eff}$  is the effective shear modulus given by:

$$G_{eff} = \left( \frac{2 - \nu_i}{G_i} + \frac{2 - \nu_j}{G_j} \right)^{-1}$$

where  $G$  is the shear modulus, related to Young's modulus  $E$  and Poisson's ratio  $\nu$  by  $G = E/(2(1 + \nu))$ . This can also be achieved by specifying *NULL* for  $k_t$ , in which case a normal contact model that specifies material parameters  $E$  and  $\nu$  is required (e.g. *hertz/material*, *dmt* or *jkr*). In this case, mixing of the shear modulus for different particle types  $i$  and  $j$  is done according to the formula above.

---

**Note:** The radius of the contact region  $a$  depends on the normal overlap. As a result, the tangential force for *mindlin* can change due to a variation in normal overlap, even with no change in tangential displacement.

---

For *tangential mindlin/force*, the accumulated elastic tangential force characterizes the contact history, instead of the accumulated tangential displacement. This prevents the dependence of the tangential force on the normal overlap as noted above. The tangential force is given by:

$$\mathbf{F}_t = -\min(\mu_t F_{n0}, \|\mathbf{F}_{te} + \mathbf{F}_{t,damp}\|)\mathbf{t}$$

The increment of the elastic component of the tangential force  $\mathbf{F}_{te}$  is given by:

$$d\mathbf{F}_{te} = -k_t a \mathbf{v}_{t,rel} d\tau$$

The changes in frame of reference of the contacting pair of particles during contact are accounted for by the same formula as above, replacing the accumulated tangential displacement  $\xi$ , by the accumulated tangential elastic force  $F_{te}$ . When the tangential force exceeds the critical force, the tangential force is directly re-scaled to match the value for the critical force:

$$\mathbf{F}_{te} = -\mu_t F_{n0} \mathbf{t} + \mathbf{F}_{t,damp}$$

The same rules as those described for *mindlin* apply regarding the tangential stiffness and mixing of the shear modulus for different particle types.

The *mindlin\_rescale* option uses the same form as *mindlin*, but the magnitude of the tangential displacement is re-scaled as the contact unloads, i.e. if  $a < a_{t_{n-1}}$ :

$$\xi = \xi_{t_{n-1}} \frac{a}{a_{t_{n-1}}}$$

Here,  $t_{n-1}$  indicates the value at the previous time step. This rescaling accounts for the fact that a decrease in the contact area upon unloading leads to the contact being unable to support the previous tangential loading, and spurious energy is created without the rescaling above ([Walton](#)).

---

**Note:** For *mindlin*, a decrease in the tangential force already occurs as the contact unloads, due to the dependence of the tangential force on the normal force described above. By re-scaling  $\xi$ , *mindlin\_rescale* effectively re-scales the tangential force twice, i.e., proportionally to  $a^2$ . This peculiar behavior results from use of the accumulated tangential displacement to characterize the contact history. Although *mindlin\_rescale* remains available for historic reasons and backward compatibility purposes, it should be avoided in favor of *mindlin\_rescale/force*.

---

The *mindlin\_rescale/force* option uses the same form as *mindlin/force*, but the magnitude of the tangential elastic force is re-scaled as the contact unloads, i.e. if  $a < a_{t_{n-1}}$ :

$$\mathbf{F}_{te} = \mathbf{F}_{te,t_{n-1}} \frac{a}{a_{t_{n-1}}}$$

This approach provides a better approximation of the *Mindlin-Deresiewicz* laws and is more consistent than *mindlin\_rescale*. See discussions in *Thornton et al, 2013*, particularly equation 18(b) of that work and associated discussion, and *Agnolin and Roux, 2007*, particularly Appendix A.

---

The optional *rolling* keyword enables rolling friction, which resists pure rolling motion of particles. The options currently supported are:

1. *none*
2. *sds* :  $k_{roll}, \gamma_{roll}, \mu_{roll}$

If the *rolling* keyword is not specified, the model defaults to *none*.

For *rolling sds*, rolling friction is computed via a spring-dashpot-slider, using a ‘pseudo-force’ formulation, as detailed by *Luding*. Unlike the formulation in *Marshall*, this allows for the required adjustment of rolling displacement due to changes in the frame of reference of the contacting pair. The rolling pseudo-force is computed analogously to the tangential force:

$$\mathbf{F}_{roll,0} = k_{roll}\xi_{roll} - \gamma_{roll}\mathbf{v}_{roll}$$

Here,  $\mathbf{v}_{roll} = -R(\boldsymbol{\Omega}_i - \boldsymbol{\Omega}_j) \times \mathbf{n}$  is the relative rolling velocity, as given in *Wang et al* and *Luding*. This differs from the expressions given by *Kuhn and Bagi* and used in *Marshall*; see *Wang et al* for details. The rolling displacement is given by:

$$\xi_{roll} = \int_{t_0}^t \mathbf{v}_{roll}(\tau) d\tau$$

A Coulomb friction criterion truncates the rolling pseudo-force if it exceeds a critical value:

$$\mathbf{F}_{roll} = \min(\mu_{roll}F_{n,0}, \|\mathbf{F}_{roll,0}\|)\mathbf{k}$$

Here,  $\mathbf{k} = \mathbf{v}_{roll}/\|\mathbf{v}_{roll}\|$  is the direction of the pseudo-force. As with tangential displacement, the rolling displacement is rescaled when the critical force is exceeded, so that the spring length corresponds the critical force. Additionally, the displacement is adjusted to account for rotations of the frame of reference of the two contacting particles in a manner analogous to the tangential displacement.

The rolling pseudo-force does not contribute to the total force on either particle (hence ‘pseudo’), but acts only to induce an equal and opposite torque on each particle, according to:

$$\tau_{roll,i} = R_{eff}\mathbf{n} \times \mathbf{F}_{roll}$$

$$\tau_{roll,j} = -\tau_{roll,i}$$

---

The optional *twisting* keyword enables twisting friction, which resists rotation of two contacting particles about the vector  $\mathbf{n}$  that connects their centers. The options currently supported are:

1. *none*
2. *sds* :  $k_{twist}, \gamma_{twist}, \mu_{twist}$
3. *marshall*

If the *twisting* keyword is not specified, the model defaults to *none*.

For both *twisting sds* and *twisting marshall*, a history-dependent spring-dashpot-slider is used to compute the twisting torque. Because twisting displacement is a scalar, there is no need to adjust for changes in the frame of reference due to rotations of the particle pair. The formulation in *Marshall* therefore provides the most straightforward treatment:

$$\tau_{twist,0} = -k_{twist}\xi_{twist} - \gamma_{twist}\Omega_{twist}$$

Here  $\xi_{twist} = \int_{t_0}^t \Omega_{twist}(\tau) d\tau$  is the twisting angular displacement, and  $\Omega_{twist} = (\Omega_i - \Omega_j) \cdot \mathbf{n}$  is the relative twisting angular velocity. The torque is then truncated according to:

$$\tau_{twist} = \min(\mu_{twist} F_{n,0}, \tau_{twist,0})$$

Similar to the sliding and rolling displacement, the angular displacement is rescaled so that it corresponds to the critical value if the twisting torque exceeds this critical value:

$$\xi_{twist} = \frac{1}{k_{twist}} (\mu_{twist} F_{n,0} \text{sgn}(\Omega_{twist}) - \gamma_{twist} \Omega_{twist})$$

For *twisting sds*, the coefficients  $k_{twist}$ ,  $\gamma_{twist}$  and  $\mu_{twist}$  are simply the user input parameters that follow the *twisting sds* keywords in the *pair\_coeff* command.

For *twisting\_marshall*, the coefficients are expressed in terms of sliding friction coefficients, as discussed in *Marshall* (see equations 32 and 33 of that work):

$$k_{twist} = 0.5 k_t a^2$$

$$\eta_{twist} = 0.5 \eta_t a^2$$

$$\mu_{twist} = \frac{2}{3} a \mu_t$$

Finally, the twisting torque on each particle is given by:

$$\tau_{twist,i} = \tau_{twist} \mathbf{n}$$

$$\tau_{twist,j} = -\tau_{twist,i}$$

If two particles are moving away from each other while in contact, there is a possibility that the particles could experience an effective attractive force due to damping. If the optional *limit\_damping* keyword is used, this option will zero out the normal component of the force if there is an effective attractive force. This keyword cannot be used with the JKR or DMT models.

The *granular* pair style can reproduce the behavior of the *pair gran/\** styles with the appropriate settings (some very minor differences can be expected due to corrections in displacement history frame-of-reference, and the application of the torque at the center of the contact rather than at each particle). The first example above is equivalent to *pair gran/hooke 1000.0 NULL 50.0 50.0 0.4 1*. The second example is equivalent to *pair gran/hooke/history 1000.0 500.0 50.0 50.0 0.4 1*. The third example is equivalent to *pair gran/hertz/history 1000.0 500.0 50.0 50.0 0.4 1*.

LAMMPS automatically sets pairwise cutoff values for *pair\_style granular* based on particle radii (and in the case of *jkr* pull-off distances). In the vast majority of situations, this is adequate. However, a cutoff value can optionally be appended to the *pair\_style granular* command to specify a global cutoff (i.e. a cutoff for all atom types). Additionally, the optional *cutoff* keyword can be passed to the *pair\_coeff* command, followed by a cutoff value. This will set a pairwise cutoff for the atom types in the *pair\_coeff* command. These options may be useful in some rare cases where the automatic cutoff determination is not sufficient, e.g. if particle diameters are being modified via the *fix adapt* command. In that case, the global cutoff specified as part of the *pair\_style granular* command is applied to all atom types, unless it is overridden for a given atom type combination by the *cutoff* value specified in the *pair coeff* command. If *cutoff* is only specified in the *pair coeff* command and no global cutoff is appended to the *pair\_style granular* command, then LAMMPS will use that cutoff for the specified atom type combination, and automatically set pairwise cutoffs for the remaining atom types.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.118.4 Mixing, shift, table, tail correction, restart, rRESPA info

The *pair\_modify* mix, shift, table, and tail options are not relevant for granular pair styles.

Mixing of coefficients is carried out using geometric averaging for most quantities, e.g. if friction coefficient for type 1-type 1 interactions is set to  $\mu_1$ , and friction coefficient for type 2-type 2 interactions is set to  $\mu_2$ , the friction coefficient for type1-type2 interactions is computed as  $\sqrt{\mu_1\mu_2}$  (unless explicitly specified to a different value by a *pair\_coeff 1 2 ...* command). The exception to this is elastic modulus, only applicable to *hertz/material*, *dmt* and *jkr* normal contact models. In that case, the effective elastic modulus is computed as:

$$E_{eff,ij} = \left( \frac{1 - \nu_i^2}{E_i} + \frac{1 - \nu_j^2}{E_j} \right)^{-1}$$

If the *i-j* coefficients  $E_{ij}$  and  $\nu_{ij}$  are explicitly specified, the effective modulus is computed as:

$$E_{eff,ij} = \left( \frac{1 - \nu_{ij}^2}{E_{ij}} + \frac{1 - \nu_{ij}^2}{E_{ij}} \right)^{-1}$$

or

$$E_{eff,ij} = \frac{E_{ij}}{2(1 - \nu_{ij})}$$

These pair styles write their information to *binary restart files*, so a *pair\_style* command does not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

The *single()* function of these pair styles returns 0.0 for the energy of a pairwise interaction, since energy is not conserved in these dissipative potentials. It also returns only the normal component of the pairwise interaction force. However, the *single()* function also calculates 12 extra pairwise quantities. The first 3 are the components of the tangential force between particles I and J, acting on particle I. The fourth is the magnitude of this tangential force. The next 3 (5-7) are the components of the rolling torque acting on particle I. The next entry (8) is the magnitude of the rolling torque. The next entry (9) is the magnitude of the twisting torque acting about the vector connecting the two particle centers. The last 3 (10-12) are the components of the vector connecting the centers of the two particles ( $x_I - x_J$ ).

These extra quantities can be accessed by the *compute pair/local* command, as *p1*, *p2*, ..., *p12*.

---

### 19.118.5 Restrictions

All the granular pair styles are part of the GRANULAR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These pair styles require that atoms store torque and angular velocity ( $\omega$ ) as defined by the *atom\_style*. They also require a per-particle radius is stored. The *sphere* atom style does all of this.

This pair style requires you to use the *comm\_modify vel yes* command so that velocities are stored by ghost atoms.

These pair styles will not restart exactly when using the *read\_restart* command, though they should provide statistically similar results. This is because the forces they compute depend on atom velocities. See the *read\_restart* command for more details.

### 19.118.6 Related commands

*pair\_coeff pair gran/\**

### 19.118.7 Default

For the *pair\_coeff* settings: *damping viscoelastic, rolling none, twisting none*.

### 19.118.8 References

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(Mindlin and Deresiewicz, 1953) Mindlin, R.D., & Deresiewicz, H (1953). Elastic Spheres in Contact under Varying Oblique Force. J. Appl. Mech., ASME 20, 327-344.

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## 19.119 pair\_style lj/gromacs command

Accelerator Variants: *lj/gromacs/gpu*, *lj/gromacs/kk*, *lj/gromacs/omp*

## 19.120 pair\_style lj/gromacs/coul/gromacs command

Accelerator Variants: *lj/gromacs/coul/gromacs/kk*, *lj/gromacs/coul/gromacs/omp*

### 19.120.1 Syntax

```
pair_style style args
```

- style = *lj/gromacs* or *lj/gromacs/coul/gromacs*
- args = list of arguments for a particular style

```
lj/gromacs args = inner outer
```

```
inner, outer = global switching cutoffs for Lennard Jones
```

```
lj/gromacs/coul/gromacs args = inner outer (inner2) (outer2)
```

```
inner, outer = global switching cutoffs for Lennard Jones (and Coulombic if only 2_
```

```
↪args)
```

```
inner2, outer2 = global switching cutoffs for Coulombic (optional)
```

### 19.120.2 Examples

```
pair_style lj/gromacs 9.0 12.0
pair_coeff * * 100.0 2.0
pair_coeff 2 2 100.0 2.0 8.0 10.0

pair_style lj/gromacs/coul/gromacs 9.0 12.0
pair_style lj/gromacs/coul/gromacs 8.0 10.0 7.0 9.0
pair_coeff * * 100.0 2.0
```



### 19.120.3 Description

The *lj/gromacs* styles compute shifted LJ and Coulombic interactions with an additional switching function  $S(r)$  that ramps the energy and force smoothly to zero between an inner and outer cutoff. It is a commonly used potential in the GROMACS MD code and for the coarse-grained models of (*Marrink*).

$$\begin{aligned}
 E_{LJ} &= 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] + S_{LJ}(r) & r < r_c \\
 E_C &= \frac{Cq_iq_j}{\epsilon r} + S_C(r) & r < r_c \\
 S(r) &= C & r < r_1 \\
 S(r) &= \frac{A}{3}(r - r_1)^3 + \frac{B}{4}(r - r_1)^4 + C & r_1 < r < r_c \\
 A &= (-3E'(r_c) + (r_c - r_1)E''(r_c))/(r_c - r_1)^2 \\
 B &= (2E'(r_c) - (r_c - r_1)E''(r_c))/(r_c - r_1)^3 \\
 C &= -E(r_c) + \frac{1}{2}(r_c - r_1)E'(r_c) - \frac{1}{12}(r_c - r_1)^2E''(r_c)
 \end{aligned}$$

$r_1$  is the inner cutoff;  $r_c$  is the outer cutoff. The coefficients A, B, and C are computed by LAMMPS to perform the shifting and smoothing. The function  $S(r)$  is actually applied once to each term of the LJ formula and once to the Coulombic formula, so there are 2 or 3 sets of A,B,C coefficients depending on which *pair\_style* is used. The boundary conditions applied to the smoothing function are as follows:  $S'(r_1) = S''(r_1) = 0$ ,  $S(r_c) = -E(r_c)$ ,  $S'(r_c) = -E'(r_c)$ , and  $S''(r_c) = -E''(r_c)$ , where  $E(r)$  is the corresponding term in the LJ or Coulombic potential energy function. Single and double primes denote first and second derivatives with respect to  $r$ , respectively.

The inner and outer cutoff for the LJ and Coulombic terms can be the same or different depending on whether 2 or 4 arguments are used in the *pair\_style* command. The inner LJ cutoff must be  $> 0$ , but the inner Coulombic cutoff can be  $\geq 0$ .

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- inner (distance units)
- outer (distance units)

Note that sigma is defined in the LJ formula as the zero-crossing distance for the potential, not as the energy minimum at  $2^{1/6}\sigma$ .

The last 2 coefficients are optional inner and outer cutoffs for style *lj/gromacs*. If not specified, the global *inner* and *outer* values are used.

The last 2 coefficients cannot be used with style *lj/gromacs/coul/gromacs* because this force field does not allow varying cutoffs for individual atom pairs; all pairs use the global cutoff(s) specified in the *pair\_style* command.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.120.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the lj/cut pair styles can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

None of the GROMACS pair styles support the *pair\_modify* shift option, since the Lennard-Jones portion of the pair interaction is already smoothed to 0.0 at the cutoff.

The *pair\_modify* table option is not relevant for this pair style.

None of the GROMACS pair styles support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure, since there are no corrections for a potential that goes to 0.0 at the cutoff.

All of the GROMACS pair styles write their information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

All of the GROMACS pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner, middle, outer* keywords.

---

#### 19.120.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 19.120.6 Related commands

*pair\_coeff*

#### 19.120.7 Default

none

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### 19.121 pair\_style gw command

### 19.122 pair\_style gw/zbl command

#### 19.122.1 Syntax

```
pair_style style
```

- style = *gw* or *gw/zbl*

### 19.122.2 Examples

```
pair_style gw
pair_coeff * * SiC.gw Si C C

pair_style gw/zbl
pair_coeff * * SiC.gw.zbl C Si
```

### 19.122.3 Description

The *gw* style computes a 3-body *Gao-Weber* potential; similarly *gw/zbl* combines this potential with a modified repulsive ZBL core function in a similar fashion as implemented in the *tersoff/zbl* pair style.

Unfortunately the author of this contributed code has not been able to submit a suitable documentation explaining the details of the potentials. The LAMMPS developers thus have finally decided to release the code anyway with only the technical explanations. For details of the model and the parameters, please refer to the linked publication.

Only a single `pair_coeff` command is used with the *gw* and *gw/zbl* styles which specifies a Gao-Weber potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of GW elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the potential file.

As an example, imagine a file `SiC.gw` has Gao-Weber values for Si and C. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Si, and the fourth to be C, you would use the following `pair_coeff` command:

```
pair_coeff * * SiC.gw Si Si Si C
```

The first 2 arguments must be `**` so as to span all LAMMPS atom types. The first three Si arguments map LAMMPS atom types 1,2,3 to the Si element in the GW file. The final C argument maps LAMMPS atom type 4 to the C element in the GW file. If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *gw* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

Gao-Weber files in the *potentials* directory of the LAMMPS distribution have a “.gw” suffix. Gao-Weber with ZBL files have a “.gz.zbl” suffix. The structure of the potential files is similar to other many-body potentials supported by LAMMPS. You have to refer to the comments in the files and the literature to learn more details.

### 19.122.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, mixing is performed by LAMMPS as described above from values in the potential file.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.122.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

The Gao-Weber potential files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use the GW potential with any LAMMPS units, but you would need to create your own GW potential file with coefficients listed in the appropriate units if your simulation does not use “metal” units.

### 19.122.6 Related commands

*pair\_coeff*

### 19.122.7 Default

none

---

**(Gao)** Gao and Weber, Nuclear Instruments and Methods in Physics Research B 191 (2012) 504.

## 19.123 pair\_style harmonic/cut command

Accelerator Variants: *harmonic/cut/omp*

### 19.123.1 Syntax

```
pair_style style
```

- style = *harmonic/cut*

## 19.123.2 Examples

```
pair_style harmonic/cut
pair_coeff * * 0.2 2.0
pair_coeff 1 1 0.5 2.5
```

## 19.123.3 Description

Style *harmonic/cut* computes pairwise repulsive-only harmonic interactions with the formula

$$E = k(r_c - r)^2 \quad r < r_c$$

where  $r_c$  is the cutoff. Note that the usual 1/2 factor is included in  $k$ .

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $k$  (energy/distance<sup>2</sup> units)
- $r_c$  (distance units)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

## 19.123.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the  $k$  and  $r_c$  coefficients can be mixed. The default mix value is *geometric*. See the “*pair\_modify*” command for details.

Since the potential is zero at and beyond the cutoff parameter by construction, there is no need to support the *pair\_modify* shift or tail options for the energy and pressure of the pair interaction.

These pair styles write their information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

### 19.123.5 Restrictions

The *harmonic/cut* pair style is only enabled if LAMMPS was built with the EXTRA-PAIR package. See the *Build package* page for more info.

### 19.123.6 Related commands

*pair\_coeff*

### 19.123.7 Default

none

## 19.124 `pair_style hbond/dreiding/lj` command

Accelerator Variants: *hbond/dreiding/lj/omp*

## 19.125 `pair_style hbond/dreiding/morse` command

Accelerator Variants: *hbond/dreiding/morse/omp*

### 19.125.1 Syntax

```
pair_style style N inner_distance_cutoff outer_distance_cutoff angle_cutof
```

- style = *hbond/dreiding/lj* or *hbond/dreiding/morse*
- n = cosine angle periodicity
- inner\_distance\_cutoff = global inner cutoff for Donor-Acceptor interactions (distance units)
- outer\_distance\_cutoff = global cutoff for Donor-Acceptor interactions (distance units)
- angle\_cutoff = global angle cutoff for Acceptor-Hydrogen-Donor interactions (degrees)

### 19.125.2 Examples

```
pair_style hybrid/overlay lj/cut 10.0 hbond/dreiding/lj 4 9.0 11.0 90  
pair_coeff 1 2 hbond/dreiding/lj 3 i 9.5 2.75 4 9.0 11.0 90.0  
  
pair_style hybrid/overlay lj/cut 10.0 hbond/dreiding/morse 2 9.0 11.0 90  
pair_coeff 1 2 hbond/dreiding/morse 3 i 3.88 1.7241379 2.9 2 9 11 90
```

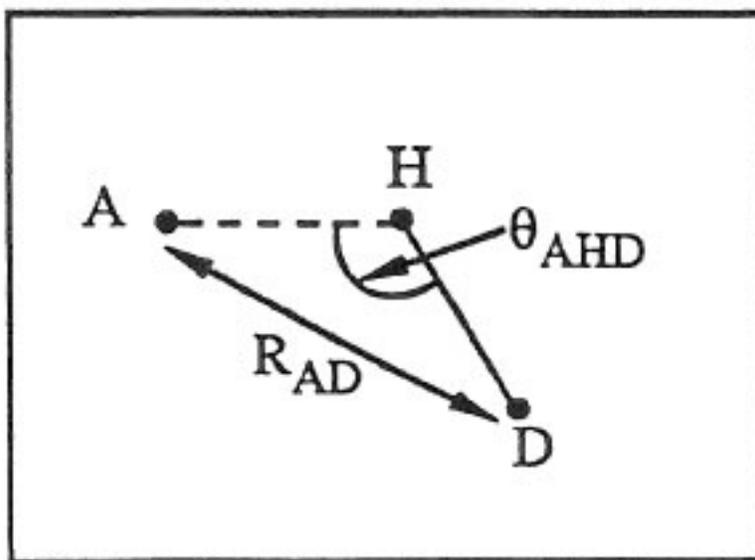
### 19.125.3 Description

The *hbond/dreiding* styles compute the Acceptor-Hydrogen-Donor (AHD) 3-body hydrogen bond interaction for the *DREIDING* force field, given by:

$$\begin{aligned}
 E &= [LJ(r)|Morse(r)] && r < r_{in} \\
 &= S(r) * [LJ(r)|Morse(r)] && r_{in} < r < r_{out} \\
 &= 0 && r > r_{out} \\
 LJ(r) &= AR^{-12} - BR^{-10} \cos^n \theta = \epsilon \left\{ 5 \left[ \frac{\sigma}{r} \right]^{12} - 6 \left[ \frac{\sigma}{r} \right]^{10} \right\} \cos^n \theta \\
 Morse(r) &= D_0 \{ \chi^2 - 2\chi \} \cos^n \theta = D_0 \left\{ e^{-2\alpha(r-r_0)} - 2e^{-\alpha(r-r_0)} \right\} \cos^n \theta \\
 S(r) &= \frac{[r_{out}^2 - r^2]^2 [r_{out}^2 + 2r^2 - 3r_{in}^2]}{[r_{out}^2 - r_{in}^2]^3}
 \end{aligned}$$

where  $r_{in}$  is the inner spline distance cutoff,  $r_{out}$  is the outer distance cutoff,  $\theta_c$  is the angle cutoff, and  $n$  is the cosine periodicity.

Here,  $r$  is the radial distance between the donor (D) and acceptor (A) atoms and  $\theta$  is the bond angle between the acceptor, the hydrogen (H) and the donor atoms:



These 3-body interactions can be defined for pairs of acceptor and donor atoms, based on atom types. For each donor/acceptor atom pair, the third atom in the interaction is a hydrogen permanently bonded to the donor atom, e.g. in a bond list read in from a data file via the *read\_data* command. The atom types of possible hydrogen atoms for each donor/acceptor type pair are specified by the *pair\_coeff* command (see below).

Style *hbond/dreiding/lj* is the original DREIDING potential of (Mayo). It uses a LJ 12/10 functional for the Donor-Acceptor interactions. To match the results in the original paper, use  $n = 4$ .

Style *hbond/dreiding/morse* is an improved version using a Morse potential for the Donor-Acceptor interactions. (Liu) showed that the Morse form gives improved results for Dendrimer simulations, when  $n = 2$ .

See the *Howto bioFF* page for more information on the DREIDING force field.

**Note:** Because the Dreiding hydrogen bond potential is only one portion of an overall force field which typically includes other pairwise interactions, it is common to use it as a sub-style in a *pair\_style hybrid/overlay* command,

where another pair style provides the repulsive core interaction between pairs of atoms, e.g. a  $1/r^{12}$  Lennard-Jones repulsion.

---

**Note:** When using the `hbond/dreiding` pair styles with `pair_style hybrid/overlay`, you should explicitly define pair interactions between the donor atom and acceptor atoms, (as well as between these atoms and ALL other atoms in your system). Whenever `pair_style hybrid/overlay` is used, ordinary mixing rules are not applied to atoms like the donor and acceptor atoms because they are typically referenced in multiple pair styles. Neglecting to do this can cause difficult-to-detect physics problems.

---

**Note:** In the original Dreiding force field paper 1-4 non-bonded interactions ARE allowed. If this is desired for your model, use the `special_bonds` command (e.g. “`special_bonds lj 0.0 0.0 1.0`”) to turn these interactions on.

---

The following coefficients must be defined for pairs of eligible donor/acceptor types via the `pair_coeff` command as in the examples above.

---

**Note:** Unlike other pair styles and their associated `pair_coeff` commands, you do not need to specify `pair_coeff` settings for all possible I,J type pairs. Only I,J type pairs for atoms which act as joint donors/acceptors need to be specified; all other type pairs are assumed to be inactive.

---

**Note:** A `pair_coeff` command can be specified multiple times for the same donor/acceptor type pair. This enables multiple hydrogen types to be assigned to the same donor/acceptor type pair. For other pair\_styles, if the `pair_coeff` command is re-used for the same I,J type pair, the settings for that type pair are overwritten. For the hydrogen bond potentials this is not the case; the settings are cumulative. This means the only way to turn off a previous setting, is to re-use the `pair_style` command and start over.

---

For the `hbond/dreiding/lj` style the list of coefficients is as follows:

- $K$  = hydrogen atom type = 1 to Ntypes
- donor flag =  $i$  or  $j$
- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- $n$  = exponent in formula above
- distance cutoff  $r_{in}$  (distance units)
- distance cutoff  $r_{out}$  (distance units)
- angle cutoff (degrees)

For the `hbond/dreiding/morse` style the list of coefficients is as follows:

- $K$  = hydrogen atom type = 1 to Ntypes
- donor flag =  $i$  or  $j$
- $D_0$  (energy units)
- $\alpha$  (1/distance units)
- $r_0$  (distance units)



- $n$  = exponent in formula above
- distance cutoff  $r_{in}$  (distance units)
- distance cutoff  $r_{out}$  (distance units)
- angle cutoff (degrees)

A single hydrogen atom type K can be specified, or a wild-card asterisk can be used in place of or in conjunction with the K arguments to select multiple types as hydrogen atoms. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. See the [pair\\_coeff](#) command page for details.

If the donor flag is  $i$ , then the atom of type I in the pair\_coeff command is treated as the donor, and J is the acceptor. If the donor flag is  $j$ , then the atom of type J in the pair\_coeff command is treated as the donor and I is the donor. This option is required because the [pair\\_coeff](#) command requires that  $I \leq J$ .

$\epsilon$  and  $\sigma$  are settings for the hydrogen bond potential based on a Lennard-Jones functional form. Note that sigma is defined as the zero-crossing distance for the potential, not as the energy minimum at  $2^{1/6}\sigma$ .

$D_0$  and  $\alpha$  and  $r_0$  are settings for the hydrogen bond potential based on a Morse functional form.

The last 3 coefficients for both styles are optional. If not specified, the global n, distance cutoff, and angle cutoff specified in the pair\_style command are used. If you wish to only override the second or third optional parameter, you must also specify the preceding optional parameters.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* [command-line switch](#) when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.125.4 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles do not support mixing. You must explicitly identify each donor/acceptor type pair.

These styles do not support the *pair\_modify* shift option for the energy of the interactions.

The *pair\_modify* table option is not relevant for these pair styles.

These pair styles do not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

These pair styles do not write their information to *binary restart files*, so pair\_style and pair\_coeff commands need to be re-specified in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

These pair styles tally a count of how many hydrogen bonding interactions they calculate each timestep and the hbond energy. These quantities can be accessed via the *compute pair* command as a vector of values of length 2.

To print these quantities to the log file (with a descriptive column heading) the following commands could be included in an input script:

```
compute hb all pair hbond/dreiding/lj
variable n_hbond equal c_hb[1] #number hbonds
variable E_hbond equal c_hb[2] #hbond energy
thermo_style custom step temp epair v_E_hbond
```

---

### 19.125.5 Restrictions

none

### 19.125.6 Related commands

*pair\_coeff*

### 19.125.7 Default

none

---

**(Mayo)** Mayo, Olfason, Goddard III, J Phys Chem, 94, 8897-8909 (1990).

**(Liu)** Liu, Bryantsev, Diallo, Goddard III, J. Am. Chem. Soc 131 (8) 2798 (2009)

## 19.126 pair\_style hdnnp command

### 19.126.1 Syntax

```
pair_style hdnnp cutoff keyword value ...
```

- cutoff = short-range cutoff of HDNNP (maximum symmetry function cutoff radius)
- zero or more keyword/value pairs may be appended
- keyword = *dir* or *showew* or *showewsum* or *maxew* or *resetew* or *cflength* or *cfenergy*
- value depends on the preceding keyword:

*dir* value = directory

directory = Path to HDNNP configuration files

*showew* value = *yes* or *no*

*showewsum* value = *summary*

summary = Write EW summary every this many timesteps (0 turns summary off)

*maxew* value = *threshold*

threshold = Maximum number of EWs allowed

*resetew* value = *yes* or *no*

*cflength* value = *length*

length = Length unit conversion factor

*cfenergy* value = *energy*

energy = Energy unit conversion factor

## 19.126.2 Examples

```
pair_style hdnnp 6.35 showew yes showewsum 100 maxew 1000 resetew yes cflength 1.
↪8897261328 cfenergy 0.0367493254
pair_coeff * * H O

pair_style hdnnp 6.01 dir "./" showewsum 10000
pair_coeff * * S Cu NULL Cu
```

## 19.126.3 Description

This pair style adds an interaction based on the high-dimensional neural network potential (HDNNP) method as presented in (*Behler and Parrinello 2007*). HDNNPs are machine learning potentials which require careful training of neural networks prior to application in MD simulations. The pair style uses an interface to the *n2p2* library (*Singraber, Behler and Dellago 2019*) which is available on Github [here](#). Please see the *n2p2* documentation for further details. *n2p2* (and hence this pair style) is compatible with neural network potentials trained with its own tools (*Singraber et al 2019*) and with RuNNer.

Only a single *pair\_coeff* command with two asterisk wild-cards is used with this pair style. Its additional arguments define the mapping of LAMMPS atom types to *n2p2* elements.

```
pair_coeff * * H O
```

In the above example LAMMPS types 1 and 2 are mapped to the elements “H” and “O” in *n2p2*, respectively. Multiple types may map to the same element, or some types may not be mapped at all. For example, if the LAMMPS simulation has four atom types, the command

```
pair_coeff * * H H O NULL
```

maps atom types 1 and 2 to the element “H”, type 3 to “O” and type 4 is not mapped (indicated by NULL). Atoms mapped to NULL are ignored by the HDNNP calculation, i.e. they do not contribute in any way to the evaluation of HDNNP energies and forces. This may be useful in a setup with *hybrid pair styles*.

The mandatory pair style argument *cutoff* must match the short-range cutoff radius of the HDNNP. This corresponds to the maximum cutoff radius of all symmetry functions (the atomic environment descriptors of HDNNPs) used.

**Note:** The cutoff must be given in LAMMPS length units, even if the neural network potential has been trained using a different unit system (see remarks about the *cflength* and *cfenergy* keywords below for details).

The numeric value may be slightly larger than the actual maximum symmetry function cutoff radius (to account for rounding errors when converting units), but must not be smaller.

Use the *dir* keyword to specify the directory containing the HDNNP configuration files. The directory must contain *input.nn* with neural network and symmetry function setup, *scaling.data* with symmetry function scaling data and *weights.??? .data* with weight parameters for each element.

The keyword *showew* can be used to turn on/off the display of extrapolation warnings (EWs) which are issued whenever a symmetry function value is out of bounds defined by minimum/maximum values in *scaling.data*. An extrapolation warning may look like this:

```
### NNP EXTRAPOLATION WARNING ### STRUCTURE:      0 ATOM:      119 ELEMENT: Cu SYMFUNC:↵
↪ 32 TYPE: 3 VALUE: 2.166E-02 MIN: 2.003E-05 MAX: 1.756E-02
```

stating that the value 2.166E-02 of symmetry function 32 of type 3 (Narrow Angular symmetry function), element Cu (see the log file for a symmetry function listing) was out of bounds (maximum in `scaling.data` is 1.756E-02) for atom 119. Here, the atom index refers to the LAMMPS tag (global index) and the structure index is used to print out the MPI rank the atom belongs to.

---

**Note:** The `showew` keyword should only be set to `yes` for debugging purposes. Extrapolation warnings may add lots of overhead as they are communicated each timestep. Also, if the simulation is run in a region where the HDNNP was not correctly trained, lots of extrapolation warnings may clog log files and the console. In a production run use `showewsum` instead.

---

The keyword `showewsum` can be used to get an overview of extrapolation warnings occurring during an MD simulation. The argument specifies the interval at which extrapolation warning summaries are displayed and logged. An EW summary may look like this:

```
### NNP EW SUMMARY ### TS:      100 EW      11 EWPERSTEP  1.100E-01
```

Here, at timestep 100 the occurrence of 11 extrapolation warnings since the last summary is reported, which corresponds to an EW rate of 0.11 per timestep. Setting `showewsum` to 0 deactivates the EW summaries.

A maximum number of allowed extrapolation warnings can be specified with the `maxew` keyword. If the number of EWs exceeds the `maxew` argument the simulation is stopped. Note however that this is merely an approximate threshold since the check is only performed at the end of each timestep and each MPI process counts individually to minimize communication overhead.

The keyword `resetew` alters the behavior of the above mentioned `maxew` threshold. If `resetew` is set to `yes` the threshold is applied on a per-timestep basis and the internal EW counters are reset at the beginning of each timestep. With `resetew` set to `no` the counters accumulate EWs along the whole trajectory.

If the training of a neural network potential has been performed with different physical units for length and energy than those set in LAMMPS, it is still possible to use the potential when the unit conversion factors are provided via the `cflength` and `cfenergy` keywords. If for example, the HDNNP was parameterized with Bohr and Hartree training data and symmetry function parameters (i.e. distances and energies in “input.nn” are given in Bohr and Hartree) but LAMMPS is set to use *metal* units (Angstrom and eV) the correct conversion factors are:

```
cflength 1.8897261328
cfenergy 0.0367493254
```

Thus, arguments of `cflength` and `cfenergy` are the multiplicative factors required to convert lengths and energies given in LAMMPS units to respective quantities in native HDNNP units (1 Angstrom = 1.8897261328 Bohr, 1 eV = 0.0367493254 Hartree).

---

## 19.126.4 Mixing, shift, table, tail correction, restart, rRESPA info

This style does not support mixing. The `pair_coeff` command should only be invoked with asterisk wild cards (see above).

This style does not support the `pair_modify` shift, table, and tail options.

This style does not write information to *binary restart files*. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

This style can only be used via the `pair` keyword of the `run_style respa` command. It does not support the *inner*, *middle*, *outer* keywords.

## 19.126.5 Restrictions

This pair style is part of the ML-HDNNP package. It is only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

Please report bugs and feature requests to the [n2p2 GitHub issue page](#).

### Related commands

*pair\_coeff*, *pair\_hybrid*, *units*

### Default

The default options are *dir* = "hdnnp", *showew* = yes, *showewsum* = 0, *maxew* = 0, *resetew* = no, *cflength* = 1.0, *cfenergy* = 1.0.

---

**(Behler and Parrinello 2007)** Behler, J.; Parrinello, M. Generalized Neural-Network Representation of High-Dimensional Potential-Energy Surfaces. *Phys. Rev. Lett.* 2007, 98 (14), 146401.

**(Singraber, Behler and Dellago 2019)** Singraber, A.; Behler, J.; Dellago, C. Library-Based LAMMPS Implementation of High-Dimensional Neural Network Potentials. *J. Chem. Theory Comput.* 2019, 15 (3), 1827-1840

**(Singraber et al 2019)** Singraber, A.; Morawietz, T.; Behler, J.; Dellago, C. Parallel Multistream Training of High-Dimensional Neural Network Potentials. *J. Chem. Theory Comput.* 2019, 15 (5), 3075-3092.

## 19.127 pair\_style hybrid command

Accelerator Variants: *hybrid/kk*

## 19.128 pair\_style hybrid/overlay command

Accelerator Variants: *hybrid/overlay/kk*

## 19.129 pair\_style hybrid/scaled command

### 19.129.1 Syntax

```
pair_style hybrid style1 args style2 args ...
pair_style hybrid/overlay style1 args style2 args ...
pair_style hybrid/scaled factor1 style1 args factor2 style 2 args ...
```

- style1,style2 = list of one or more pair styles and their arguments
- factor1,factor2 = scale factors for pair styles, may be a variable

## 19.129.2 Examples

```

pair_style hybrid lj/cut/coul/cut 10.0 eam lj/cut 5.0
pair_coeff 1*2 1*2 eam niu3
pair_coeff 3 3 lj/cut/coul/cut 1.0 1.0
pair_coeff 1*2 3 lj/cut 0.5 1.2

pair_style hybrid/overlay lj/cut 2.5 coul/long 2.0
pair_coeff * * lj/cut 1.0 1.0
pair_coeff * * coul/long

pair_style hybrid/scaled 0.5 tersoff 0.5 sw
pair_coeff * * tersoff Si.tersoff Si
pair_coeff * * sw Si.sw Si

variable one equal ramp(1.0,0.0)
variable two equal 1.0-v_one
pair_style hybrid/scaled v_one lj/cut 2.5 v_two morse 2.5
pair_coeff 1 1 lj/cut 1.0 1.0 2.5
pair_coeff 1 1 morse 1.0 1.0 1.0 2.5

```

## 19.129.3 Description

The *hybrid*, *hybrid/overlay*, and *hybrid/scaled* styles enable the use of multiple pair styles in one simulation. With the *hybrid* style, exactly one pair style is assigned to each pair of atom types. With the *hybrid/overlay* and *hybrid/scaled* styles, one or more pair styles can be assigned to each pair of atom types. The assignment of pair styles to type pairs is made via the *pair\_coeff* command. The major difference between the *hybrid/overlay* and *hybrid/scaled* styles is that the *hybrid/scaled* adds a scale factor for each sub-style contribution to forces, energies and stresses. Because of the added complexity, the *hybrid/scaled* style has more overhead and thus may be slower than *hybrid/overlay*.

Here are two examples of hybrid simulations. The *hybrid* style could be used for a simulation of a metal droplet on a LJ surface. The metal atoms interact with each other via an *eam* potential, the surface atoms interact with each other via a *lj/cut* potential, and the metal/surface interaction is also computed via a *lj/cut* potential. The *hybrid/overlay* style could be used as in the second example above, where multiple potentials are superimposed in an additive fashion to compute the interaction between atoms. In this example, using *lj/cut* and *coul/long* together gives the same result as if the *lj/cut/coul/long* potential were used by itself. In this case, it would be more efficient to use the single combined potential, but in general any combination of pair potentials can be used together in to produce an interaction that is not encoded in any single *pair\_style* file, e.g. adding Coulombic forces between granular particles. Another limitation of using the *hybrid/overlay* variant, that it does not generate *lj/cut* parameters for mixed atom types from a mixing rule due to restrictions discussed below.

If the *hybrid/scaled* style is used instead of *hybrid/overlay*, contributions from sub-styles are weighted by their scale factors, which may be fractional or even negative. Furthermore the scale factors may be variables that may change during a simulation. This enables switching smoothly between two different pair styles or two different parameter sets during a run.

All pair styles that will be used are listed as “sub-styles” following the *hybrid* or *hybrid/overlay* keyword, in any order. In case of the *hybrid/scaled* pair style, each sub-style is prefixed with a scale factor. The scale factor is either a floating point number or an equal style (or equivalent) variable. Each sub-style’s name is followed by its usual arguments, as illustrated in the examples above. See the doc pages of the individual pair styles for a listing and explanation of the appropriate arguments for them.

Note that an individual pair style can be used multiple times as a sub-style. For efficiency reasons this should only be done if your model requires it. E.g. if you have different regions of Si and C atoms and wish to use a Tersoff potential

for pure Si for one set of atoms, and a Tersoff potential for pure C for the other set (presumably with some third potential for Si-C interactions), then the sub-style *tersoff* could be listed twice. But if you just want to use a Lennard-Jones or other pairwise potential for several different atom type pairs in your model, then you should just list the sub-style once and use the `pair_coeff` command to assign parameters for the different type pairs.

**Note:** There is one exception to this option to list an individual pair style multiple times: GPU-enabled pair styles in the GPU package. This is because the GPU package currently assumes that only one instance of a pair style is being used.

In the `pair_coeff` commands, the name of a pair style must be added after the I,J type specification, with the remaining coefficients being those appropriate to that style. If the pair style is used multiple times in the `pair_style` command, then an additional numeric argument must also be specified which is a number from 1 to M where M is the number of times the sub-style was listed in the pair style command. The extra number indicates which instance of the sub-style these coefficients apply to.

For example, consider a simulation with 3 atom types: types 1 and 2 are Ni atoms, type 3 are LJ atoms with charges. The following commands would set up a hybrid simulation:

```
pair_style hybrid eam/alloy lj/cut/coul/cut 10.0 lj/cut 8.0
pair_coeff * * eam/alloy nialhjea Ni Ni NULL
pair_coeff 3 3 lj/cut/coul/cut 1.0 1.0
pair_coeff 1*2 3 lj/cut 0.8 1.3
```

As an example of using the same pair style multiple times, consider a simulation with 2 atom types. Type 1 is Si, type 2 is C. The following commands would model the Si atoms with Tersoff, the C atoms with Tersoff, and the cross-interactions with Lennard-Jones:

```
pair_style hybrid lj/cut 2.5 tersoff tersoff
pair_coeff * * tersoff 1 Si.tersoff Si NULL
pair_coeff * * tersoff 2 C.tersoff NULL C
pair_coeff 1 2 lj/cut 1.0 1.5
```

It is not recommended to read pair coefficients for a hybrid style from a “Pair Coeffs” or “PairIJ Coeffs” section of a data file via the `read_data` command, since those sections expect a fixed number of lines, either one line per atom type or one line pair pair of atom types, respectively. When reading from a data file, the lines of the “Pair Coeffs” and “PairIJ Coeffs” are changed in the same way as the `pair_coeff` command, i.e. the name of the pair style to which the parameters apply must follow the atom type (or atom types), e.g.

```
Pair Coeffs
1 lj/cut/coul/cut 1.0 1.0
...

PairIJ Coeffs
1 1 lj/cut/coul/cut 1.0 1.0
...
```

Note that the `pair_coeff` command for some potentials such as `pair_style eam/alloy` includes a mapping specification of elements to all atom types, which in the hybrid case, can include atom types not assigned to the `eam/alloy` potential. The NULL keyword is used by many such potentials (`eam/alloy`, Tersoff, AIREBO, etc), to denote an atom type that will be assigned to a different sub-style.

For the *hybrid* style, each atom type pair I,J is assigned to exactly one sub-style. Just as with a simulation using a single pair style, if you specify the same atom type pair in a second `pair_coeff` command, the previous assignment will be

overwritten.

For the *hybrid/overlay* and *hybrid/scaled* styles, each atom type pair I,J can be assigned to one or more sub-styles. If you specify the same atom type pair in a second `pair_coeff` command with a new sub-style, then the second sub-style is added to the list of potentials that will be calculated for two interacting atoms of those types. If you specify the same atom type pair in a second `pair_coeff` command with a sub-style that has already been defined for that pair of atoms, then the new pair coefficients simply override the previous ones, as in the normal usage of the `pair_coeff` command. E.g. these two sets of commands are the same:

```
pair_style lj/cut 2.5
pair_coeff * * 1.0 1.0
pair_coeff 2 2 1.5 0.8

pair_style hybrid/overlay lj/cut 2.5
pair_coeff * * lj/cut 1.0 1.0
pair_coeff 2 2 lj/cut 1.5 0.8
```

Coefficients must be defined for each pair of atoms types via the `pair_coeff` command as described above, or in the “Pair Coeffs” or “PairIJ Coeffs” section of the data file read by the `read_data` command, or by mixing as described below.

For all of the *hybrid*, *hybrid/overlay*, and *hybrid/scaled* styles, every atom type pair I,J (where  $I \leq J$ ) must be assigned to at least one sub-style via the `pair_coeff` command as in the examples above, or in the data file read by the `read_data`, or by mixing as described below. Also all sub-styles must be used at least once in a `pair_coeff` command.

**Warning:** With hybrid pair styles the use of mixing to generate pair coefficients is significantly limited compared to the individual pair styles. LAMMPS **never** performs mixing of parameters from different sub-styles, **even** if they use the same type of coefficients, e.g. contain a Lennard-Jones potential variant. Those parameters must be provided explicitly. Also for *hybrid/overlay* and *hybrid/scaled* mixing is **only** performed for pairs of atom types for which only a single pair style is assigned.

Thus it is strongly recommended to provide all mixed terms explicitly. For non-hybrid styles those could be generated and written out using the `write_coeff_command` and then edited as needed to comply with the requirements for hybrid styles as explained above.

If you want there to be no interactions between a particular pair of atom types, you have 3 choices. You can assign the pair of atom types to some sub-style and use the `neigh_modify exclude type` command. You can assign it to some sub-style and set the coefficients so that there is effectively no interaction (e.g.  $\epsilon = 0.0$  in a LJ potential). Or, for *hybrid*, *hybrid/overlay*, or *hybrid/scaled* simulations, you can use this form of the `pair_coeff` command in your input script or the “PairIJ Coeffs” section of your data file:

```
pair_coeff 2 3 none
```

or this form in the “Pair Coeffs” section of the data file:

```
3 none
```

If an assignment to *none* is made in a simulation with the *hybrid/overlay* or *hybrid/scaled* pair style, it wipes out all previous assignments of that pair of atom types to sub-styles.

Note that you may need to use an `atom_style hybrid` command in your input script, if atoms in the simulation will need attributes from several atom styles, due to using multiple pair styles with different requirements.

Different force fields (e.g. CHARMM vs. AMBER) may have different rules for applying exclusions or weights that change the strength of pairwise non-bonded interactions between pairs of atoms that are also 1-2, 1-3, and 1-4 neighbors



in the molecular bond topology. This is normally a global setting defined the *special\_bonds* command. However, different weights can be assigned to different hybrid sub-styles via the *pair\_modify special* command. This allows multiple force fields to be used in a model of a hybrid system, however, there is no consistent approach to determine parameters automatically for the interactions **between** atoms of the two force fields, thus this approach this is only recommended when particles described by the different force fields do not mix.

Here is an example for combining CHARMM and AMBER: The global *amber* setting sets the 1-4 interactions to non-zero scaling factors and then overrides them with 0.0 only for CHARMM:

```
special_bonds amber
pair_style hybrid lj/charmm/coul/long 8.0 10.0 lj/cut/coul/long 10.0
pair_modify pair lj/charmm/coul/long special lj/coul 0.0 0.0 0.0
```

This input achieves the same effect:

```
special_bonds 0.0 0.0 0.1
pair_style hybrid lj/charmm/coul/long 8.0 10.0 lj/cut/coul/long 10.0
pair_modify pair lj/cut/coul/long special lj 0.0 0.0 0.5
pair_modify pair lj/cut/coul/long special coul 0.0 0.0 0.83333333
pair_modify pair lj/charmm/coul/long special lj/coul 0.0 0.0 0.0
```

Here is an example for combining Tersoff with OPLS/AA based on a data file that defines bonds for all atoms where - for the Tersoff part of the system - the force constants for the bonded interactions have been set to 0. Note the global settings are effectively *lj/coul 0.0 0.0 0.5* as required for OPLS/AA:

```
special_bonds lj/coul 1e-20 1e-20 0.5
pair_style hybrid tersoff lj/cut/coul/long 12.0
pair_modify pair tersoff special lj/coul 1.0 1.0 1.0
```

For use with the various *compute \*/tally* computes, the *pair\_modify compute/tally* command can be used to selectively turn off processing of the compute tally styles, for example, if those pair styles (e.g. many-body styles) do not support this feature.

See the *pair\_modify* page for details on the specific syntax, requirements and restrictions.

The potential energy contribution to the overall system due to an individual sub-style can be accessed and output via the *compute pair* command. Note that in the case of pair style *hybrid/scaled* this is the **unscaled** potential energy of the selected sub-style.

**Note:** Several of the potentials defined via the *pair\_style* command in LAMMPS are really many-body potentials, such as Tersoff, AIREBO, MEAM, ReaxFF, etc. The way to think about using these potentials in a hybrid setting is as follows.

A subset of atom types is assigned to the many-body potential with a single *pair\_coeff* command, using “\* \*” to include all types and the NULL keywords described above to exclude specific types not assigned to that potential. If types 1,3,4 were assigned in that way (but not type 2), this means that all many-body interactions between all atoms of types 1,3,4 will be computed by that potential. Pair\_style hybrid allows interactions between type pairs 2-2, 1-2, 2-3, 2-4 to be specified for computation by other pair styles. You could even add a second interaction for 1-1 to be computed by another pair style, assuming pair\_style hybrid/overlay is used.

But you should not, as a general rule, attempt to exclude the many-body interactions for some subset of the type pairs within the set of 1,3,4 interactions, e.g. exclude 1-1 or 1-3 interactions. That is not conceptually well-defined for many-body interactions, since the potential will typically calculate energies and forces for small groups of atoms, e.g. 3 or 4 atoms, using the neighbor lists of the atoms to find the additional atoms in the group.

However, you can still use the `pair_coeff none` setting or the `neigh_modify exclude` command to exclude certain type pairs from the neighbor list that will be passed to a many-body sub-style. This will alter the calculations made by a many-body potential beyond the specific pairs, since it builds its list of 3-body, 4-body, etc interactions from the pair lists. You will need to think **carefully** as to whether excluding such pairs produces a physically meaningful result for your model.

For example, imagine you have two atom types in your model, type 1 for atoms in one surface, and type 2 for atoms in the other, and you wish to use a Tersoff potential to compute interactions within each surface, but not between the surfaces. Then either of these two command sequences would implement that model:

```
pair_style hybrid tersoff
pair_coeff * * tersoff SiC.tersoff C C
pair_coeff 1 2 none

pair_style tersoff
pair_coeff * * SiC.tersoff C C
neigh_modify exclude type 1 2
```

Either way, only neighbor lists with 1-1 or 2-2 interactions would be passed to the Tersoff potential, which means it would compute no 3-body interactions containing both type 1 and 2 atoms.

Here is another example to use 2 many-body potentials together in an overlapping manner using hybrid/overlay. Imagine you have CNT (C atoms) on a Si surface. You want to use Tersoff for Si/Si and Si/C interactions, and AIREBO for C/C interactions. Si atoms are type 1; C atoms are type 2. Something like this will work:

```
pair_style hybrid/overlay tersoff airebo 3.0
pair_coeff * * tersoff SiC.tersoff.custom Si C
pair_coeff * * airebo CH.airebo NULL C
```

Note that to prevent the Tersoff potential from computing C/C interactions, you would need to **modify** the `SiC.tersoff` potential file to turn off C/C interaction, i.e. by setting the appropriate coefficients to 0.0.

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix` *command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

**Note:** Since the `hybrid`, `hybrid/overlay`, `hybrid/scaled` styles delegate computation to the individual sub-styles, the suffix versions of the `hybrid` and `hybrid/overlay` styles are used to propagate the corresponding suffix to all sub-styles, if those versions exist. Otherwise the non-accelerated version will be used. The individual accelerated sub-styles are part of the GPU, KOKKOS, INTEL, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

---

### 19.129.4 Mixing, shift, table, tail correction, restart, rRESPA info

Any pair potential settings made via the *pair\_modify* command are passed along to all sub-styles of the hybrid potential.

For atom type pairs I,J and I != J, if the sub-style assigned to I,I and J,J is the same, and if the sub-style allows for mixing, then the coefficients for I,J can be mixed. This means you do not have to specify a *pair\_coeff* command for I,J since the I,J type pair will be assigned automatically to the sub-style defined for both I,I and J,J and its coefficients generated by the mixing rule used by that sub-style. For the *hybrid/overlay* and *hybrid/scaled* style, there is an additional requirement that both the I,I and J,J pairs are assigned to a single sub-style. If this requirement is not met, no I,J coeffs will be generated, even if the sub-styles support mixing, and I,J pair coefficients must be explicitly defined.

See the *pair\_modify* command for details of mixing rules. See the See the page for the sub-style to see if allows for mixing.

The hybrid pair styles supports the *pair\_modify* shift, table, and tail options for an I,J pair interaction, if the associated sub-style supports it.

For the hybrid pair styles, the list of sub-styles and their respective settings are written to *binary restart files*, so a *pair\_style* command does not need to be specified in an input script that reads a restart file. However, the coefficient information is not stored in the restart file. Thus, *pair\_coeff* commands need to be re-specified in the restart input script. For pair style *hybrid/scaled* also the names of any variables used as scale factors are restored, but not the variables themselves, so those may need to be redefined when continuing from a restart.

These pair styles support the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command, if their sub-styles do.

### 19.129.5 Restrictions

When using a long-range Coulombic solver (via the *kpace\_style* command) with a hybrid pair\_style, one or more sub-styles will be of the “long” variety, e.g. *lj/cut/coul/long* or *buck/coul/long*. You must insure that the short-range Coulombic cutoff used by each of these long pair styles is the same or else LAMMPS will generate an error.

Pair style *hybrid/scaled* currently only works for non-accelerated pair styles and pair styles from the OPT package.

When using pair styles from the GPU package they must not be listed multiple times. LAMMPS will detect this and abort.

### 19.129.6 Related commands

*pair\_coeff*

### 19.129.7 Default

none

## 19.130 pair\_style ilp/graphene/hbn command

Accelerator Variant: *ilp/graphene/hbn/opt*

### 19.130.1 Syntax

```
pair_style [hybrid/overlay ...] ilp/graphene/hbn cutoff tap_flag
```

- cutoff = global cutoff (distance units)
- tap\_flag = 0/1 to turn off/on the taper function

### 19.130.2 Examples

```
pair_style hybrid/overlay ilp/graphene/hbn 16.0 1
pair_coeff * * ilp/graphene/hbn BNCH.ILP B N C

pair_style hybrid/overlay rebo tersoff ilp/graphene/hbn 16.0 coul/shield 16.0
pair_coeff * * rebo CH.rebo NULL NULL C
pair_coeff * * tersoff BNC.tersoff B N NULL
pair_coeff * * ilp/graphene/hbn BNCH.ILP B N C
pair_coeff 1 1 coul/shield 0.70
pair_coeff 1 2 coul/shield 0.695
pair_coeff 2 2 coul/shield 0.69
```

### 19.130.3 Description

The *ilp/graphene/hbn* style computes the registry-dependent interlayer potential (ILP) potential as described in (*Leven1*), (*Leven2*) and (*Maaravi*). The normals are calculated in the way as described in (*Kolmogorov*).

$$E = \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij}$$

$$V_{ij} = \text{Tap}(r_{ij}) \left\{ e^{-\alpha(r_{ij}/\beta-1)} [\epsilon + f(\rho_{ij}) + f(\rho_{ji})] - \frac{1}{1 + e^{-d[(r_{ij}/(s_R \cdot r_{eff})) - 1]}} \cdot \frac{C_6}{r_{ij}^6} \right\}$$

$$\rho_{ij}^2 = r_{ij}^2 - (\mathbf{r}_{ij} \cdot \mathbf{n}_i)^2$$

$$\rho_{ji}^2 = r_{ij}^2 - (\mathbf{r}_{ij} \cdot \mathbf{n}_j)^2$$

$$f(\rho) = C e^{-(\rho/\delta)^2}$$

$$\text{Tap}(r_{ij}) = 20 \left( \frac{r_{ij}}{R_{cut}} \right)^7 - 70 \left( \frac{r_{ij}}{R_{cut}} \right)^6 + 84 \left( \frac{r_{ij}}{R_{cut}} \right)^5 - 35 \left( \frac{r_{ij}}{R_{cut}} \right)^4 + 1$$

Where  $\text{Tap}(r_{ij})$  is the taper function which provides a continuous cutoff (up to third derivative) for interatomic separations larger than  $r_c$  (*Maaravi*). The definitions of each parameter in the above equation can be found in (*Leven1*) and (*Maaravi*).

It is important to include all the pairs to build the neighbor list for calculating the normals.

**Note:** This potential (ILP) is intended for interlayer interactions between two different layers of graphene, hexagonal boron nitride (h-BN) and their hetero-junction. To perform a realistic simulation, this potential must be used in combination with intralayer potential, such as *AIREBO* or *Tersoff* potential. To keep the intralayer properties unaffected, the

interlayer interaction within the same layers should be avoided. Hence, each atom has to have a layer identifier such that atoms residing on the same layer interact via the appropriate intralayer potential and atoms residing on different layers interact via the ILP. Here, the molecule id is chosen as the layer identifier, thus a data file with the “full” atom style is required to use this potential.

The parameter file (e.g. BNCH.ILP), is intended for use with *metal units*, with energies in meV. Two additional parameters, *S*, and *rcut* are included in the parameter file. *S* is designed to facilitate scaling of energies. *rcut* is designed to build the neighbor list for calculating the normals for each atom pair.

**Note:** The parameters presented in the parameter file (e.g. BNCH.ILP), are fitted with taper function by setting the cutoff equal to 16.0 Angstrom. Using different cutoff or taper function should be careful. The parameters for atoms pairs between Boron and Nitrogen are fitted with a screened Coulomb interaction *coul/shield*. Therefore, to simulated the properties of h-BN correctly, this potential must be used in combination with the pair style *coul/shield*.

**Note:** Four new sets of parameters of ILP for 2D layered Materials with bilayer and bulk configurations are presented in (*Ouyang1*) and (*Ouyang2*), respectively. These parameters provide a good description in both short- and long-range interaction regimes. While the old ILP parameters published in (*Leven2*) and (*Maaravi*) are only suitable for long-range interaction regime. This feature is essential for simulations in high pressure regime (i.e., the interlayer distance is smaller than the equilibrium distance). The benchmark tests and comparison of these parameters can be found in (*Ouyang1*) and (*Ouyang2*).

This potential must be used in combination with hybrid/overlay. Other interactions can be set to zero using pair\_style *none*.

This pair style tallies a breakdown of the total interlayer potential energy into sub-categories, which can be accessed via the *compute pair* command as a vector of values of length 2. The 2 values correspond to the following sub-categories:

1.  $E_{vdW}$  = vdW (attractive) energy
2.  $E_{Rep}$  = Repulsive energy

To print these quantities to the log file (with descriptive column headings) the following commands could be included in an input script:

```
compute 0 all pair ilp/graphene/hbn
variable Evdw equal c_0[1]
variable Erep equal c_0[2]
thermo_style custom step temp epair v_Erep v_Evdw
```

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

### 19.130.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the `pair_modify` mix, shift, table, and tail options.

This pair style does not write their information to binary restart files, since it is stored in potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

### 19.130.5 Restrictions

This pair style is part of the INTERLAYER package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the newton setting to be *on* for pair interactions.

The BNCH.ILP potential file provided with LAMMPS (see the potentials directory) are parameterized for *metal* units. You can use this potential with any LAMMPS units, but you would need to create your BNCH.ILP potential file with coefficients listed in the appropriate units, if your simulation does not use *metal* units.

### 19.130.6 Related commands

*pair\_coeff*, *pair\_none*, *pair\_style hybrid/overlay*, *pair\_style drip*, *pair\_style ilp\_tmd*, *pair\_style saip\_metal*, *pair\_style pair\_kolmogorov\_crespi\_z*, *pair\_style pair\_kolmogorov\_crespi\_full*, *pair\_style pair\_lebedeva\_z*, *pair\_style pair\_coul\_shield*.

### 19.130.7 Default

`tap_flag = 1`

---

**(Leven1)** I. Leven, I. Azuri, L. Kronik and O. Hod, J. Chem. Phys. 140, 104106 (2014).

**(Leven2)** I. Leven et al, J. Chem.Theory Comput. 12, 2896-905 (2016).

**(Maaravi)** T. Maaravi et al, J. Phys. Chem. C 121, 22826-22835 (2017).

**(Kolmogorov)** A. N. Kolmogorov, V. H. Crespi, Phys. Rev. B 71, 235415 (2005).

**(Ouyang1)** W. Ouyang, D. Mandelli, M. Urbakh and O. Hod, Nano Lett. 18, 6009-6016 (2018).

**(Ouyang2)** W. Ouyang et al., J. Chem. Theory Comput. 16(1), 666-676 (2020).

## 19.131 `pair_style ilp/tmd` command

Accelerator Variant: *ilp/tmd/opt*

### 19.131.1 Syntax

```
pair_style [hybrid/overlay ...] ilp/tmd cutoff tap_flag
```

- cutoff = global cutoff (distance units)
- tap\_flag = 0/1 to turn off/on the taper function

### 19.131.2 Examples

```
pair_style hybrid/overlay ilp/tmd 16.0 1
pair_coeff * * ilp/tmd TMD.ILP Mo S S

pair_style hybrid/overlay sw/mod sw/mod ilp/tmd 16.0
pair_coeff * * sw/mod 1 tmd.sw.mod Mo S S NULL NULL NULL
pair_coeff * * sw/mod 2 tmd.sw.mod NULL NULL NULL Mo S S
pair_coeff * * ilp/tmd TMD.ILP Mo S S Mo S S
```

### 19.131.3 Description

The *ilp/tmd* style computes the registry-dependent interlayer potential (ILP) potential for transition metal dichalcogenides (TMD) as described in (Ouyang7).

$$E = \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij}$$

$$V_{ij} = \text{Tap}(r_{ij}) \left\{ e^{-\alpha(r_{ij}/\beta-1)} [\epsilon + f(\rho_{ij}) + f(\rho_{ji})] - \frac{1}{1 + e^{-d[(r_{ij}/(s_R \cdot r^{eff})-1)]}} \cdot \frac{C_6}{r_{ij}^6} \right\}$$

$$\rho_{ij}^2 = r_{ij}^2 - (\mathbf{r}_{ij} \cdot \mathbf{n}_i)^2$$

$$\rho_{ji}^2 = r_{ij}^2 - (\mathbf{r}_{ij} \cdot \mathbf{n}_j)^2$$

$$f(\rho) = C e^{-(\rho/\delta)^2}$$

$$\text{Tap}(r_{ij}) = 20 \left( \frac{r_{ij}}{R_{cut}} \right)^7 - 70 \left( \frac{r_{ij}}{R_{cut}} \right)^6 + 84 \left( \frac{r_{ij}}{R_{cut}} \right)^5 - 35 \left( \frac{r_{ij}}{R_{cut}} \right)^4 + 1$$

Where  $\text{Tap}(r_{ij})$  is the taper function which provides a continuous cutoff (up to third derivative) for interatomic separations larger than  $r_c$  *pair\_style ilp\_graphene\_hbn*.

It is important to include all the pairs to build the neighbor list for calculating the normals.

---

**Note:** Since each MX<sub>2</sub> (M = Mo, W and X = S, Se Te) layer contains two sub-layers of X atoms and one sub-layer of M atoms, the definition of the normal vectors used for graphene and h-BN is no longer valid for TMDs. In (Ouyang7), a new definition is proposed, where for each atom *i*, its six nearest neighboring atoms belonging to the same sub-layer are chosen to define the normal vector *{bf n}\_i*.

---

The parameter file (e.g. TMD.ILP), is intended for use with *metal units*, with energies in meV. Two additional parameters, *S*, and *rcut* are included in the parameter file. *S* is designed to facilitate scaling of energies. *rcut* is designed to build the neighbor list for calculating the normals for each atom pair.

---

**Note:** The parameters presented in the parameter file (e.g. TMD.ILP), are fitted with taper function by setting the cutoff equal to 16.0 Angstrom. Using different cutoff or taper function should be careful. These parameters provide a good

description in both short- and long-range interaction regimes. This feature is essential for simulations in high pressure regime (i.e., the interlayer distance is smaller than the equilibrium distance). The benchmark tests and comparison of these parameters can be found in ([Ouyang7](#)).

---

This potential must be used in combination with hybrid/overlay. Other interactions can be set to zero using `pair_style none`.

This pair style tallies a breakdown of the total interlayer potential energy into sub-categories, which can be accessed via the `compute pair` command as a vector of values of length 2. The 2 values correspond to the following sub-categories:

1.  $E_{vdW}$  = vdW (attractive) energy
2.  $E_{Rep}$  = Repulsive energy

To print these quantities to the log file (with descriptive column headings) the following commands could be included in an input script:

```
compute 0 all pair ilp/tmd
variable Evdw equal c_0[1]
variable Erep equal c_0[2]
thermo_style custom step temp epair v_Erep v_Evdw
```

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix` *command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.131.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the `pair_modify` `mix`, `shift`, `table`, and `tail` options.

This pair style does not write their information to binary restart files, since it is stored in potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

#### 19.131.5 Restrictions

This pair style is part of the INTERLAYER package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This pair style requires the `newton` setting to be `on` for pair interactions.

The TMD.ILP potential file provided with LAMMPS (see the potentials directory) are parameterized for *metal* units. You can use this potential with any LAMMPS units, but you would need to create your BNCH.ILP potential file with coefficients listed in the appropriate units, if your simulation does not use *metal* units.



### 19.131.6 Related commands

*pair\_coeff*, *pair\_none*, *pair\_style hybrid/overlay*, *pair\_style drip*, *pair\_style saip\_metal*, *pair\_style ilp\_graphene\_hbn*, *pair\_style pair\_kolmogorov\_crespi\_z*, *pair\_style pair\_kolmogorov\_crespi\_full*, *pair\_style pair\_lebedeva\_z*, *pair\_style pair\_coul\_shield*.

### 19.131.7 Default

tap\_flag = 1

(Ouyang<sup>7</sup>) W. Ouyang, et al., J. Chem. Theory Comput. 17, 7237 (2021).

## 19.132 pair\_style kim command

### 19.132.1 Syntax

```
pair_style kim model
```

model = name of a KIM model (the KIM ID for models archived in OpenKIM)

### 19.132.2 Examples

```
pair_style kim SW_StillingerWeber_1985_Si__MO_405512056662_005
pair_coeff * * Si
```

### 19.132.3 Description

This pair style is a wrapper on the [Open Knowledgebase of Interatomic Models \(OpenKIM\)](#) repository of interatomic potentials to enable their use in LAMMPS scripts.

The preferred interface for using interatomic models archived in OpenKIM is the *kim command* interface. That interface supports both “KIM Portable Models” (PMs) that conform to the KIM API Portable Model Interface (PMI) and can be used by any simulation code that conforms to the KIM API/PMI, and “KIM Simulator Models” (SMs) that are natively implemented within a single simulation code (like LAMMPS) and can only be used with it. The *pair\_style kim* command is limited to KIM PMs. It is used by the *kim command* interface as needed.

**Note:** Since *pair\_style kim* is called by *kim interactions* as needed, it is not recommended to be directly used in input scripts.

The argument *model* is the name of the KIM PM. For potentials archived in OpenKIM this is the extended KIM ID (see *kim command* for details). LAMMPS can invoke any KIM PM, however there can be incompatibilities (for example due to unit matching issues). In the event of an incompatibility, the code will terminate with an error message. Check both the LAMMPS and KIM log files for details.

Only a single *pair\_coeff* command is used with the *kim* style, which specifies the mapping of LAMMPS atom types to the species supported by the KIM PM. This is done by specifying *N* additional arguments after the *\*\** in the *pair\_coeff* command, where *N* is the number of LAMMPS atom types:

- N element names = mapping of KIM elements to atom types

For example, consider a KIM PM that supports Si and C species. If the LAMMPS simulation has four atom types, where the first three are Si, and the fourth is C, the following *pair\_coeff* command would be used:

```
pair_coeff * * Si Si Si C
```

The first two arguments must be *\*\** so as to span all LAMMPS atom types. The first three Si arguments map LAMMPS atom types 1, 2, and 3 to Si as defined within KIM PM. The final C argument maps LAMMPS atom type 4 to C.

---

In addition to the usual LAMMPS error messages, the KIM library itself may generate errors, which should be printed to the screen. In this case it is also useful to check the *kim.log* file for additional error information. The file *kim.log* should be generated in the same directory where LAMMPS is running.

To download, build, and install the KIM library on your system, see the *lib/kim/README* file. Once you have done this and built LAMMPS with the KIM package installed you can run the example input scripts in *examples/kim*.

---

#### 19.132.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* mix, shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since KIM stores the potential parameters. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.132.5 Restrictions

This pair style is part of the KIM package. See details on restrictions in *kim command*.

This current version of *pair\_style kim* is compatible with the *kim-api* package version 2.0.0 and higher.

#### 19.132.6 Related commands

*pair\_coeff*, *kim command*

#### 19.132.7 Default

none

## 19.133 pair\_style kolmogorov/crespi/full command

### 19.133.1 Syntax

```
pair_style hybrid/overlay kolmogorov/crespi/full cutoff tap_flag
```

- cutoff = global cutoff (distance units)
- tap\_flag = 0/1 to turn off/on the taper function

### 19.133.2 Examples

```
pair_style hybrid/overlay kolmogorov/crespi/full 20.0 0
pair_coeff * * none
pair_coeff * * kolmogorov/crespi/full CH.KC C C

pair_style hybrid/overlay rebo kolmogorov/crespi/full 16.0 1
pair_coeff * * rebo CH.rebo C H
pair_coeff * * kolmogorov/crespi/full CH_taper.KC C H
```

### 19.133.3 Description

The *kolmogorov/crespi/full* style computes the Kolmogorov-Crespi (KC) interaction potential as described in (*Kolmogorov*). No simplification is made,

$$E = \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij}$$

$$V_{ij} = e^{-\lambda(r_{ij}-z_0)} [C + f(\rho_{ij}) + f(\rho_{ji})] - A \left( \frac{r_{ij}}{z_0} \right)^{-6}$$

$$\rho_{ij}^2 = r_{ij}^2 - (\mathbf{r}_{ij} \cdot \mathbf{n}_i)^2$$

$$\rho_{ji}^2 = r_{ij}^2 - (\mathbf{r}_{ij} \cdot \mathbf{n}_j)^2$$

$$f(\rho) = e^{-(\rho/\delta)^2} \sum_{n=0}^2 C_{2n} (\rho/\delta)^{2n}$$

It is important to have a sufficiently large cutoff to ensure smooth forces and to include all the pairs to build the neighbor list for calculating the normals. Energies are shifted so that they go continuously to zero at the cutoff assuming that the exponential part of  $V_{ij}$  (first term) decays sufficiently fast. This shift is achieved by the last term in the equation for  $V_{ij}$  above. This is essential only when the taper function is turned off. The formula of taper function can be found in pair style *ilp/graphene/hbn*.

**Note:** This potential (ILP) is intended for interlayer interactions between two different layers of graphene. To perform a realistic simulation, this potential must be used in combination with intralayer potential, such as *AIREBO* or *Tersoff* potential. To keep the intralayer properties unaffected, the interlayer interaction within the same layers should be avoided. Hence, each atom has to have a layer identifier such that atoms residing on the same layer interact via the appropriate intralayer potential and atoms residing on different layers interact via the ILP. Here, the molecule id is chosen as the layer identifier, thus a data file with the “full” atom style is required to use this potential.

The parameter file (e.g. CH.KC), is intended for use with *metal units*, with energies in meV. Two additional parameters, *S*, and *rcut* are included in the parameter file. *S* is designed to facilitate scaling of energies. *rcut* is designed to build the neighbor list for calculating the normals for each atom pair.

---

**Note:** Two new sets of parameters of KC potential for hydrocarbons, CH.KC (without the taper function) and CH\_taper.KC (with the taper function) are presented in (*Ouyang1*). The energy for the KC potential with the taper function goes continuously to zero at the cutoff. The parameters in both CH.KC and CH\_taper.KC provide a good description in both short- and long-range interaction regimes. While the original parameters (CC.KC) published in (*Kolmogorov*) are only suitable for long-range interaction regime. This feature is essential for simulations in high pressure regime (i.e., the interlayer distance is smaller than the equilibrium distance). The benchmark tests and comparison of these parameters can be found in (*Ouyang1*) and (*Ouyang2*).

---

This potential must be used in combination with hybrid/overlay. Other interactions can be set to zero using `pair_style none`.

This pair style tallies a breakdown of the total interlayer potential energy into sub-categories, which can be accessed via the `compute pair` command as a vector of values of length 2. The 2 values correspond to the following sub-categories:

1.  $E_{vdW}$  = vdW (attractive) energy
2.  $E_{Rep}$  = Repulsive energy

To print these quantities to the log file (with descriptive column headings) the following commands could be included in an input script:

```
compute 0 all pair kolmogorov/crespi/full
variable Evdw equal c_0[1]
variable Erep equal c_0[2]
thermo_style custom step temp epair v_Erep v_Evdw
```

---

#### 19.133.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the `pair_modify` mix, shift, table, and tail options.

This pair style does not write their information to binary restart files, since it is stored in potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

#### 19.133.5 Restrictions

This pair style is part of the INTERLAYER package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the newton setting to be *on* for pair interactions.

The CH.KC potential file provided with LAMMPS (see the potentials folder) is parameterized for metal units. You can use this pair style with any LAMMPS units, but you would need to create your own custom CC.KC potential file with all coefficients converted to the appropriate units.

### 19.133.6 Related commands

*pair\_coeff*, *pair\_none*, *pair\_style hybrid/overlay*, *pair\_style drip*, *pair\_style pair\_lebedeva\_z*, *pair\_style kolmogorov/crespi/z*, *pair\_style ilp/graphene/hbn*.

### 19.133.7 Default

tap\_flag = 0

**(Kolmogorov)** A. N. Kolmogorov, V. H. Crespi, Phys. Rev. B 71, 235415 (2005)

**(Ouyang1)** W. Ouyang, D. Mandelli, M. Urbakh and O. Hod, Nano Lett. 18, 6009-6016 (2018).

**(Ouyang2)** W. Ouyang et al., J. Chem. Theory Comput. 16(1), 666-676 (2020).

## 19.134 pair\_style kolmogorov/crespi/z command

### 19.134.1 Syntax

```
pair_style [hybrid/overlay ...] kolmogorov/crespi/z cutoff
```

### 19.134.2 Examples

```
pair_style hybrid/overlay kolmogorov/crespi/z 20.0
pair_coeff * * none
pair_coeff 1 2 kolmogorov/crespi/z CC.KC C C

pair_style hybrid/overlay rebo kolmogorov/crespi/z 14.0
pair_coeff * * rebo CH.rebo C C
pair_coeff 1 2 kolmogorov/crespi/z CC.KC C C
```

### 19.134.3 Description

The *kolmogorov/crespi/z* style computes the Kolmogorov-Crespi interaction potential as described in (Kolmogorov). An important simplification is made, which is to take all normals along the z-axis.

$$E = \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij}$$

$$V_{ij} = e^{-\lambda(r_{ij}-z_0)} [C + f(\rho_{ij}) + f(\rho_{ji})] - A \left( \frac{r_{ij}}{z_0} \right)^{-6} + A \left( \frac{\text{cutoff}}{z_0} \right)^{-6}$$

$$\rho_{ij}^2 = \rho_{ji}^2 = x_{ij}^2 + y_{ij}^2 \quad (\mathbf{n}_i \equiv \hat{\mathbf{z}})$$

$$f(\rho) = e^{-(\rho/\delta)^2} \sum_{n=0}^2 C_{2n} (\rho/\delta)^{2n}$$

It is important to have a sufficiently large cutoff to ensure smooth forces. Energies are shifted so that they go continuously to zero at the cutoff assuming that the exponential part of  $V_{ij}$  (first term) decays sufficiently fast. This shift is achieved by the last term in the equation for  $V_{ij}$  above.

This potential is intended for interactions between two layers of graphene. Therefore, to avoid interaction between layers in multi-layered materials, each layer should have a separate atom type and interactions should only be computed between atom types of neighboring layers.

The parameter file (e.g. CC.KC), is intended for use with metal *units*, with energies in meV. An additional parameter, *S*, is available to facilitate scaling of energies in accordance with (*vanWijk*).

This potential must be used in combination with hybrid/overlay. Other interactions can be set to zero using *pair\_style none*.

#### 19.134.4 Restrictions

This fix is part of the INTERLAYER package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 19.134.5 Related commands

*pair\_coeff*, *pair\_none*, *pair\_style hybrid/overlay*, *pair\_style drip*, *pair\_style ilp/graphene/hbn*. *pair\_style kolmogorov/crespi/full*, *pair\_style lebedeva/z*

#### 19.134.6 Default

none

---

(**Kolmogorov**) A. N. Kolmogorov, V. H. Crespi, Phys. Rev. B 71, 235415 (2005)

(**vanWijk**) M. M. van Wijk, A. Schuring, M. I. Katsnelson, and A. Fasolino, Physical Review Letters, 113, 135504 (2014)

### 19.135 *pair\_style lcbop* command

#### 19.135.1 Syntax

```
pair_style lcbop
```

#### 19.135.2 Examples

```
pair_style lcbop  
pair_coeff * * ../potentials/C.lcbop C
```

### 19.135.3 Description

The *lcbop* pair style computes the long-range bond-order potential for carbon (LCBOP) of (*Los and Fasolino*). See section II in that paper for the analytic equations associated with the potential.

Only a single `pair_coeff` command is used with the *lcbop* style which specifies an LCBOP potential file with parameters for specific elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of LCBOP elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the potential file.

As an example, if your LAMMPS simulation has 4 atom types and you want the first 3 to be C you would use the following `pair_coeff` command:

```
pair_coeff * * C.lcbop C C C NULL
```

The first 2 arguments must be `**` so as to span all LAMMPS atom types. The first C argument maps LAMMPS atom type 1 to the C element in the LCBOP file. If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *lcbop* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

The parameters/coefficients for the LCBOP potential as applied to C are listed in the C.lcbop file to agree with the original (*Los and Fasolino*) paper. Thus the parameters are specific to this potential and the way it was fit, so modifying the file should be done carefully.

### 19.135.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the `pair_modify` mix, shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

This pair style can only be used via the `pair` keyword of the `run_style respa` command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.135.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This pair potential requires the `newton` setting to be “on” for pair interactions.

The C.lcbop potential file provided with LAMMPS (see the potentials directory) is parameterized for *metal units*. You can use the LCBOP potential with any LAMMPS units, but you would need to create your own LCBOP potential file with coefficients listed in the appropriate units if your simulation does not use “metal” units.

## 19.135.6 Related commands

*pair\_airebo*, *pair\_coeff*

## 19.135.7 Default

none

(Los and Fasolino) J. H. Los and A. Fasolino, Phys. Rev. B 68, 024107 (2003).

## 19.136 pair\_style lebedeva/z command

### 19.136.1 Syntax

```
pair_style [hybrid/overlay ...] lebedeva/z cutoff
```

### 19.136.2 Examples

```
pair_style hybrid/overlay lebedeva/z 20.0
pair_coeff * * none
pair_coeff 1 2 lebedeva/z CC.Lebedeva C C

pair_style hybrid/overlay rebo lebedeva/z 14.0
pair_coeff * * rebo CH.rebo C C
pair_coeff 1 2 lebedeva/z CC.Lebedeva C C
```

### 19.136.3 Description

The *lebedeva/z* pair style computes the Lebedeva interaction potential as described in (*Lebedeva1*) and (*Lebedeva2*). An important simplification is made, which is to take all normals along the z-axis.

The Lebedeva potential is intended for the description of the interlayer interaction between graphene layers. To perform a realistic simulation, this potential must be used in combination with an intralayer potential such as *AIREBO* or *Tersoff* facilitated by using pair style *hybrid/overlay*. To keep the intralayer properties unaffected, the interlayer interaction within the same layers should be avoided. This can be achieved by assigning different atom types to atoms of different layers (e.g. 1 and 2 in the examples above).

Other interactions can be set to zero using pair\_style *none*.

$$E = \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij}$$

$$V_{ij} = B e^{-\alpha(r_{ij} - z_0)}$$

$$+ C(1 + D_1 \rho_{ij}^2 + D_2 \rho_{ij}^4) e^{-\lambda_1 \rho_{ij}^2} e^{-\lambda_2 (z_{ij}^2 - z_0^2)}$$

$$- A \left( \frac{z_0}{r_{ij}} \right)^6 + A \left( \frac{z_0}{r_c} \right)^6$$

$$\rho_{ij}^2 = x_{ij}^2 + y_{ij}^2 \quad (\mathbf{n}_i \equiv \hat{\mathbf{z}})$$



It is important to have a sufficiently large cutoff to ensure smooth forces. Energies are shifted so that they go continuously to zero at the cutoff assuming that the exponential part of  $V_{ij}$  (first term) decays sufficiently fast. This shift is achieved by the last term in the equation for  $V_{ij}$  above.

The provided parameter file (CC.Lebedeva) contains two sets of parameters.

- The first set (element name “C”) is suitable for normal conditions and is taken from (*Popov1*)
- The second set (element name “C1”) is suitable for high-pressure conditions and is taken from (*Koziol1*)

Both sets contain an additional parameter,  $S$ , that can be used to facilitate scaling of energies and is set to 1.0 by default.

### 19.136.4 Restrictions

This pair style is part of the INTERLAYER package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.136.5 Related commands

*pair\_coeff*, *pair\_style none*, *pair\_style hybrid/overlay*, *pair\_style drip*, *pair\_style ilp/graphene/hbd*, *pair\_style kolmogorov/crespi/z*, *pair\_style kolmogorov/crespi/full*.

### 19.136.6 Default

none

---

**(Lebedeva1)** I. V. Lebedeva, A. A. Knizhnik, A. M. Popov, Y. E. Lozovik, B. V. Potapkin, Phys. Rev. B, 84, 245437 (2011)

**(Lebedeva2)** I. V. Lebedeva, A. A. Knizhnik, A. M. Popov, Y. E. Lozovik, B. V. Potapkin, Physica E: 44, 949-954 (2012)

**(Popov1)** A.M. Popov, I. V. Lebedeva, A. A. Knizhnik, Y. E. Lozovik and B. V. Potapkin, Chem. Phys. Lett. 536, 82-86 (2012).

**(Koziol1)** Z. Koziol, G. Gawlik and J. Jagielski, Chinese Phys. B 28, 096101 (2019).

## 19.137 pair\_style line/lj command

### 19.137.1 Syntax

```
pair_style line/lj cutoff
```

cutoff = global cutoff for interactions (distance units)

## 19.137.2 Examples

```
pair_style line/lj 3.0
pair_coeff * * 1.0 1.0 1.0 0.8 1.12
pair_coeff 1 2 1.0 2.0 1.0 1.5 1.12 5.0
pair_coeff 1 2 1.0 0.0 1.0 1.0 2.5
```

## 19.137.3 Description

Style *line/lj* treats particles which are line segments as a set of small spherical particles that tile the line segment length as explained below. Interactions between two line segments, each with  $N_1$  and  $N_2$  spherical particles, are calculated as the pairwise sum of  $N_1 * N_2$  Lennard-Jones interactions. Interactions between a line segment with  $N$  spherical particles and a point particle are treated as the pairwise sum of  $N$  Lennard-Jones interactions. See the *pair\_style lj/cut* page for the definition of Lennard-Jones interactions.

The set of non-overlapping spherical sub-particles that represent a line segment are generated in the following manner. Their size is a function of the line segment length and the specified sub-particle size for that particle type. If a line segment has a length  $L$  and is of type  $I$ , then the number of spheres  $N$  that represent the segment is calculated as  $N = L/sizeI$ , rounded up to an integer value. Thus if  $L$  is not evenly divisible by  $sizeI$ ,  $N$  is incremented to include one extra sphere. The centers of the spheres are spaced equally along the line segment. Imagine  $N+1$  equally-space points, which include the 2 end points of the segment. The sphere centers are halfway between each pair of points.

The LJ interaction between 2 spheres on different line segments (or a sphere on a line segment and a point particles) is computed with sub-particle  $\epsilon$ ,  $\sigma$ , and *cutoff* values that are set by the *pair\_coeff* command, as described below. If the distance between the 2 spheres is greater than the sub-particle cutoff, there is no interaction. This means that some pairs of sub-particles on 2 line segments may interact, but others may not.

For purposes of creating the neighbor list for pairs of interacting line segments or lines/point particles, a regular particle-particle cutoff is used, as defined by the *cutoff* setting above in the *pair\_style* command or overridden with an optional argument in the *pair\_coeff* command for a type pair as discussed below. The distance between the centers of 2 line segments, or the center of a line segment and a point particle, must be less than this distance (plus the neighbor skin; see the *neighbor* command), for the pair of particles to be included in the neighbor list.

---

**Note:** This means that a too-short value for the *cutoff* setting can exclude a pair of particles from the neighbor list even if pairs of their sub-particle spheres would interact, based on the sub-particle cutoff specified in the *pair\_coeff* command. E.g. sub-particles at the ends of the line segments that are close to each other. Which may not be what you want, since it means the ends of 2 line segments could pass through each other. It is up to you to specify a *cutoff* setting that is consistent with the length of the line segments you are using and the sub-particle cutoff settings.

---

For style *line/lj*, the following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- *sizeI* (distance units)
- *sizeJ* (distance units)
- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- *subcutoff* (distance units)
- *cutoff* (distance units)

The *sizeI* and *sizeJ* coefficients are the sub-particle sizes for line particles of type  $I$  and type  $J$ . They are used to define the  $N$  sub-particles per segment as described above. These coefficients are actually stored on a per-type basis. Thus

if there are multiple `pair_coeff` commands that involve type I, as either the first or second atom type, you should use consistent values for `sizeI` or `sizeJ` in all of them. If you do not do this, the last value specified for `sizeI` will apply to all segments of type I. If `typeI` or `typeJ` refers to point particles, the corresponding `sizeI` or `sizeJ` is ignored; it can be set to 0.0.

The  $\epsilon$ ,  $\sigma$ , and *subcutoff* coefficients are used to compute LJ interactions between a pair of sub-particles on 2 line segments (of type I and J), or between a sub particle/point particle pair. As discussed above, the *subcutoff* and *cutoff* params are different. The latter is only used for building the neighbor list when the distance between centers of two line segments or one segment and a point particle is calculated.

The *cutoff* coefficient is optional. If not specified, the global cutoff is used.

---

### 19.137.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , coefficients must be specified. No default mixing rules are used.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.137.5 Restrictions

This style is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

Defining particles to be line segments so they participate in line/line or line/particle interactions requires the use the *atom\_style line* command.

### 19.137.6 Related commands

*pair\_coeff*, *pair\_style tri/lj*

### 19.137.7 Default

none

## 19.138 pair\_style list command

### 19.138.1 Syntax

```
pair_style list listfile cutoff keyword
```

- listfile = name of file with list of pairwise interactions
- cutoff = global cutoff (distance units)
- keyword = optional flag *nocheck* or *check* (default is *check*)

## 19.138.2 Examples

```
pair_style list restraints.txt 200.0
pair_coeff * *

pair_style hybrid/overlay lj/cut 1.1225 list pair_list.txt 300.0
pair_coeff * * lj/cut 1.0 1.0
pair_coeff 3* 3* list
```

## 19.138.3 Description

Style *list* computes interactions between explicitly listed pairs of atoms with the option to select functional form and parameters for each individual pair. Because the parameters are set in the list file, the `pair_coeff` command has no parameters (but still needs to be provided). The *check* and *nocheck* keywords enable/disable a test that checks whether all listed bonds were present and computed.

This pair style can be thought of as a hybrid between bonded, non-bonded, and restraint interactions. It will typically be used as an additional interaction within the *hybrid/overlay* pair style. It currently supports three interaction styles: a 12-6 Lennard-Jones, a Morse and a harmonic potential.

The format of the list file is as follows:

- one line per pair of atoms
- empty lines will be ignored
- comment text starts with a '#' character
- line syntax: *ID1 ID2 style coeffs cutoff*

```
ID1 = atom ID of first atom
ID2 = atom ID of second atom
style = style of interaction
coeffs = list of coeffs
cutoff = cutoff for interaction (optional)
```

The cutoff parameter is optional. If not specified, the global cutoff is used.

Here is an example file:

```
# this is a comment

15 259 lj126      1.0 1.0      50.0
15 603 morse     10.0 1.2 2.0   10.0 # and another comment
18 470 harmonic 50.0 1.2      5.0
```

The style *lj126* computes pairwise interactions with the formula

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_c$$

and the coefficients:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)

The style *morse* computes pairwise interactions with the formula

$$E = D_0 \left[ e^{-2\alpha(r-r_0)} - 2e^{-\alpha(r-r_0)} \right] \quad r < r_c$$

and the coefficients:

- $D_0$  (energy units)
- $\alpha$  (1/distance units)
- $r_0$  (distance units)

The style *harmonic* computes pairwise interactions with the formula

$$E = K(r - r_0)^2$$

and the coefficients:

- $K$  (energy units)
- $r_0$  (distance units)

Note that the usual 1/2 factor is included in  $K$ .

---

#### 19.138.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support mixing since all parameters are explicit for each pair.

The *pair\_modify* shift option is supported by this pair style.

The *pair\_modify* table and tail options are not relevant for this pair style.

This pair style does not write its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.138.5 Restrictions

This pair style does not use a neighbor list and instead identifies atoms by their IDs. This has two consequences: 1) The cutoff has to be chosen sufficiently large, so that the second atom of a pair has to be a ghost atom on the same node on which the first atom is local; otherwise the interaction will be skipped. You can use the *check* option to detect, if interactions are missing. 2) Unlike other pair styles in LAMMPS, an atom I will not interact with multiple images of atom J (assuming the images are within the cutoff distance), but only with the nearest image.

This style is part of the MISC package. It is only enabled if LAMMPS is build with that package. See the *Build package* page on for more info.

## 19.138.6 Related commands

*pair\_coeff*, *pair\_style hybrid/overlay*, *pair\_style lj/cut*, *pair\_style morse*, *bond\_style harmonic*

## 19.138.7 Default

none

## 19.139 pair\_style lj/cut command

Accelerator Variants: *lj/cut/gpu*, *lj/cut/intel*, *lj/cut/kk*, *lj/cut/opt*, *lj/cut/omp*

### 19.139.1 Syntax

```
pair_style style args
```

- style = *lj/cut*
- args = list of arguments for a particular style

*lj/cut* args = cutoff

cutoff = global cutoff for Lennard Jones interactions (distance units)

### 19.139.2 Examples

```
pair_style lj/cut 2.5
pair_coeff * * 1 1
pair_coeff 1 1 1 1.1 2.8
```

### 19.139.3 Description

The *lj/cut* styles compute the standard 12/6 Lennard-Jones potential, given by

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_c$$

$r_c$  is the cutoff.

See the *lj/cut/coul* styles to add a Coulombic pairwise interaction and the *lj/cut/tip4p* styles to add the TIP4P water model.

### 19.139.4 Coefficients

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- LJ cutoff (distance units)

Note that  $\sigma$  is defined in the LJ formula as the zero-crossing distance for the potential, not as the energy minimum at  $2^{\frac{1}{6}}\sigma$ .

The last coefficient is optional. If not specified, the global LJ cutoff specified in the *pair\_style* command are used.

---

A version of these styles with a soft core, *lj/cut/soft*, suitable for use in free energy calculations, is part of the FEP package and is documented with the *pair\_style \*/soft* styles.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

### 19.139.5 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the *lj/cut* pair styles can be mixed. The default mix value is *geometric*. See the “*pair\_modify*” command for details.

All of the *lj/cut* pair styles support the *pair\_modify* shift option for the energy of the Lennard-Jones portion of the pair interaction.

All of the *lj/cut* pair styles support the *pair\_modify* tail option for adding a long-range tail correction to the energy and pressure for the Lennard-Jones portion of the pair interaction.

All of the *lj/cut* pair styles write their information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

The *lj/cut* pair styles support the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command, meaning the pairwise forces can be partitioned by distance at different levels of the rRESPA hierarchy. The other styles only support the *pair* keyword of *run\_style respa*. See the *run\_style* command for details.

### 19.139.6 Related commands

- `pair_coeff`
- `pair_style lj/cut/coul/cut`
- `pair_style lj/cut/coul/debye`
- `pair_style lj/cut/coul/dsf`
- `pair_style lj/cut/coul/long`
- `pair_style lj/cut/coul/msm`
- `pair_style lj/cut/coul/wolf`
- `pair_style lj/cut/tip4p/cut`
- `pair_style lj/cut/tip4p/long`

### 19.139.7 Default

none

## 19.140 pair\_style lj96/cut command

Accelerator Variants: `lj96/cut/gpu`, `lj96/cut/omp`

### 19.140.1 Syntax

```
pair_style lj96/cut cutoff
```

- `cutoff` = global cutoff for `lj96/cut` interactions (distance units)

### 19.140.2 Examples

```
pair_style lj96/cut 2.5
pair_coeff * * 1.0 1.0 4.0
pair_coeff 1 1 1.0 1.0
```

### 19.140.3 Description

The `lj96/cut` style compute a 9/6 Lennard-Jones potential, instead of the standard 12/6 potential, given by

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^9 - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_c$$

$r_c$  is the cutoff.

The following coefficients must be defined for each pair of atoms types via the `pair_coeff` command as in the examples above, or in the data file or restart files read by the `read_data` or `read_restart` commands, or by mixing as described below:



- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global LJ cutoff specified in the `pair_style` command is used.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* [command-line switch](#) when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.140.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the lj/cut pair styles can be mixed. The default mix value is *geometric*. See the “`pair_modify`” command for details.

This pair style supports the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style supports the *pair\_modify* tail option for adding a long-range tail correction to the energy and pressure of the pair interaction.

This pair style writes its information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

This pair style supports the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command, meaning the pairwise forces can be partitioned by distance at different levels of the rRESPA hierarchy. See the *run\_style* command for details.

---

#### 19.140.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 19.140.6 Related commands

*pair\_coeff*

## 19.140.7 Default

none

## 19.141 pair\_style lj/cubic command

Accelerator Variants: *lj/cubic/gpu*, *lj/cubic/omp*

### 19.141.1 Syntax

```
pair_style lj/cubic
```

### 19.141.2 Examples

```
pair_style lj/cubic
pair_coeff * * 1.0 0.8908987
```

### 19.141.3 Description

The *lj/cubic* style computes a truncated LJ interaction potential whose energy and force are continuous everywhere. Inside the inflection point the interaction is identical to the standard 12/6 *Lennard-Jones* potential. The LJ function outside the inflection point is replaced with a cubic function of distance. The energy, force, and second derivative are continuous at the inflection point. The cubic coefficient  $A_3$  is chosen so that both energy and force go to zero at the cutoff distance. Outside the cutoff distance the energy and force are zero.

$$\begin{aligned} E &= u_{LJ}(r) & r \leq r_s \\ &= u_{LJ}(r_s) + (r - r_s)u'_{LJ}(r_s) - \frac{1}{6}A_3(r - r_s)^3 & r_s < r \leq r_c \\ &= 0 & r > r_c \end{aligned}$$

The location of the inflection point  $r_s$  is defined by the LJ diameter,  $r_s/\sigma = (26/7)^{1/6}$ . The cutoff distance is defined by  $r_c/r_s = 67/48$  or  $r_c/\sigma = 1.737\dots$ . The analytic expression for the cubic coefficient  $A_3 r_{min}^3/\epsilon = 27.93\dots$  is given in the paper by Holian and Ravelo (*Holian*).

This potential is commonly used to study the shock mechanics of FCC solids, as in Ravelo et al. (*Ravelo*).

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)

Note that  $\sigma$  is defined in the LJ formula as the zero-crossing distance for the potential, not as the energy minimum, which is located at  $r_{min} = 2^{\frac{1}{6}}\sigma$ . In the above example,  $\sigma = 0.8908987$ , so  $r_{min} = 1.0$ .

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* [command-line switch](#) when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.141.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the lj/cut pair styles can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

The lj/cubic pair style does not support the *pair\_modify* shift option, since pair interaction is already smoothed to 0.0 at the cutoff.

The *pair\_modify* table option is not relevant for this pair style.

The lj/cubic pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure, since there are no corrections for a potential that goes to 0.0 at the cutoff.

The lj/cubic pair style writes its information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

The lj/cubic pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.141.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

#### 19.141.6 Related commands

*pair\_coeff*

### 19.141.7 Default

none

---

**(Holian)** Holian and Ravelo, Phys Rev B, 51, 11275 (1995).

**(Ravelo)** Ravelo, Holian, Germann and Lomdahl, Phys Rev B, 70, 014103 (2004).

### 19.142 pair\_style lj/cut/coul/cut command

Accelerator Variants: *lj/cut/coul/cut/gpu*, *lj/cut/coul/cut/kk*, *lj/cut/coul/cut/omp*

### 19.143 pair\_style lj/cut/coul/debye command

Accelerator Variants: *lj/cut/coul/debye/gpu*, *lj/cut/coul/debye/kk*, *lj/cut/coul/debye/omp*

### 19.144 pair\_style lj/cut/coul/dsf command

Accelerator Variants: *lj/cut/coul/dsf/gpu*, *lj/cut/coul/dsf/kk*, *lj/cut/coul/dsf/omp*

### 19.145 pair\_style lj/cut/coul/long command

Accelerator Variants: *lj/cut/coul/long/gpu*, *lj/cut/coul/long/kk*, *lj/cut/coul/long/intel*, *lj/cut/coul/long/opt*, *lj/cut/coul/long/omp*

### 19.146 pair\_style lj/cut/coul/msm command

Accelerator Variants: *lj/cut/coul/msm/gpu*, *lj/cut/coul/msm/omp*

### 19.147 pair\_style lj/cut/coul/wolf command

Accelerator Variants: *lj/cut/coul/wolf/omp*

#### 19.147.1 Syntax

<code>pair_style</code> style args
------------------------------------

- style = *lj/cut/coul/cut* or *lj/cut/coul/debye* or *lj/cut/coul/dsf* or *lj/cut/coul/long* *lj/cut/coul/msm* or *lj/cut/coul/wolf*
- args = list of arguments for a particular style

```

lj/cut/coul/cut args = cutoff (cutoff2)
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)
lj/cut/coul/debye args = kappa cutoff (cutoff2)
  kappa = inverse of the Debye length (inverse distance units)
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)
lj/cut/coul/dsf args = alpha cutoff (cutoff2)
  alpha = damping parameter (inverse distance units)
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (distance units)
lj/cut/coul/long args = cutoff (cutoff2)
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)
lj/cut/coul/msm args = cutoff (cutoff2)
  cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)
lj/cut/coul/wolf args = alpha cutoff (cutoff2)
  alpha = damping parameter (inverse distance units)
  cutoff = global cutoff for LJ (and Coulombic if only 2 arg) (distance units)
  cutoff2 = global cutoff for Coulombic (optional) (distance units)

```

## 19.147.2 Examples

```

pair_style lj/cut/coul/cut 10.0
pair_style lj/cut/coul/cut 10.0 8.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0
pair_coeff 1 1 100.0 3.5 9.0 9.0

pair_style lj/cut/coul/debye 1.5 3.0
pair_style lj/cut/coul/debye 1.5 2.5 5.0
pair_coeff * * 1.0 1.0
pair_coeff 1 1 1.0 1.5 2.5
pair_coeff 1 1 1.0 1.5 2.5 5.0

pair_style lj/cut/coul/dsf 0.05 2.5 10.0
pair_coeff * * 1.0 1.0
pair_coeff 1 1 1.0 1.0 2.5

pair_style lj/cut/coul/long 10.0
pair_style lj/cut/coul/long 10.0 8.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0

pair_style lj/cut/coul/msm 10.0
pair_style lj/cut/coul/msm 10.0 8.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0

pair_style lj/cut/coul/wolf 0.2 5. 10.0
pair_coeff * * 1.0 1.0

```

(continues on next page)

```
pair_coeff 1 1 1.0 1.0 2.5
```

### 19.147.3 Description

The *lj/cut/coul* styles compute the standard 12/6 Lennard-Jones potential, given by

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_c$$

$r_c$  is the cutoff.

Style *lj/cut/coul/cut* adds a Coulombic pairwise interaction given by

$$E = \frac{Cq_iq_j}{\epsilon r} \quad r < r_c$$

where  $C$  is an energy-conversion constant,  $q_i$  and  $q_j$  are the charges on the 2 atoms, and  $\epsilon$  is the dielectric constant which can be set by the *dielectric* command. If one cutoff is specified in the *pair\_style* command, it is used for both the LJ and Coulombic terms. If two cutoffs are specified, they are used as cutoffs for the LJ and Coulombic terms respectively.

Style *lj/cut/coul/debye* adds an additional  $\exp()$  damping factor to the Coulombic term, given by

$$E = \frac{Cq_iq_j}{\epsilon r} \exp(-\kappa r) \quad r < r_c$$

where  $\kappa$  is the inverse of the Debye length. This potential is another way to mimic the screening effect of a polar solvent.

Style *lj/cut/coul/dsf* computes the Coulombic term via the damped shifted force model described in *Fennell*, given by:

$$E = q_iq_j \left[ \frac{\operatorname{erfc}(\alpha r)}{r} - \frac{\operatorname{erfc}(\alpha r_c)}{r_c} + \left( \frac{\operatorname{erfc}(\alpha r_c)}{r_c^2} + \frac{2\alpha \exp(-\alpha^2 r_c^2)}{\sqrt{\pi} r_c} \right) (r - r_c) \right] \quad r < r_c$$

where  $\alpha$  is the damping parameter and  $\operatorname{erfc}()$  is the complementary error-function. This potential is essentially a short-range, spherically-truncated, charge-neutralized, shifted, pairwise  $1/r$  summation. The potential is based on Wolf summation, proposed as an alternative to Ewald summation for condensed phase systems where charge screening causes electrostatic interactions to become effectively short-ranged. In order for the electrostatic sum to be absolutely convergent, charge neutralization within the cutoff radius is enforced by shifting the potential through placement of image charges on the cutoff sphere. Convergence can often be improved by setting  $\alpha$  to a small non-zero value.

Styles *lj/cut/coul/long* and *lj/cut/coul/msm* compute the same Coulombic interactions as style *lj/cut/coul/cut* except that an additional damping factor is applied to the Coulombic term so it can be used in conjunction with the *kpace\_style* command and its *ewald* or *pppm* option. The Coulombic cutoff specified for this style means that pairwise interactions within this distance are computed directly; interactions outside that distance are computed in reciprocal space.

Style *coul/wolf* adds a Coulombic pairwise interaction via the Wolf summation method, described in *Wolf*, given by:

$$E_i = \frac{1}{2} \sum_{j \neq i} \frac{q_i q_j \operatorname{erfc}(\alpha r_{ij})}{r_{ij}} + \frac{1}{2} \sum_{j \neq i} \frac{q_i q_j \operatorname{erf}(\alpha r_{ij})}{r_{ij}} \quad r < r_c$$

where  $\alpha$  is the damping parameter, and  $\operatorname{erfc}()$  is the complementary error-function terms. This potential is essentially a short-range, spherically-truncated, charge-neutralized, shifted, pairwise  $1/r$  summation. With a manipulation of adding and subtracting a self term (for  $i = j$ ) to the first and second term on the right-hand-side, respectively, and a small enough  $\alpha$  damping parameter, the second term shrinks and the potential becomes a rapidly-converging real-space summation. With a long enough cutoff and small enough  $\alpha$  parameter, the energy and forces calculated by the Wolf summation method approach those of the Ewald sum. So it is a means of getting effective long-range interactions with a short-range potential.

### 19.147.4 Coefficients

For all of the *lj/cut/coul* pair styles, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- cutoff1 (distance units)
- cutoff2 (distance units)

Note that  $\sigma$  is defined in the LJ formula as the zero-crossing distance for the potential, not as the energy minimum at  $2^{\frac{1}{6}}\sigma$ .

The latter 2 coefficients are optional. If not specified, the global LJ and Coulombic cutoffs specified in the *pair\_style* command are used. If only one cutoff is specified, it is used as the cutoff for both LJ and Coulombic interactions for this type pair. If both coefficients are specified, they are used as the LJ and Coulombic cutoffs for this type pair.

For *lj/cut/coul/long* and *lj/cut/coul/msm* only the LJ cutoff can be specified since a Coulombic cutoff cannot be specified for an individual I,J type pair. All type pairs use the same global Coulombic cutoff specified in the *pair\_style* command.

A version of these styles with a soft core, *lj/cut/coul/soft* and *lj/cut/coul/long/soft*, suitable for use in free energy calculations, is part of the FEP package and is documented with the *pair\_style \*/soft* styles.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

### 19.147.5 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the *lj/cut* pair styles can be mixed. The default mix value is *geometric*. See the “*pair\_modify*” command for details.

All of the *lj/cut* pair styles support the *pair\_modify* shift option for the energy of the Lennard-Jones portion of the pair interaction.

The *lj/cut/coul/long* pair styles support the *pair\_modify* table option since they can tabulate the short-range portion of the long-range Coulombic interaction.

All of the *lj/cut* pair styles support the *pair\_modify* tail option for adding a long-range tail correction to the energy and pressure for the Lennard-Jones portion of the pair interaction.

All of the *lj/cut* pair styles write their information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

The *lj/cut/coul/long* pair styles support the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command, meaning the pairwise forces can be partitioned by distance at different levels of the rRESPA hierarchy. The other styles only support the *pair* keyword of *run\_style respa*. See the *run\_style* command for details.

---

### 19.147.6 Restrictions

The *lj/cut/coul/long* and *lj/cut/coul/msm* styles are part of the KSPACE package.

The *lj/cut/coul/debye*, *lj/cut/coul/dsf*, and *lj/cut/coul/wolf* styles are part of the EXTRA-PAIR package.

These styles are only enabled if LAMMPS was built with those respective packages. See the *Build package* page for more info.

### 19.147.7 Related commands

*pair\_coeff*

### 19.147.8 Default

none

---

(**Wolf**) D. Wolf, P. Keblinski, S. R. Phillpot, J. Eggebrecht, J Chem Phys, 110, 8254 (1999).

(**Fennell**) C. J. Fennell, J. D. Gezelter, J Chem Phys, 124, 234104 (2006).

## 19.148 pair\_style lj/cut/tip4p/cut command

Accelerator Variants: *lj/cut/tip4p/cut/omp*

## 19.149 pair\_style lj/cut/tip4p/long command

Accelerator Variants: *lj/cut/tip4p/long/gpu*, *lj/cut/tip4p/long/omp*, *lj/cut/tip4p/long/opt*

### 19.149.1 Syntax

<b>pair_style</b> style args
------------------------------

- style = *lj/cut/tip4p/cut* or *lj/cut/tip4p/long*
- args = list of arguments for a particular style

*lj/cut/tip4p/cut* args = otype htype btype atype qdist cutoff (cutoff2)  
otype, htype = atom types for TIP4P O and H  
btype, atype = bond and angle types for TIP4P waters  
qdist = distance from O atom to massless charge (distance units)  
cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)  
cutoff2 = global cutoff for Coulombic (optional) (distance units)



*lj/cut/tip4p/long* args = otype htype btype atype qdist cutoff (cutoff2)  
 otype,htype = atom types for TIP4P O and H  
 btype,atype = bond and angle types for TIP4P waters  
 qdist = distance from O atom to massless charge (distance units)  
 cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)  
 cutoff2 = global cutoff for Coulombic (optional) (distance units)

## 19.149.2 Examples

```
pair_style lj/cut/tip4p/cut 1 2 7 8 0.15 12.0
pair_style lj/cut/tip4p/cut 1 2 7 8 0.15 12.0 10.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0

pair_style lj/cut/tip4p/long 1 2 7 8 0.15 12.0
pair_style lj/cut/tip4p/long 1 2 7 8 0.15 12.0 10.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0
```

## 19.149.3 Description

The *lj/cut/tip4p* styles compute the standard 12/6 Lennard-Jones potential, given by

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_c$$

$r_c$  is the cutoff.

They add Coulombic pairwise interactions given by

$$E = \frac{Cq_iq_j}{\epsilon r} \quad r < r_c$$

where  $C$  is an energy-conversion constant,  $q_i$  and  $q_j$  are the charges on the 2 atoms, and  $\epsilon$  is the dielectric constant which can be set by the *dielectric* command. If one cutoff is specified in the *pair\_style* command, it is used for both the LJ and Coulombic terms. If two cutoffs are specified, they are used as cutoffs for the LJ and Coulombic terms respectively.

Style *lj/cut/tip4p/long* compute the same Coulombic interactions as style *lj/cut/tip4p/cut* except that an additional damping factor is applied to the Coulombic term so it can be used in conjunction with the *kpspace\_style* command and its *ewald* or *pppm* option. The Coulombic cutoff specified for this style means that pairwise interactions within this distance are computed directly; interactions outside that distance are computed in reciprocal space.

The *lj/cut/tip4p* styles implement the TIP4P water model of (*Jorgensen*), which introduces a massless site located a short distance away from the oxygen atom along the bisector of the HOH angle. The atomic types of the oxygen and hydrogen atoms, the bond and angle types for OH and HOH interactions, and the distance to the massless charge site are specified as *pair\_style* arguments. Style *lj/cut/tip4p/cut* uses a cutoff for Coulomb interactions; style *lj/cut/tip4p/long* is for use with a long-range Coulombic solver (Ewald or PPPM).

---

**Note:** For each TIP4P water molecule in your system, the atom IDs for the O and 2 H atoms must be consecutive, with the O atom first. This is to enable LAMMPS to “find” the 2 H atoms associated with each O atom. For example, if the atom ID of an O atom in a TIP4P water molecule is 500, then its 2 H atoms must have IDs 501 and 502.

---

See the [Howto tip4p](#) page for more information on how to use the TIP4P pair styles and lists of parameters to set. Note that the neighbor list cutoff for Coulomb interactions is effectively extended by a distance  $2*qdist$  when using the TIP4P pair style, to account for the offset distance of the fictitious charges on O atoms in water molecules. Thus it is typically best in an efficiency sense to use a LJ cutoff  $\geq$  Coulombic cutoff +  $2*qdist$ , to shrink the size of the neighbor list. This leads to slightly larger cost for the long-range calculation, so you can test the trade-off for your model.

### 19.149.4 Coefficients

For all of the *lj/cut* pair styles, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- LJ cutoff (distance units)

Note that  $\sigma$  is defined in the LJ formula as the zero-crossing distance for the potential, not as the energy minimum at  $2^{\frac{1}{6}}\sigma$ .

The last coefficient is optional. If not specified, the global LJ cutoff specified in the *pair\_style* command is used.

For *lj/cut/tip4p/cut* and *lj/cut/tip4p/long* only the LJ cutoff can be specified since a Coulombic cutoff cannot be specified for an individual I,J type pair. All type pairs use the same global Coulombic cutoff specified in the *pair\_style* command.

**Warning:** Because of how these pair styles implement the coulomb interactions by implicitly defining a fourth site for the negative charge of the TIP4P and similar water models, special care must be taken when using these pair styles with other computations that also use charges. Unless they are specially set up to also handle the implicit definition of the 4th site, results are likely incorrect. Example: [compute dipole/chunk](#). For the same reason, when using one of these pair styles with *pair\_style hybrid*, **all** coulomb interactions should be handled by a single sub-style with TIP4P support. All other instances and styles will “see” the M point charges at the position of the Oxygen atom and thus compute incorrect forces and energies. LAMMPS will print a warning when it detects one of these issues.

---

A version of these styles with a soft core, *lj/cut/tip4p/long/soft*, suitable for use in free energy calculations, is part of the FEP package and is documented with the *pair\_style \*/soft* styles.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 19.149.5 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the lj/cut pair styles can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

All of the *lj/cut* pair styles support the *pair\_modify* shift option for the energy of the Lennard-Jones portion of the pair interaction.

The *lj/cut/coul/long* and *lj/cut/tip4p/long* pair styles support the *pair\_modify* table option since they can tabulate the short-range portion of the long-range Coulombic interaction.

All of the *lj/cut* pair styles support the *pair\_modify* tail option for adding a long-range tail correction to the energy and pressure for the Lennard-Jones portion of the pair interaction.

All of the *lj/cut* pair styles write their information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

The *lj/cut* and *lj/cut/coul/long* pair styles support the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command, meaning the pairwise forces can be partitioned by distance at different levels of the rRESPA hierarchy. The other styles only support the *pair* keyword of run\_style respa. See the *run\_style* command for details.

---

### 19.149.6 Restrictions

The *lj/cut/tip4p/long* styles are part of the KSPACE package. The *lj/cut/tip4p/cut* style is part of the MOLECULE package. These styles are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

### 19.149.7 Related commands

*pair\_coeff*

### 19.149.8 Default

none

---

(Jorgensen) Jorgensen, Chandrasekhar, Madura, Impey, Klein, J Chem Phys, 79, 926 (1983).

## 19.150 pair\_style lj/expand command

Accelerator Variants: *lj/expand/gpu*, *lj/expand/kk*, *lj/expand/omp*

## 19.151 pair\_style lj/expand/coul/long command

Accelerator Variants: *lj/expand/coul/long/gpu*

### 19.151.1 Syntax

```
pair_style lj/expand cutoff
```

- cutoff = global cutoff for lj/expand interactions (distance units)

### 19.151.2 Examples

```
pair_style lj/expand 2.5
pair_coeff * * 1.0 1.0 0.5
pair_coeff 1 1 1.0 1.0 -0.2 2.0

pair_style lj/expand/coul/long 2.5
pair_style lj/expand/coul/long 2.5 4.0
pair_coeff * * 1.0 1.0 0.5
pair_coeff 1 1 1.0 1.0 -0.2 3.0
```

### 19.151.3 Description

Style *lj/expand* computes a LJ interaction with a distance shifted by delta which can be useful when particles are of different sizes, since it is different that using different sigma values in a standard LJ formula:

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r - \Delta} \right)^{12} - \left( \frac{\sigma}{r - \Delta} \right)^6 \right] \quad r < r_c + \Delta$$

$r_c$  is the cutoff which does not include the  $\Delta$  distance. I.e. the actual force cutoff is the sum of  $r_c + \Delta$ .

For all of the *lj/expand* pair styles, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- $\Delta$  (distance units)
- cutoff (distance units)

The  $\Delta$  values can be positive or negative. The last coefficient is optional. If not specified, the global LJ cutoff is used.

For *lj/expand/coul/long* only the LJ cutoff can be specified since a Coulombic cutoff cannot be specified for an individual I,J type pair. All type pairs use the same global Coulombic cutoff specified in the *pair\_style* command.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.151.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon, sigma, and shift coefficients and cutoff distance for this pair style can be mixed. Shift is always mixed via an *arithmetic* rule. The other coefficients are mixed according to the *pair\_modify* mix value. The default mix value is *geometric*. See the “*pair\_modify*” command for details.

This pair style supports the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style supports the *pair\_modify* tail option for adding a long-range tail correction to the energy and pressure of the pair interaction.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.151.5 Restrictions

none

#### 19.151.6 Related commands

*pair\_coeff*

#### 19.151.7 Default

none

### 19.152 pair\_style lj/long/coul/long command

Accelerator Variants: *lj/long/coul/long/intel*, *lj/long/coul/long/omp*, *lj/long/coul/long/opt*

## 19.153 pair\_style lj/long/tip4p/long command

Accelerator Variants: *lj/long/tip4p/long/omp*

### 19.153.1 Syntax

```
pair_style style args
```

- style = *lj/long/coul/long* or *lj/long/tip4p/long*
- args = list of arguments for a particular style

*lj/long/coul/long* args = flag\_lj flag\_coul cutoff (cutoff2)

flag\_lj = *long* or *cut* or *off*

*long* = use Kspace long-range summation for dispersion  $1/r^6$  term

*cut* = use a cutoff on dispersion  $1/r^6$  term

*off* = omit dispersion  $1/r^6$  term entirely

flag\_coul = *long* or *off*

*long* = use Kspace long-range summation for Coulombic  $1/r$  term

*off* = omit Coulombic term

cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

*lj/long/tip4p/long* args = flag\_lj flag\_coul otype htype btype atype qdist cutoff\_↪(cutoff2)

flag\_lj = *long* or *cut*

*long* = use Kspace long-range summation for dispersion  $1/r^6$  term

*cut* = use a cutoff

flag\_coul = *long* or *off*

*long* = use Kspace long-range summation for Coulombic  $1/r$  term

*off* = omit Coulombic term

otype,htype = atom types for TIP4P O and H

btype,atype = bond and angle types for TIP4P waters

qdist = distance from O atom to massless charge (distance units)

cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

### 19.153.2 Examples

```
pair_style lj/long/coul/long cut off 2.5
pair_style lj/long/coul/long cut long 2.5 4.0
pair_style lj/long/coul/long long long 2.5 4.0
pair_coeff * * 1 1
pair_coeff 1 1 1 3 4

pair_style lj/long/tip4p/long long long 1 2 7 8 0.15 12.0
pair_style lj/long/tip4p/long long long 1 2 7 8 0.15 12.0 10.0
pair_coeff * * 100.0 3.0
pair_coeff 1 1 100.0 3.5 9.0
```

### 19.153.3 Description

Style *lj/long/coul/long* computes the standard 12/6 Lennard-Jones potential:

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_c$$

with  $\epsilon$  and  $\sigma$  being the usual Lennard-Jones potential parameters, plus the Coulomb potential, given by:

$$E = \frac{Cq_iq_j}{\epsilon r} \quad r < r_c$$

where  $C$  is an energy-conversion constant,  $q_i$  and  $q_j$  are the charges on the 2 atoms,  $\epsilon$  is the dielectric constant which can be set by the *dielectric* command, and  $r_c$  is the cutoff. If one cutoff is specified in the *pair\_style* command, it is used for both the LJ and Coulombic terms. If two cutoffs are specified, they are used as cutoffs for the LJ and Coulombic terms respectively.

The purpose of this pair style is to capture long-range interactions resulting from both attractive  $1/r^6$  Lennard-Jones and Coulombic  $1/r$  interactions. This is done by use of the *flag\_lj* and *flag\_coul* settings. The *In 't Veld* paper has more details on when it is appropriate to include long-range  $1/r^6$  interactions, using this potential.

Style *lj/long/tip4p/long* implements the TIP4P water model of (*Jorgensen*), which introduces a massless site located a short distance away from the oxygen atom along the bisector of the HOH angle. The atomic types of the oxygen and hydrogen atoms, the bond and angle types for OH and HOH interactions, and the distance to the massless charge site are specified as *pair\_style* arguments.

---

**Note:** For each TIP4P water molecule in your system, the atom IDs for the O and 2 H atoms must be consecutive, with the O atom first. This is to enable LAMMPS to “find” the 2 H atoms associated with each O atom. For example, if the atom ID of an O atom in a TIP4P water molecule is 500, then its 2 H atoms must have IDs 501 and 502.

---

See the *Howto tip4p* page for more information on how to use the TIP4P pair style. Note that the neighbor list cutoff for Coulomb interactions is effectively extended by a distance  $2*qd_{\text{dist}}$  when using the TIP4P pair style, to account for the offset distance of the fictitious charges on O atoms in water molecules. Thus it is typically best in an efficiency sense to use a LJ cutoff  $\geq$  Coulombic cutoff +  $2*qd_{\text{dist}}$ , to shrink the size of the neighbor list. This leads to slightly larger cost for the long-range calculation, so you can test the trade-off for your model.

If *flag\_lj* is set to *long*, no cutoff is used on the LJ  $1/r^6$  dispersion term. The long-range portion can be calculated by using the *kpspace\_style ewald/disp or ppm/disp* commands. The specified LJ cutoff then determines which portion of the LJ interactions are computed directly by the pair potential versus which part is computed in reciprocal space via the Kspace style. If *flag\_lj* is set to *cut*, the LJ interactions are simply cutoff, as with *pair\_style lj/cut*.

If *flag\_coul* is set to *long*, no cutoff is used on the Coulombic interactions. The long-range portion can be calculated by using any of several *kpspace\_style* command options such as *pppm* or *ewald*. Note that if *flag\_lj* is also set to *long*, then the *ewald/disp* or *pppm/disp* Kspace style needs to be used to perform the long-range calculations for both the LJ and Coulombic interactions. If *flag\_coul* is set to *off*, Coulombic interactions are not computed.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- cutoff1 (distance units)
- cutoff2 (distance units)

Note that  $\sigma$  is defined in the LJ formula as the zero-crossing distance for the potential, not as the energy minimum at  $2^{1/6}\sigma$ .

The latter 2 coefficients are optional. If not specified, the global LJ and Coulombic cutoffs specified in the `pair_style` command are used. If only one cutoff is specified, it is used as the cutoff for both LJ and Coulombic interactions for this type pair. If both coefficients are specified, they are used as the LJ and Coulombic cutoffs for this type pair.

Note that if you are using `flag_lj` set to `long`, you cannot specify a LJ cutoff for an atom type pair, since only one global LJ cutoff is allowed. Similarly, if you are using `flag_coul` set to `long`, you cannot specify a Coulombic cutoff for an atom type pair, since only one global Coulombic cutoff is allowed.

For `lj/long/tip4p/long` only the LJ cutoff can be specified since a Coulombic cutoff cannot be specified for an individual I,J type pair. All type pairs use the same global Coulombic cutoff specified in the `pair_style` command.

---

A version of these styles with a soft core, `lj/cut/soft`, suitable for use in free energy calculations, is part of the FEP package and is documented with the `pair_style */soft` styles. The version with soft core is only available if LAMMPS was built with that package. See the [Build package](#) page for more info.

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix` *command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.153.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the `lj/long` pair styles can be mixed. The default mix value is *geometric*. See the “`pair_modify`” command for details.

These pair styles support the `pair_modify` shift option for the energy of the Lennard-Jones portion of the pair interaction, assuming `flag_lj` is `cut`.

These pair styles support the `pair_modify` table and table/disp options since they can tabulate the short-range portion of the long-range Coulombic and dispersion interactions.

These pair styles do not support the `pair_modify` tail option for adding a long-range tail correction to the Lennard-Jones portion of the energy and pressure.

These pair styles write their information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

The pair `lj/long/coul/long` styles support the use of the *inner*, *middle*, and *outer* keywords of the `run_style respa` command, meaning the pairwise forces can be partitioned by distance at different levels of the rRESPA hierarchy. See the `run_style` command for details.

---



### 19.153.5 Restrictions

These styles are part of the KSPACE package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.153.6 Related commands

*pair\_coeff*

### 19.153.7 Default

none

(In 't Veld) In 't Veld, Ismail, Grest, J Chem Phys, 127, 144711 (2007).

(Jorgensen) Jorgensen, Chandrasekhar, Madura, Impey, Klein, J Chem Phys, 79, 926 (1983).

## 19.154 pair\_style lj/relres command

Accelerator Variants: *lj/relres/omp*

### 19.154.1 Syntax

```
pair_style lj/relres Rsi Rso Rci Rco
```

- Rsi = inner switching cutoff between the fine-grained and coarse-grained potentials (distance units)
- Rso = outer switching cutoff between the fine-grained and coarse-grained potentials (distance units)
- Rci = inner cutoff beyond which the force smoothing for all interactions is applied (distance units)
- Rco = outer cutoff for all interactions (distance units)

### 19.154.2 Examples

```
pair_style lj/relres 4.0 5.0 8.0 10.0
pair_coeff 1 1 0.5 1.0 1.5 1.1
pair_coeff 2 2 0.5 1.0 0.0 0.0 3.0 3.5 6.0 7.0
```

### 19.154.3 Description

Pair style *lj/relres* computes a LJ interaction using the Relative Resolution (RelRes) framework which applies a fine-grained (FG) potential between near neighbors and a coarse-grained (CG) potential between far neighbors

(*Chaimovich1*). This approach can improve the computational efficiency by almost an order of magnitude, while maintaining the correct static and dynamic behavior of a reference system (*Chaimovich2*).

$$E = \begin{cases} 4\epsilon^{FG} \left[ \left( \frac{\sigma^{FG}}{r} \right)^{12} - \left( \frac{\sigma^{FG}}{r} \right)^6 \right] - \Gamma_{si}, & \text{if } r < r_{si}, \\ \sum_{m=0}^4 \gamma_{sm} (r - r_{si})^m - \Gamma_{so}, & \text{if } r_{si} \leq r < r_{so}, \\ 4\epsilon^{CG} \left[ \left( \frac{\sigma^{CG}}{r} \right)^{12} - \left( \frac{\sigma^{CG}}{r} \right)^6 \right] - \Gamma_c, & \text{if } r_{so} \leq r < r_{ci}, \\ \sum_{m=0}^4 \gamma_{cm} (r - r_{ci})^m - \Gamma_c, & \text{if } r_{ci} \leq r < r_{co}, \\ 0, & \text{if } r \geq r_{co}. \end{cases}$$

The FG parameters of the LJ potential ( $\epsilon^{FG}$  and  $\sigma^{FG}$ ) are applied up to the inner switching cutoff,  $r_{si}$ , while the CG parameters of the LJ potential ( $\epsilon^{CG}$  and  $\sigma^{CG}$ ) are applied beyond the outer switching cutoff,  $r_{so}$ . Between  $r_{si}$  and  $r_{so}$  a polynomial smoothing function is applied so that the force and its derivative are continuous between the FG and CG potentials. An analogous smoothing function is applied between the inner and outer cutoffs ( $r_{ci}$  and  $r_{co}$ ). The offsets  $\Gamma_{si}$ ,  $\Gamma_{so}$  and  $\Gamma_c$  ensure the continuity of the energy over the entire domain. The corresponding polynomial coefficients  $\gamma_{sm}$  and  $\gamma_{cm}$ , as well as the offsets are automatically computed by LAMMPS.

---

**Note:** Energy and force resulting from this methodology can be plotted via the `pair_write` command.

---

The following coefficients must be defined for each pair of atom types via the `pair_coeff` command as in the examples above, or in the data file or restart files read by the `read_data` or `read_restart` commands, or by mixing as will be described below:

- $\epsilon^{FG}$  (energy units)
- $\sigma^{FG}$  (distance units)
- $\epsilon^{CG}$  (energy units)
- $\sigma^{CG}$  (distance units)

Additional parameters can be defined to specify different  $r_{si}$ ,  $r_{so}$ ,  $r_{ci}$ ,  $r_{co}$  for a particular set of atom types:

- $r_{si}$  (distance units)
- $r_{so}$  (distance units)
- $r_{ci}$  (distance units)
- $r_{co}$  (distance units)

These parameters are optional, and they are used to override the global cutoffs as defined in the `pair_style` command. If not specified, the global values for  $r_{si}$ ,  $r_{so}$ ,  $r_{ci}$ , and  $r_{co}$  are used. If this override option is employed, all four arguments must be specified.

---

Here are some guidelines for using the `pair_style lj/relres` command.

In general, RelRes focuses on the speedup of pairwise interactions between all LJ sites. Importantly, it works with any settings and flags (e.g., `special_bonds` settings and `newton` flags) that can be used in a molecular simulation with the conventional LJ potential. In particular, all intramolecular topology with its energetics (i.e., bonds, angles, etc.) remains unaltered.

At the most basic level in the RelRes framework, all sites are mapped into clusters. Each cluster is just a collection of sites bonded together (the bonds themselves are not part of the cluster). In general, a molecule may be comprised of several clusters, and preferably, no two sites in a cluster are separated by more than two bonds. There are two categories of sites in RelRes: “hybrid” sites embody both FG and CG models, while “ordinary” sites embody just FG characteristics with no CG features. A given cluster has a single hybrid site (typically its central site) and several

ordinary sites (typically its peripheral sites). Notice that while clusters are necessary for the RelRes parameterization (discussed below), they are not actually defined in LAMMPS. Besides, the total number of sites in the cluster are called the “mapping ratio”, and this substantially impacts the computational efficiency of RelRes: For a mapping ratio of 3, the efficiency factor is around 4, and for a mapping ratio of 5, the efficiency factor is around 5 (*Chaimovich2*).

The flexibility of LAMMPS allows placing any values for the LJ parameters in the input script. However, here are the optimal recommendations for the RelRes parameters, which yield the correct structural and thermal behavior in a system of interest (*Chaimovich1*). One must first assign a complete set of parameters for the FG interactions that are applicable to all atom types. Regarding the parameters for the CG interactions, the rules rely on the site category (if it is a hybrid or an ordinary site). For atom types of ordinary sites,  $\epsilon^{CG}$  must be set to 0 (zero) while the specific value of  $\sigma^{CG}$  is irrelevant. For atom types of hybrid sites, the CG parameters should be generally calculated using the following equations:

$$\sigma_I^{CG} = \frac{\left(\sum_{\alpha \in A} \sqrt{\epsilon_{\alpha}^{FG} (\sigma_{\alpha}^{FG})^{12}}\right)^{1/2}}{\left(\sum_{\alpha \in A} \sqrt{\epsilon_{\alpha}^{FG} (\sigma_{\alpha}^{FG})^6}\right)^{1/3}} \quad \text{and} \quad \epsilon_I^{CG} = \frac{\left(\sum_{\alpha \in A} \sqrt{\epsilon_{\alpha}^{FG} (\sigma_{\alpha}^{FG})^6}\right)^4}{\left(\sum_{\alpha \in A} \sqrt{\epsilon_{\alpha}^{FG} (\sigma_{\alpha}^{FG})^{12}}\right)^2}$$

where  $I$  is an atom type of a hybrid site of a particular cluster  $A$ , and corresponding with this cluster, the summation proceeds over all of its sites  $\alpha$ . These equations are derived from the monopole term in the underlying Taylor series, and they are indeed relevant only if geometric mixing is applicable for the FG model; if this is not the case, Ref. (*Chaimovich2*) discusses the alternative formula, and in such a situation, the `pair_coeff` command should be explicitly used for all combinations of atom types  $I \neq J$ .

The switching distance (the midpoint between inner and outer switching cutoffs) is another crucial factor in RelRes: decreasing it improves the computational efficiency, yet if it is too small, the molecular simulations may not capture the system behavior correctly. As a rule of thumb, the switching distance should be approximately  $\sim 1.5\sigma$  (*Chaimovich1*); recommendations can be found in Ref. (*Chaimovich2*). Regarding the switching smoothing zone,  $\sim 0.1\sigma$  is recommended; if desired, smoothing can be eliminated by setting the inner switching cutoff,  $r_{si}$ , equal to the outer switching cutoff,  $r_{so}$  (the same is true for the other cutoffs  $r_{ci}$  and  $r_{co}$ ).

As an example, imagine that in your system, a molecule is comprised just of one cluster such that one atom type (#1) is associated with its hybrid site, and another atom type (#2) is associated with its ordinary sites (in total, there are 2 atom types). If geometric mixing is applicable, the following commands should be used:

```
pair_style lj/relres Rsi Rso Rci Rco
pair_coeff 1 1 epsilon_FG1 sigma_FG1 epsilon_CG1 sigma_CG1
pair_coeff 2 2 epsilon_FG2 sigma_FG2 0.0 0.0
pair_modify shift yes
```

In a more complex situation, there may be two distinct clusters in a system (these two clusters may be on same molecule or on different molecules), each with its own switching cutoffs. If there are still two atom types in each cluster as in the earlier example, the commands should be:

```
pair_style lj/relres Rsi Rso Rci Rco
pair_coeff 1 1 epsilon_FG1 sigma_FG1 epsilon_CG1 sigma_CG1 Rsi1 Rso1 Rci Rco
pair_coeff 2 2 epsilon_FG2 sigma_FG2 0.0 0.0 Rsi1 Rso1 Rci Rco
pair_coeff 3 3 epsilon_FG3 sigma_FG3 epsilon_CG3 sigma_CG3
pair_coeff 4 4 epsilon_FG4 sigma_FG4 0.0 0.0
pair_modify shift yes
```

In this example, the switching cutoffs for the first cluster (atom types 1 and 2) is defined explicitly in the `pair_coeff` command which overrides the global values, while the second cluster (atom types 3 and 4) uses the global definition from the `pair_style` command. The emphasis here is that the atom types that belong to a specific cluster should have the same switching/cutoff arguments.

In the case that geometric mixing is not applicable, for simulating the system from the previous example, we recommend using the following commands:

```
pair_style lj/relres Rsi Rso Rci Rco
pair_coeff 1 1 epsilon_FG1 sigma_FG1 epsilon_CG1 sigma_CG1 Rsi1 Rso1 Rci Rco
pair_coeff 1 2 epsilon_FG12 sigma_FG12 0.0 0.0 Rsi1 Rso1 Rci Rco
pair_coeff 1 3 epsilon_FG13 sigma_FG13 epsilon_CG13 sigma_CG13 Rsi13 Rso13 Rci Rco
pair_coeff 1 4 epsilon_FG14 sigma_FG14 0.0 0.0 Rsi13 Rso13 Rci Rco
pair_coeff 2 2 epsilon_FG2 sigma_FG2 0.0 0.0 Rsi1 Rso1 Rci Rco
pair_coeff 2 3 epsilon_FG23 sigma_FG23 0.0 0.0 Rsi13 Rso13 Rci Rco
pair_coeff 2 4 epsilon_FG24 sigma_FG24 0.0 0.0 Rsi13 Rso13 Rci Rco
pair_coeff 3 3 epsilon_FG3 sigma_FG3 epsilon_CG3 sigma_CG3
pair_coeff 3 4 epsilon_FG34 sigma_FG34 0.0 0.0
pair_coeff 4 4 epsilon_FG4 sigma_FG4 0.0 0.0
pair_modify shift yes
```

Notice that the CG parameters are mixed only for interactions between atom types associated with hybrid sites, and that the cutoffs are mixed on the cluster basis.

More examples can be found in the *examples/relres* folder.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.154.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs  $I, J$  with  $I \neq J$ , the  $\epsilon^{FG}$ ,  $\sigma^{FG}$ ,  $\epsilon^{CG}$ ,  $\sigma^{CG}$ ,  $r_{si}$ ,  $r_{so}$ ,  $r_{ci}$ , and  $r_{co}$  parameters for this pair style can be mixed, if not defined explicitly. All parameters are mixed according to the `pair_modify mix` option. The default mix value is *geometric*, and it is recommended to use with this *lj/relres* style. See the “`pair_modify`” command for details.

This pair style supports the `pair_modify shift` option for the energy of the pair interaction. It is recommended to set this option to *yes*. Otherwise, the offset  $\Gamma_c$  is set to zero. Constants  $\Gamma_{si}$  and  $\Gamma_{so}$  are not impacted by this option.

The `pair_modify table` option is not relevant for this pair style.

This pair style does not support the `pair_modify tail` option for adding long-range tail corrections to energy and pressure, since the energy of the pair interaction is smoothed to 0.0 at the cutoff.

This pair style writes its information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the `pair` keyword of the `run_style respa` command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.154.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.154.6 Related commands

*pair\_coeff*

### 19.154.7 Default

none

(Chaimovich1) A. Chaimovich, C. Peter and K. Kremer, J. Chem. Phys. 143, 243107 (2015).

(Chaimovich2) M. Chaimovich and A. Chaimovich, J. Chem. Theory Comput. 17, 1045-1059 (2021).

## 19.155 pair\_style lj/smooth command

Accelerator Variants: *lj/smooth/gpu*, *lj/smooth/omp*

### 19.155.1 Syntax

```
pair_style lj/smooth Rin Rc
```

- Rin = inner cutoff beyond which force smoothing will be applied (distance units)
- Rc = outer cutoff for lj/smooth interactions (distance units)

### 19.155.2 Examples

```
pair_style lj/smooth 8.0 10.0
pair_coeff * * 10.0 1.5
pair_coeff 1 1 20.0 1.3 7.0 9.0
```

### 19.155.3 Description

Style *lj/smooth* computes a LJ interaction with a force smoothing applied between the inner and outer cutoff.

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] \quad r < r_{in}$$

$$F = C_1 + C_2(r - r_{in}) + C_3(r - r_{in})^2 + C_4(r - r_{in})^3 \quad r_{in} < r < r_c$$

The polynomial coefficients C1, C2, C3, C4 are computed by LAMMPS to cause the force to vary smoothly from the inner cutoff  $r_{in}$  to the outer cutoff  $r_c$ .

At the inner cutoff the force and its first derivative will match the non-smoothed LJ formula. At the outer cutoff the force and its first derivative will be 0.0. The inner cutoff cannot be 0.0.

**Note:** this force smoothing causes the energy to be discontinuous both in its values and first derivative. This can lead to poor energy conservation and may require the use of a thermostat. Plot the energy and force resulting from this formula via the *pair\_write* command to see the effect.

---

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- $r_{in}$  (distance units)
- $r_c$  (distance units)

The last 2 coefficients are optional inner and outer cutoffs. If not specified, the global values for  $r_{in}$  and  $r_c$  are used.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.155.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon, sigma, Rin coefficients and the cutoff distance for this pair style can be mixed. Rin is a cutoff value and is mixed like the cutoff. The other coefficients are mixed according to the *pair\_modify* mix option. The default mix value is *geometric*. See the “*pair\_modify*” command for details.

This pair style supports the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure, since the energy of the pair interaction is smoothed to 0.0 at the cutoff.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.155.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.155.6 Related commands

*pair\_coeff*, *pair lj/smooth/linear*

### 19.155.7 Default

none

## 19.156 pair\_style lj/smooth/linear command

Accelerator Variants: *lj/smooth/linear/omp*

### 19.156.1 Syntax

```
pair_style lj/smooth/linear cutoff
```

- cutoff = global cutoff for Lennard-Jones interactions (distance units)

### 19.156.2 Examples

```
pair_style lj/smooth/linear 2.5
pair_coeff * * 1.0 1.0
pair_coeff 1 1 0.3 3.0 9.0
```

### 19.156.3 Description

Style *lj/smooth/linear* computes a truncated and force-shifted LJ interaction (aka Shifted Force Lennard-Jones) that combines the standard 12/6 Lennard-Jones function and subtracts a linear term based on the cutoff distance, so that both, the potential and the force, go continuously to zero at the cutoff  $R_c$  (*Toxvaerd*):

$$\phi(r) = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right]$$

$$E(r) = \phi(r) - \phi(R_c) - (r - R_c) \left. \frac{d\phi}{dr} \right|_{r=R_c} \quad r < R_c$$

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)

- cutoff (distance units)

The last coefficient is optional. If not specified, the global LJ cutoff specified in the `pair_style` command is used.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.156.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs  $I, J$  and  $I \neq J$ , the epsilon and sigma coefficients and cutoff distance can be mixed. The default mix value is geometric. See the “`pair_modify`” command for details.

This pair style does not support the *pair\_modify* shift option for the energy of the pair interaction, since it goes to 0.0 at the cutoff by construction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure, since the energy of the pair interaction is smoothed to 0.0 at the cutoff.

This pair style writes its information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.156.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

#### 19.156.6 Related commands

*pair\_coeff*, *pair lj/smooth*



## 19.156.7 Default

none

(Toxvaerd) Toxvaerd, Dyre, J Chem Phys, 134, 081102 (2011).

## 19.157 pair\_style lj/switch3/coulgauss/long command

## 19.158 pair\_style mm3/switch3/coulgauss/long command

### 19.158.1 Syntax

```
pair_style style args
```

- style = *lj/switch3/coulgauss/long* or *mm3/switch3/coulgauss/long*
- args = list of arguments for a particular style

*lj/switch3/coulgauss/long* args = cutoff (cutoff2) width  
 cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)  
 cutoff2 = global cutoff for Coulombic (optional) (distance units)  
 width = width parameter of the smoothing function (distance units)

*mm3/switch3/coulgauss/long* args = cutoff (cutoff2) width  
 cutoff = global cutoff for MM3 (and Coulombic if only 1 arg) (distance units)  
 cutoff2 = global cutoff for Coulombic (optional) (distance units)  
 width = width parameter of the smoothing function (distance units)

### 19.158.2 Examples

```
pair_style lj/switch3/coulgauss/long 12.0 3.0
pair_coeff 1 0.2 2.5 1.2

pair_style lj/switch3/coulgauss/long 12.0 10.0 3.0
pair_coeff 1 0.2 2.5 1.2

pair_style mm3/switch3/coulgauss/long 12.0 3.0
pair_coeff 1 0.2 2.5 1.2

pair_style mm3/switch3/coulgauss/long 12.0 10.0 3.0
pair_coeff 1 0.2 2.5 1.2
```

### 19.158.3 Description

The *lj/switch3/coulgauss* style evaluates the LJ vdW potential

$$E = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right]$$

The *mm3/switch3/coulgauss/long* style evaluates the MM3 vdW potential (*Allinger*)

$$E = \epsilon_{ij} \left[ -2.25 \left( \frac{r_{v,ij}}{r_{ij}} \right)^6 + 1.84(10)^5 \exp[-12.0r_{ij}/r_{v,ij}] \right] S_3(r_{ij})$$
$$r_{v,ij} = r_{v,i} + r_{v,j}$$
$$\epsilon_{ij} = \sqrt{\epsilon_i \epsilon_j}$$

Both potentials go smoothly to zero at the cutoff  $r_c$  as defined by the switching function

$$S_3(r) = \begin{cases} 1 & \text{if } r < r_c - w \\ 3x^2 - 2x^3 & \text{if } r < r_c \text{ with } x = \frac{r_c - r}{w} \\ 0 & \text{if } r \geq r_c \end{cases}$$

where  $w$  is the width defined in the arguments. This potential is combined with Coulomb interaction between Gaussian charge densities:

$$E = \frac{q_i q_j \operatorname{erf} \left( r / \sqrt{\gamma_1^2 + \gamma_2^2} \right)}{\epsilon r_{ij}}$$

where  $q_i$  and  $q_j$  are the charges on the 2 atoms,  $\epsilon$  is the dielectric constant which can be set by the *dielectric* command,  $\gamma_i$  and  $\gamma_j$  are the widths of the Gaussian charge distribution and  $\operatorname{erf}()$  is the error-function. This style has to be used in conjunction with the *kspace\_style* command

If one cutoff is specified it is used for both the vdW and Coulomb terms. If two cutoffs are specified, the first is used as the cutoff for the vdW terms, and the second is the cutoff for the Coulombic term.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $\epsilon$  (energy)
- $\sigma$  (distance)
- $\gamma$  (distance)

---

### 19.158.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the *lj/long* pair styles can be mixed. The default mix value is *geometric*. See the “*pair\_modify*” command for details.

Shifting the potential energy is not necessary because the switching function ensures that the potential is zero at the cut-off.

These pair styles support the *pair\_modify* table and options since they can tabulate the short-range portion of the long-range Coulombic interactions.

These pair styles do not support the *pair\_modify* tail option for adding a long-range tail correction to the Lennard-Jones portion of the energy and pressure.

These pair styles write their information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the `pair` keyword of the `run_style respa` command. They do not support the `inner`, `middle`, `outer` keywords.

### 19.158.5 Restrictions

These styles are part of the YAFF package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.158.6 Related commands

*pair\_coeff*

### 19.158.7 Default

none

## 19.159 pair\_style local/density command

### 19.159.1 Syntax

```
pair_style style arg
```

- `style` = *local/density*
- `arg` = name of file containing tabulated values of local density and the potential

### 19.159.2 Examples

```
pair_style local/density benzene_water.localdensity.table
pair_style hybrid/overlay table spline 500 local/density
pair_coeff * * local/density benzene_water.localdensity.table
```

### 19.159.3 Description

The local density (LD) potential is a mean-field manybody potential, and, in some way, a generalization of embedded atom models (EAM). The name “local density potential” arises from the fact that it assigns an energy to an atom depending on the number of neighboring atoms of a given type around it within a predefined spherical volume (i.e., within the cutoff). The bottom-up coarse-graining (CG) literature suggests that such potentials can be widely useful in capturing effective manybody forces in a computationally efficient manner and thus improve the quality of CG models of implicit solvation (*Sanyal1*) and phase-segregation in liquid mixtures (*Sanyal2*), and provide guidelines to determine the extent of manybody correlations present in a CG model (*Rosenberger*). The LD potential in LAMMPS is primarily intended to be used as a corrective potential over traditional pair potentials in bottom-up CG models via *hybrid/overlay pair style* with other explicit pair interaction terms (e.g., tabulated, Lennard-Jones, Morse etc.). Because

the LD potential is not a pair potential per se, it is implemented simply as a single auxiliary file with all specifications that will be read upon initialization.

---

**Note:** Thus when used as the only interaction in the system, there is no corresponding pair\_coeff command and when used with other pair styles using the hybrid/overlay option, the corresponding pair\_coeff command must be supplied \* as placeholders for the atom types.

---

### System with a single CG atom type:

A system of a single atom type (e.g., LJ argon) with a single local density (LD) potential would have an energy given by:

$$U_{LD} = \sum_i F(\rho_i)$$

where  $\rho_i$  is the LD at atom  $i$  and  $F(\rho)$  is similar in spirit to the embedding function used in EAM potentials. The LD at atom  $i$  is given by the sum

$$\rho_i = \sum_{j \neq i} \varphi(r_{ij})$$

where  $\varphi$  is an indicator function that is one at  $r=0$  and zero beyond a cutoff distance  $R_2$ . The choice of the functional form of  $\varphi$  is somewhat arbitrary, but the following piecewise cubic function has proven sufficiently general: (*Sanyal1*), (*Sanyal2*) (*Rosenberger*)

$$\varphi(r) = \begin{cases} 1 & r \leq R_1 \\ c_0 + c_2 r^2 + c_4 r^4 + c_6 r^6 & r \in (R_1, R_2) \\ 0 & r \geq R_2 \end{cases}$$

The constants  $c$  are chosen so that the indicator function smoothly interpolates between 1 and 0 between the distances  $R_1$  and  $R_2$ , which are called the inner and outer cutoffs, respectively. Thus  $\varphi(R_1) = 1$ ,  $\varphi(R_2) = 0$ ,  $d\varphi/dr @ (r=R_1) = d\varphi/dr @ (r=R_2) = 0$ . The embedding function  $F(\rho)$  may or may not have a closed-form expression. To maintain generality, it is practically represented with a spline-interpolated table over a predetermined range of  $\rho$ . Outside of that range it simply adopts zero values at the endpoints.

It can be shown that the total force between two atoms due to the LD potential takes the form of a pair force, which motivates its designation as a LAMMPS pair style. Please see (*Sanyal1*) for details of the derivation.

---

### Systems with arbitrary numbers of atom types:

The potential is easily generalized to systems involving multiple atom types:

$$U_{LD} = \sum_i a_\alpha F(\rho_i)$$

with the LD expressed as

$$\rho_i = \sum_{j \neq i} b_\beta \varphi(r_{ij})$$

where  $\alpha$  gives the type of atom  $i$ ,  $\beta$  the type of atom  $j$ , and the coefficients  $a$  and  $b$  filter for atom types as specified by the user.  $a$  is called the central atom filter as it determines to which atoms the potential applies;  $a_\alpha = 1$  if the LD potential applies to atom type  $\alpha$  else zero. On the other hand,  $b$  is called the neighbor atom filter because it specifies which atom types to use in the calculation of the LD;  $b_\beta = 1$  if atom type  $\beta$  contributes to the LD and zero otherwise.

---

**Note:** Note that the potentials need not be symmetric with respect to atom types, which is the reason for two distinct sets of coefficients  $a$  and  $b$ . An atom type may contribute to the LD but not the potential, or to the potential but not the LD. Such decisions are made by the user and should (ideally) be motivated on physical grounds for the problem at hand.

---

### General form for implementation in LAMMPS:

Of course, a system with many atom types may have many different possible LD potentials, each with their own atom type filters, cutoffs, and embedding functions. The most general form of this potential as implemented in the pair\_style local/density is:

$$U_{LD} = \sum_k U_{LD}^{(k)} = \sum_i \left[ \sum_k a_\alpha^{(k)} F^{(k)}(\rho_i^{(k)}) \right]$$

where,  $k$  is an index that spans the (arbitrary) number of applied LD potentials  $N\_LD$ . Each LD is calculated as before with:

$$\rho_i^{(k)} = \sum_j b_\beta^{(k)} \varphi^{(k)}(r_{ij})$$

The superscript on the indicator function phi simply indicates that it is associated with specific values of the cutoff distances  $R1(k)$  and  $R2(k)$ . In summary, there may be  $N\_LD$  distinct LD potentials. With each potential type ( $k$ ), one must specify:

- the inner and outer cutoffs as  $R1$  and  $R2$
  - the central type filter  $a(k)$ , where  $k = 1, 2, \dots, N\_LD$
  - the neighbor type filter  $b(k)$ , where  $k = 1, 2, \dots, N\_LD$
  - the LD potential function  $F(k)(\rho)$ , typically as a table that is later spline-interpolated
- 

### Tabulated input file format:

```

Line 1:          comment or blank (ignored)
Line 2:          comment or blank (ignored)
Line 3:          N_LD N_rho (# of LD potentials and # of tabulated values, single_
↳space separated)
Line 4:          blank (ignored)
Line 5:          R1(k) R2(k) (lower and upper cutoffs, single space separated)
Line 6:          central-types (central atom types, single space separated)
Line 7:          neighbor-types (neighbor atom types single space separated)
Line 8:          rho_min rho_max drho (min, max and diff. in tabulated rho values,
↳single space separated)
Line 9:          F(k)(rho_min + 0.drho)
Line 10:         F(k)(rho_min + 1.drho)
Line 11:         F(k)(rho_min + 2.drho)
...
Line 9+N_rho:    F(k)(rho_min + N_rho . drho)
Line 10+N_rho:   blank (ignored)

Block 2

```

(continues on next page)

(continued from previous page)

Block 3

Block N\_LD

Lines 5 to 9+N\_rho constitute the first block. Thus the input file is separated (by blank lines) into N\_LD blocks each representing a separate LD potential and each specifying its own upper and lower cutoffs, central and neighbor atoms, and potential. In general, blank lines anywhere are ignored.

---

#### 19.159.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support automatic mixing. For atom type pairs  $\alpha$ ,  $\beta$  and  $\alpha \neq \beta$ , even if LD potentials of type  $(\alpha, \alpha)$  and  $(\beta, \beta)$  are provided, you will need to explicitly provide LD potential types  $(\alpha, \beta)$  and  $(\beta, \alpha)$  if need be (Here, the notation  $(\alpha, \beta)$  means that  $\alpha$  is the central atom to which the LD potential is applied and  $\beta$  is the neighbor atom which contributes to the LD potential on  $\alpha$ ).

This pair style does not support the *pair\_modify* shift, table, and tail options.

The local/density pair style does not write its information to *binary restart files*, since it is stored in tabulated potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

---

#### 19.159.5 Restrictions

The local/density pair style is a part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 19.159.6 Related commands

*pair\_coeff*

#### 19.159.7 Default

none

---

**(Sanyal1)** Sanyal and Shell, Journal of Chemical Physics, 2016, 145 (3), 034109.

**(Sanyal2)** Sanyal and Shell, Journal of Physical Chemistry B, 122 (21), 5678-5693.

**(Rosenberger)** Rosenberger, Sanyal, Shell and van der Vegt, Journal of Chemical Physics, 2019, 151 (4), 044111.

## 19.160 pair\_style lubricate command

Accelerator Variants: *lubricate/omp*

## 19.161 pair\_style lubricate/poly command

Accelerator Variants: *lubricate/poly/omp*

### 19.161.1 Syntax

```
pair_style style mu flaglog flagfld cutinner cutoff flagHI flagVF
```

- style = *lubricate* or *lubricate/poly*
- mu = dynamic viscosity (dynamic viscosity units)
- flaglog = 0/1 to exclude/include log terms in the lubrication approximation
- flagfld = 0/1 to exclude/include Fast Lubrication Dynamics (FLD) effects
- cutinner = inner cutoff distance (distance units)
- cutoff = outer cutoff for interactions (distance units)
- flagHI (optional) = 0/1 to exclude/include 1/r hydrodynamic interactions
- flagVF (optional) = 0/1 to exclude/include volume fraction corrections in the long-range isotropic terms

### 19.161.2 Examples

(all assume radius = 1)

```
pair_style lubricate 1.5 1 1 2.01 2.5
pair_coeff 1 1 2.05 2.8
pair_coeff * *

pair_style lubricate 1.5 1 1 2.01 2.5
pair_coeff * *
variable mu equal ramp(1,2)
fix 1 all adapt 1 pair lubricate mu * * v_mu
```

### 19.161.3 Description

Styles *lubricate* and *lubricate/poly* compute hydrodynamic interactions between mono-disperse finite-size spherical particles in a pairwise fashion. The interactions have 2 components. The first is Ball-Melrose lubrication terms via the formulas in (*Ball and Melrose*)

$$W = -a_{sq}|(v_1 - v_2) \bullet \mathbf{nn}|^2 - a_{sh}(\omega_1 + \omega_2) \bullet (\mathbf{I} - \mathbf{nn}) - 2\Omega_N|^2 - a_{pu}|(\omega_1 - \omega_2) \bullet (\mathbf{I} - \mathbf{nn})|^2 - a_{tw}|(\omega_1 - \omega_2) \bullet \mathbf{nn}|^2 \quad r < r_c$$

$$\Omega_N = \mathbf{n} \times (v_1 - v_2)/r$$

which represents the dissipation  $W$  between two nearby particles due to their relative velocities in the presence of a background solvent with viscosity  $\mu$ . Note that this is dynamic viscosity which has units of mass/distance/time, not kinematic viscosity.

The  $Asq$  (squeeze) term is the strongest and is included if  $flagHI$  is set to 1 (default). It scales as  $1/gap$  where  $gap$  is the separation between the surfaces of the 2 particles. The  $Ash$  (shear) and  $Apu$  (pump) terms are only included if  $flaglog$  is set to 1. They are the next strongest interactions, and the only other singular interaction, and scale as  $\log(gap)$ . Note that  $flaglog = 1$  and  $flagHI = 0$  is invalid, and will result in a warning message, after which  $flagHI$  will be set to 1. The  $Atw$  (twist) term is currently not included. It is typically a very small contribution to the lubrication forces.

The  $flagHI$  and  $flagVF$  settings are optional. Neither should be used, or both must be defined.

$Cutinner$  sets the minimum center-to-center separation that will be used in calculations irrespective of the actual separation.  $Cutoff$  is the maximum center-to-center separation at which an interaction is computed. Using a  $cutoff$  less than 3 radii is recommended if  $flaglog$  is set to 1.

The other component is due to the Fast Lubrication Dynamics (FLD) approximation, described in ([Kumar](#)), which can be represented by the following equation

$$F^H = -R_{FU}(U - U^\infty) + R_{FE}E^\infty$$

where  $U$  represents the velocities and angular velocities of the particles,  $U^\infty$  represents the velocity and the angular velocity of the undisturbed fluid, and  $E^\infty$  represents the rate of strain tensor of the undisturbed fluid with viscosity  $\mu$ . Again, note that this is dynamic viscosity which has units of mass/distance/time, not kinematic viscosity. Volume fraction corrections to  $R_{FU}$  are included as long as  $flagVF$  is set to 1 (default).

---

**Note:** When using the FLD terms, these pair styles are designed to be used with explicit time integration and a correspondingly small timestep. Thus either `fix nve/sphere` or `fix nve/asphere` should be used for time integration. To perform implicit FLD, see the `pair_style lubricateU` command.

---

Style `lubricate` requires monodisperse spherical particles; style `lubricate/poly` allows for polydisperse spherical particles.

The viscosity  $\mu$  can be varied in a time-dependent manner over the course of a simulation, in which case in which case the `pair_style` setting for  $\mu$  will be overridden. See the `fix adapt` command for details.

If the suspension is sheared via the `fix deform` command then the pair style uses the shear rate to adjust the hydrodynamic interactions accordingly. Volume changes due to `fix deform` are accounted for when computing the volume fraction corrections to  $R_{FU}$ .

When computing the volume fraction corrections to  $R_{FU}$ , the presence of walls (whether moving or stationary) will affect the volume fraction available to colloidal particles. This is currently accounted for with the following types of walls: `wall/lj93`, `wall/lj126`, `wall/colloid`, and `wall/harmonic`. For these wall styles, the correct volume fraction will be used when walls do not coincide with the box boundary, as well as when walls move and thereby cause a change in the volume fraction. Other wall styles will still work, but they will result in the volume fraction being computed based on the box boundaries.

Since lubrication forces are dissipative, it is usually desirable to thermostat the system at a constant temperature. If Brownian motion (at a constant temperature) is desired, it can be set using the `pair_style brownian` command. These pair styles and the brownian style should use consistent parameters for  $\mu$ ,  $flaglog$ ,  $flagfld$ ,  $cutinner$ ,  $cutoff$ ,  $flagHI$  and  $flagVF$ .

---

The following coefficients must be defined for each pair of atoms types via the `pair_coeff` command as in the examples above, or in the data file or restart files read by the `read_data` or `read_restart` commands, or by mixing as described below:

- `cutinner` (distance units)



- cutoff (distance units)

The two coefficients are optional. If neither is specified, the two cutoffs specified in the `pair_style` command are used. Otherwise both must be specified.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* [command-line switch](#) when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.161.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs  $I, J$  and  $I \neq J$ , the two cutoff distances for this pair style can be mixed. The default mix value is *geometric*. See the “`pair_modify`” command for details.

This pair style does not support the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.161.5 Restrictions

These styles are part of the COLLOID package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

Only spherical monodisperse particles are allowed for `pair_style lubricate`.

Only spherical particles are allowed for `pair_style lubricate/poly`.

These pair styles will not restart exactly when using the *read\_restart* command, though they should provide statistically similar results. This is because the forces they compute depend on atom velocities. See the *read\_restart* command for more details.

### 19.161.6 Related commands

*pair\_coeff*, *pair\_style lubricateU*

### 19.161.7 Default

The default settings for the optional args are `flagHI = 1` and `flagVF = 1`.

---

**(Ball)** Ball and Melrose, *Physica A*, 247, 444-472 (1997).

**(Kumar)** Kumar and Higdon, *Phys Rev E*, 82, 051401 (2010). See also his thesis for more details: A. Kumar, "Microscale Dynamics in Suspensions of Non-spherical Particles", Thesis, University of Illinois Urbana-Champaign, (2010). (<https://www.ideals.illinois.edu/handle/2142/16032>)

## 19.162 *pair\_style lubricateU* command

## 19.163 *pair\_style lubricateU/poly* command

### 19.163.1 Syntax

```
pair_style style mu flaglog cutinner cutoff gdot flagHI flagVF
```

- `style = lubricateU` or `lubricateU/poly`
- `mu` = dynamic viscosity (dynamic viscosity units)
- `flaglog = 0/1` to exclude/include log terms in the lubrication approximation
- `cutinner` = inner cut off distance (distance units)
- `cutoff` = outer cutoff for interactions (distance units)
- `gdot` = shear rate (1/time units)
- `flagHI` (optional) = 0/1 to exclude/include 1/r hydrodynamic interactions
- `flagVF` (optional) = 0/1 to exclude/include volume fraction corrections in the long-range isotropic terms

### 19.163.2 Examples

(all assume radius = 1)

```
pair_style lubricateU 1.5 1 2.01 2.5 0.01 1 1  
pair_coeff 1 1 2.05 2.8  
pair_coeff * *
```

### 19.163.3 Description

Styles *lubricateU* and *lubricateU/poly* compute velocities and angular velocities for finite-size spherical particles such that the hydrodynamic interaction balances the force and torque due to all other types of interactions.

The interactions have 2 components. The first is Ball-Melrose lubrication terms via the formulas in (*Ball and Melrose*)

$$W = -a_{sq}|(v_1 - v_2) \bullet \mathbf{nn}|^2 - a_{sh}|(\omega_1 + \omega_2) \bullet (\mathbf{I} - \mathbf{nn}) - 2\Omega_N|^2 - a_{pu}|(\omega_1 - \omega_2) \bullet (\mathbf{I} - \mathbf{nn})|^2 - a_{tw}|(\omega_1 - \omega_2) \bullet \mathbf{nn}|^2 \quad r < r_c$$

$$\Omega_N = \mathbf{n} \times (v_1 - v_2)/r$$

which represents the dissipation  $W$  between two nearby particles due to their relative velocities in the presence of a background solvent with viscosity  $\mu$ . Note that this is dynamic viscosity which has units of mass/distance/time, not kinematic viscosity.

The *Asq* (squeeze) term is the strongest and is included as long as *flagHI* is set to 1 (default). It scales as  $1/\text{gap}$  where *gap* is the separation between the surfaces of the 2 particles. The *Ash* (shear) and *Apu* (pump) terms are only included if *flaglog* is set to 1. They are the next strongest interactions, and the only other singular interaction, and scale as  $\log(\text{gap})$ . Note that *flaglog* = 1 and *flagHI* = 0 is invalid, and will result in a warning message, after which *flagHI* will be set to 1. The *Atw* (twist) term is currently not included. It is typically a very small contribution to the lubrication forces.

The *flagHI* and *flagVF* settings are optional. Neither should be used, or both must be defined.

*Cutinner* sets the minimum center-to-center separation that will be used in calculations irrespective of the actual separation. *Cutoff* is the maximum center-to-center separation at which an interaction is computed. Using a *cutoff* less than 3 radii is recommended if *flaglog* is set to 1.

The other component is due to the Fast Lubrication Dynamics (FLD) approximation, described in (*Kumar*). The equation being solved to balance the forces and torques is

$$-R_{FU}(U - U^\infty) = -R_{FE}E^\infty - F^{rest}$$

where  $U$  represents the velocities and angular velocities of the particles,  $U^\infty$  represents the velocities and the angular velocities of the undisturbed fluid, and  $E^\infty$  represents the rate of strain tensor of the undisturbed fluid flow with viscosity  $\mu$ . Again, note that this is dynamic viscosity which has units of mass/distance/time, not kinematic viscosity. Volume fraction corrections to  $R_{FU}$  are included if *flagVF* is set to 1 (default).

*Frest* represents the forces and torques due to all other types of interactions, e.g. Brownian, electrostatic etc. Note that this algorithm neglects the inertial terms, thereby removing the restriction of resolving the small inertial time scale, which may not be of interest for colloidal particles. This pair style solves for the velocity such that the hydrodynamic force balances all other types of forces, thereby resulting in a net zero force (zero inertia limit). When defining this pair style, it must be defined last so that when this style is invoked all other types of forces have already been computed. For the same reason, it won't work if additional non-pair styles are defined (such as bond or Kspace forces) as they are calculated in LAMMPS after the pairwise interactions have been computed.

---

**Note:** When using these styles, the these pair styles are designed to be used with implicit time integration and a correspondingly larger timestep. Thus either *fix nve/noforce* should be used for spherical particles defined via *atom\_style sphere* or *fix nve/asphere/noforce* should be used for spherical particles defined via *atom\_style ellipsoid*. This is because the velocity and angular momentum of each particle is set by the pair style, and should not be reset by the time integration fix.

---

Style *lubricateU* requires monodisperse spherical particles; style *lubricateU/poly* allows for polydisperse spherical particles.

If the suspension is sheared via the *fix deform* command then the pair style uses the shear rate to adjust the hydrodynamic interactions accordingly. Volume changes due to fix deform are accounted for when computing the volume fraction corrections to R\_FU.

When computing the volume fraction corrections to R\_FU, the presence of walls (whether moving or stationary) will affect the volume fraction available to colloidal particles. This is currently accounted for with the following types of walls: *wall/lj93*, *wall/lj126*, *wall/colloid*, and *wall/harmonic*. For these wall styles, the correct volume fraction will be used when walls do not coincide with the box boundary, as well as when walls move and thereby cause a change in the volume fraction. To use these wall styles with pair\_style *lubricateU* or *lubricateU/poly*, the *fld yes* option must be specified in the fix wall command.

Since lubrication forces are dissipative, it is usually desirable to thermostat the system at a constant temperature. If Brownian motion (at a constant temperature) is desired, it can be set using the *pair\_style brownian* command. These pair styles and the brownian style should use consistent parameters for *mu*, *flaglog*, *flagfld*, *cutinner*, *cutoff*, *flagHI* and *flagVF*.

---

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- cutinner (distance units)
- cutoff (distance units)

The two coefficients are optional. If neither is specified, the two cutoffs specified in the pair\_style command are used. Otherwise both must be specified.

---

#### 19.163.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the two cutoff distances for this pair style can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

This pair style does not support the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.163.5 Restrictions

These styles are part of the COLLOID package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Currently, these pair styles assume that all other types of forces/torques on the particles have been already been computed when it is invoked. This requires this style to be defined as the last of the pair styles, and that no fixes apply additional constraint forces. One exception is the *fix wall/colloid* commands, which has an “fld” option to apply their wall forces correctly.

---

Only spherical monodisperse particles are allowed for pair\_style lubricateU.

Only spherical particles are allowed for pair\_style lubricateU/poly.

For sheared suspensions, it is assumed that the shearing is done in the xy plane, with x being the velocity direction and y being the velocity-gradient direction. In this case, one must use *fix deform* with the same rate of shear (erate).

### 19.163.6 Related commands

*pair\_coeff*, *pair\_style lubricate*

### 19.163.7 Default

The default settings for the optional args are flagHI = 1 and flagVF = 1.

---

**(Ball)** Ball and Melrose, Physica A, 247, 444-472 (1997).

**(Kumar)** Kumar and Higdon, Phys Rev E, 82, 051401 (2010).

## 19.164 pair\_style lj/mdf command

## 19.165 pair\_style buck/mdf command

## 19.166 pair\_style lennard/mdf command

### 19.166.1 Syntax

**pair\_style** style args

- style = *lj/mdf* or *buck/mdf* or *lennard/mdf*
- args = list of arguments for a particular style

*lj/mdf* args = cutoff1 cutoff2

cutoff1 = inner cutoff for the start of the tapering function

cutoff2 = out cutoff for the end of the tapering function

*buck/mdf* args = cutoff1 cutoff2

cutoff1 = inner cutoff for the start of the tapering function

cutoff2 = out cutoff for the end of the tapering function

*lennard/mdf* args = cutoff1 cutoff2

cutoff1 = inner cutoff for the start of the tapering function

cutoff2 = out cutoff for the end of the tapering function

## 19.166.2 Examples

```

pair_style lj/mdf 2.5 3.0
pair_coeff * * 1.0 1.0
pair_coeff 1 1 1.1 2.8 3.0 3.2

pair_style buck/mdf 2.5 3.0
pair_coeff * * 100.0 1.5 200.0
pair_coeff * * 100.0 1.5 200.0 3.0 3.5

pair_style lennard/mdf 2.5 3.0
pair_coeff * * 1.0 1.0
pair_coeff 1 1 1021760.3664 2120.317338 3.0 3.2

```

## 19.166.3 Description

The *lj/mdf*, *buck/mdf* and *lennard/mdf* compute the standard 12-6 Lennard-Jones and Buckingham potential with the addition of a taper function that ramps the energy and force smoothly to zero between an inner and outer cutoff.

$$E_{smooth}(r) = E(r) * f(r)$$

The tapering,  $f(r)$ , is done by using the Mei, Davenport, Fernando function (*Mei*).

$$\begin{aligned}
 f(r) &= 1.0 && \text{for } r < r_m \\
 f(r) &= (1 - x)^3 * (1 + 3x + 6x^2) && \text{for } r_m < r < r_{cut} \\
 f(r) &= 0.0 && \text{for } r \geq r_{cut}
 \end{aligned}$$

where

$$x = \frac{(r - r_m)}{(r_{cut} - r_m)}$$

Here  $r_m$  is the inner cutoff radius and  $r_{cut}$  is the outer cutoff radius.

For the *lj/mdf* pair\_style, the potential energy,  $E(r)$ , is the standard 12-6 Lennard-Jones written in the epsilon/sigma form:

$$E(r) = 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right]$$

Either the first two or all of the following coefficients must be defined for each pair of atoms types via the pair\_coeff command as in the examples above, or in the data file read by the *read\_data*. The two cutoffs default to the global values and  $\epsilon$  and  $\sigma$  can also be determined by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- $r_m$  (distance units)
- $r_{cut}$  (distance units)

For the *buck/mdf* pair\_style, the potential energy,  $E(r)$ , is the standard Buckingham potential with three required coefficients. The two cutoffs can be omitted and default to the corresponding global values:

$$E(r) = A e^{(-r/\rho)} - \frac{C}{r^6}$$

- $A$  (energy units)
  - $\rho$  (distance units)
  - $C$  (energy-distance<sup>6</sup> units)
  - $r_m$  (distance units)
  - $r_{cut}$  (distance units)
- 

For the *lennard/mdf* pair\_style, the potential energy,  $E(r)$ , is the standard 12-6 Lennard-Jones written in the A/B form:

$$E(r) = \frac{A}{r^{12}} - \frac{B}{r^6}$$

The following coefficients must be defined for each pair of atoms types via the pair\_coeff command as in the examples above, or in the data file read by the read\_data commands, or by mixing as described below. The two cutoffs default to their global values and must be either both given or both left out:

- $A$  (energy-distance<sup>12</sup> units)
  - $B$  (energy-distance<sup>6</sup> units)
  - $r_m$  (distance units)
  - $r_{cut}$  (distance units)
- 

#### 19.166.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the  $\epsilon$  and  $\sigma$  coefficients and cutoff distances for the lj/mdf pair style can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details. The other two pair styles buck/mdf and lennard/mdf do not support mixing, so all I,J pairs of coefficients must be specified explicitly.

None of the lj/mdf, buck/mdf, or lennard/mdf pair styles supports the *pair\_modify* shift option or long-range tail corrections to pressure and energy.

These styles write their information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

These styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

---

#### 19.166.5 Restrictions

These pair styles can only be used if LAMMPS was built with the EXTRA-PAIR package. See the *Build package* doc page for more info.

## 19.166.6 Related commands

*pair\_coeff*

## 19.166.7 Default

none

---

(Mei) Mei, Davenport, Fernando, Phys Rev B, 43 4653 (1991)

## 19.167 pair\_style meam command

Accelerator Variants: *meam/kk*

### 19.167.1 Syntax

```
pair_style meam
```

### 19.167.2 Examples

```
pair_style meam
pair_coeff * * ../potentials/library.meam Si ../potentials/si.meam Si
pair_coeff * * ../potentials/library.meam Ni Al NULL Ni Al Ni Ni
```

### 19.167.3 Description

---

**Note:** The behavior of the MEAM potential for alloy systems has changed as of November 2010; see description below of the `mixture_ref_t` parameter

---

Pair style *meam* computes non-bonded interactions for a variety of materials using the modified embedded-atom method (MEAM) (*Baskes*). Conceptually, it is an extension to the original *EAM method* which adds angular forces. It is thus suitable for modeling metals and alloys with fcc, bcc, hcp and diamond cubic structures, as well as materials with covalent interactions like silicon and carbon. This *meam* pair style is a translation of the original Fortran version to C++. It is functionally equivalent but more efficient and has additional features. The Fortran version of the *meam* pair style has been removed from LAMMPS after the 12 December 2018 release.

In the MEAM formulation, the total energy  $E$  of a system of atoms is given by:

$$E = \sum_i \left\{ F_i(\bar{\rho}_i) + \frac{1}{2} \sum_{i \neq j} \phi_{ij}(r_{ij}) \right\}$$

where  $F$  is the embedding energy which is a function of the atomic electron density  $\rho$ , and  $\phi$  is a pair potential interaction. The pair interaction is summed over all neighbors  $J$  of atom  $I$  within the cutoff distance. As with EAM, the multi-body nature of the MEAM potential is a result of the embedding energy term. Details of the computation of the embedding and pair energies, as implemented in LAMMPS, are given in (*Gullet*) and references therein.



The various parameters in the MEAM formulas are listed in two files which are specified by the `pair_coeff` command. These are ASCII text files in a format consistent with other MD codes that implement MEAM potentials, such as the serial DYNAMO code and Warp. Several MEAM potential files with parameters for different materials are included in the “potentials” directory of the LAMMPS distribution with a “.meam” suffix. All of these are parameterized in terms of LAMMPS *metal units*.

Note that unlike for other potentials, cutoffs for MEAM potentials are not set in the `pair_style` or `pair_coeff` command; they are specified in the MEAM potential files themselves.

Only a single `pair_coeff` command is used with the `meam` style which specifies two MEAM files and the element(s) to extract information for. The MEAM elements are mapped to LAMMPS atom types by specifying N additional arguments after the second filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- MEAM library file
- Element1, Element2, ...
- MEAM parameter file
- N element names = mapping of MEAM elements to atom types

See the `pair_coeff` page for alternate ways to specify the path for the potential files.

As an example, the `potentials/library.meam` file has generic MEAM settings for a variety of elements. The `potentials/SiC.meam` file has specific parameter settings for a Si and C alloy system. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Si, and the fourth to be C, you would use the following `pair_coeff` command:

```
pair_coeff * * library.meam Si C sic.meam Si Si Si C
```

The first 2 arguments must be `**` so as to span all LAMMPS atom types. The first filename is the element library file. The list of elements following it extracts lines from the library file and assigns numeric indices to these elements. The second filename is the alloy parameter file, which refers to elements using the numeric indices assigned before. The arguments after the parameter file map LAMMPS atom types to elements, i.e. LAMMPS atom types 1,2,3 to the MEAM Si element. The final C argument maps LAMMPS atom type 4 to the MEAM C element.

If the second filename is specified as NULL, no parameter file is read, which simply means the generic parameters in the library file are used. Use of the NULL specification for the parameter file is discouraged for systems with more than a single element type (e.g. alloys), since the parameter file is expected to set element interaction terms that are not captured by the information in the library file.

If a mapping value is specified as NULL, the mapping is not performed. This can be used when a `meam` potential is used as part of the `hybrid` pair style. The NULL values are placeholders for atom types that will be used with other potentials.

---

**Note:** If the second filename is NULL, the element names between the two filenames can appear in any order, e.g. “Si C” or “C Si” in the example above. However, if the second filename is **not** NULL (as in the example above), it contains settings that are indexed **by numbers** for the elements that precede it. Thus you need to insure that you list the elements between the filenames in an order consistent with how the values in the second filename are indexed. See details below on the syntax for settings in the second file.

---

The MEAM library file provided with LAMMPS has the name `potentials/library.meam`. It is the “meamf” file used by other MD codes. Aside from blank and comment lines (starting with # which can appear anywhere), it is formatted as a series of entries, each of which has 19 parameters and can span multiple lines:

```
elt, lat, z, ielement, atwt, alpha, b0, b1, b2, b3, alat, esub, asub, t0, t1, t2, t3, rozero, ibar
```

The `elt` and `lat` parameters are text strings, such as `elt = Si` or `Cu` and `lat = dia` or `fcc`. Because the library file is used by Fortran MD codes, these strings may be enclosed in single quotes, but this is not required. The other numeric

parameters match values in the formulas above. The value of the *elt* string is what is used in the `pair_coeff` command to identify which settings from the library file you wish to read in. There can be multiple entries in the library file with the same *elt* value; LAMMPS reads the first matching entry it finds and ignores the rest.

Other parameters in the MEAM library file correspond to single-element potential parameters:

```
lat      = lattice structure of reference configuration
z        = number of nearest neighbors in the reference structure
ielement = atomic number
atwt     = atomic weight
alat     = lattice constant of reference structure
esub     = energy per atom (eV) in the reference structure at equilibrium
asub     = "A" parameter for MEAM (see e.g. (Baskes))
```

The *alpha*, *b0*, *b1*, *b2*, *b3*, *t0*, *t1*, *t2*, *t3* parameters correspond to the standard MEAM parameters in the literature (*Baskes*) (the *b* parameters are the standard beta parameters). Note that only parameters normalized to *t0* = 1.0 are supported. The *rozero* parameter is an element-dependent density scaling that weights the reference background density (see e.g. equation 4.5 in (*Gullet*)) and is typically 1.0 for single-element systems. The *ibar* parameter selects the form of the function *G*(*Gamma*) used to compute the electron density; options are

```
0 => G = sqrt(1+Gamma)
1 => G = exp(Gamma/2)
2 => not implemented
3 => G = 2/(1+exp(-Gamma))
4 => G = sqrt(1+Gamma)
-5 => G = +-sqrt(abs(1+Gamma))
```

If used, the MEAM parameter file contains settings that override or complement the library file settings. Examples of such parameter files are in the potentials directory with a “.meam” suffix. Their format is the same as is read by other Fortran MD codes. Aside from blank and comment lines (start with # which can appear anywhere), each line has one of the following forms. Each line can also have a trailing comment (starting with #) which is ignored.

```
keyword = value
keyword(I) = value
keyword(I,J) = value
keyword(I,J,K) = value
```

The indices I, J, K correspond to the elements selected from the MEAM library file numbered in the order of how those elements were selected starting from 1. Thus for the example given before

```
pair_coeff * * library.meam Si C sic.meam Si Si Si C
```

an index of 1 would refer to Si and an index of 2 to C.

The recognized keywords for the parameter file are as follows:

```
rc          = cutoff radius for cutoff function; default = 4.0
delr        = length of smoothing distance for cutoff function; default = 0.1
rho0(I)     = relative density for element I (overwrites value
              read from meamf file)
Ec(I,J)     = cohesive energy of reference structure for I-J mixture
delta(I,J)  = heat of formation for I-J alloy; if Ec_IJ is input as
              zero, then LAMMPS sets Ec_IJ = (Ec_II + Ec_JJ)/2 - delta_IJ
alpha(I,J)  = alpha parameter for pair potential between I and J (can
              be computed from bulk modulus of reference structure)
re(I,J)     = equilibrium distance between I and J in the reference
              structure
```

`Cmax(I,J,K)` = Cmax screening parameter when I-J pair is screened  
 by K ( $I \leq J$ ); default = 2.8  
`Cmin(I,J,K)` = Cmin screening parameter when I-J pair is screened  
 by K ( $I \leq J$ ); default = 2.0  
`lattce(I,J)` = lattice structure of I-J reference structure:  
     fcc = face centered cubic  
     bcc = body centered cubic  
     hcp = hexagonal close-packed  
     dim = dimer  
     dia = diamond (interlaced fcc for alloy)  
     dia3 = diamond structure with primary 1NN and secondary 3NN interaction  
     b1 = rock salt (NaCl structure)  
     c11 = MoSi2 structure  
     l12 = Cu3Au structure (lower case L, followed by 12)  
     b2 = CsCl structure (interpenetrating simple cubic)  
     ch4 = methane-like structure, only for binary system  
     lin = linear structure (180 degree angle)  
     zig = zigzag structure with a uniform angle  
     tri = H2O-like structure that has an angle  
     sc = simple cubic  
`mn2(I,J)` = turn on second-nearest neighbor MEAM formulation for  
 I-J pair (see for example [Lee](#)).  
     0 = second-nearest neighbor formulation off  
     1 = second-nearest neighbor formulation on  
     default = 0  
`attrac(I,J)` = additional cubic attraction term in Rose energy I-J pair potential  
     default = 0  
`repuls(I,J)` = additional cubic repulsive term in Rose energy I-J pair potential  
     default = 0  
`zbl(I,J)` = blend the MEAM I-J pair potential with the ZBL potential for small  
 atom separations ([ZBL](#))  
     default = 1  
`theta(I,J)` = angle between three atoms in line, zigzag, and trimer reference structures,  
 ↪ in degrees  
     default = 180  
`gsmooth_factor` = factor determining the length of the G-function smoothing  
 region; only significant for `ibar=0` or `ibar=4`.  
     99.0 = short smoothing region, sharp step  
     0.5 = long smoothing region, smooth step  
     default = 99.0  
`augt1` = integer flag for whether to augment t1 parameter by  
 $3/5 * t3$  to account for old vs. new meam formulations;  
     0 = don't augment t1  
     1 = augment t1  
     default = 1  
`ialloy` = integer flag to use alternative averaging rule for t parameters,  
 for comparison with the DYNAMO MEAM code  
     0 = standard averaging (matches `ialloy=0` in DYNAMO)  
     1 = alternative averaging (matches `ialloy=1` in DYNAMO)  
     2 = no averaging of t (use single-element values)  
     default = 0  
`mixture_ref_t` = integer flag to use mixture average of t to compute the background  
 reference density for alloys, instead of the single-element values  
 (see description and warning elsewhere in this doc page)

```
0 = do not use mixture averaging for t in the reference density
1 = use mixture averaging for t in the reference density
default = 0
erose_form = integer value to select the form of the Rose energy function
              (see description below).
              default = 0
emb_lin_neg = integer value to select embedding function for negative densities
              0 = F(rho)=0
              1 = F(rho) = -asub*esub*rho (linear in rho, matches DYNAMO)
              default = 0
bkgd_dyn    = integer value to select background density formula
              0 = rho_bkgd = rho_ref_meam(a) (as in the reference structure)
              1 = rho_bkgd = rho0_meam(a)*Z_meam(a) (matches DYNAMO)
              default = 0
```

*Rc*, *delr*, *re* are in distance units (Angstroms in the case of metal units). *Ec* and *delta* are in energy units (eV in the case of metal units).

Each keyword represents a quantity which is either a scalar, vector, 2d array, or 3d array and must be specified with the correct corresponding array syntax. The indices I,J,K each run from 1 to N where N is the number of MEAM elements being used.

Thus these lines

```
rho0(2) = 2.25
alpha(1,2) = 4.37
```

set *rho0* for the second element to the value 2.25 and set *alpha* for the alloy interaction between elements 1 and 2 to 4.37.

The *augt1* parameter is related to modifications in the MEAM formulation of the partial electron density function. In recent literature, an extra term is included in the expression for the third-order density in order to make the densities orthogonal (see for example (*Wang*), equation 3d); this term is included in the MEAM implementation in lammps. However, in earlier published work this term was not included when deriving parameters, including most of those provided in the `library.meam` file included with lammps, and to account for this difference the parameter *t1* must be augmented by  $3/5*t^3$ . If *augt1* = 1, the default, this augmentation is done automatically. When parameter values are fit using the modified density function, as in more recent literature, *augt1* should be set to 0.

The *mixture\_ref\_t* parameter is available to match results with those of previous versions of lammps (before January 2011). Newer versions of lammps, by default, use the single-element values of the *t* parameters to compute the background reference density. This is the proper way to compute these parameters. Earlier versions of lammps used an alloy mixture averaged value of *t* to compute the background reference density. Setting *mixture\_ref\_t* = 1 gives the old behavior. WARNING: using *mixture\_ref\_t* = 1 will give results that are demonstrably incorrect for second-neighbor MEAM, and non-standard for first-neighbor MEAM; this option is included only for matching with previous versions of lammps and should be avoided if possible.

The parameters *attrac* and *repuls*, along with the integer selection parameter *erose\_form*, can be used to modify the Rose energy function used to compute the pair potential. This function gives the energy of the reference state as a function of interatomic spacing. The form of this function is:

```
astar = alpha * (r/re - 1.d0)
if erose_form = 0: erose = -Ec*(1+astar+a3*(astar**3)/(r/re))*exp(-astar)
if erose_form = 1: erose = -Ec*(1+astar+(-attrac+repuls/r)*(astar**3))*exp(-astar)
if erose_form = 2: erose = -Ec*(1 +astar + a3*(astar**3))*exp(-astar)
a3 = repuls, astar < 0
a3 = attrac, astar >= 0
```

Most published MEAM parameter sets use the default values  $attrac = repulse = 0$ . Setting  $repuls = attrac = delta$  corresponds to the form used in several recent published MEAM parameter sets, such as (*Valone*)

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

**Note:** The default form of the *erose* expression in LAMMPS was corrected in March 2009. The current version is correct, but may show different behavior compared with earlier versions of lammmps with the *attrac* and/or *repuls* parameters are non-zero. To obtain the previous default form, use *erose\_form = 1* (this form does not seem to appear in the literature). An alternative form (see e.g. (*Lee2*)) is available using *erose\_form = 2*.

---

#### 19.167.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , where types I and J correspond to two different element types, mixing is performed by LAMMPS with user-specifiable parameters as described above.

This pair style does not support the *pair\_modify shift, table, and tail* options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner, middle, outer* keywords.

---

#### 19.167.5 Restrictions

The *meam* style is provided in the MEAM package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The maximum number of elements, that can be read from the MEAM library file, is determined at compile time. The default is 5. If you need support for more elements, you have to change the define for the constant 'maxelt' at the beginning of the file *src/MEAM/meam.h* and update/recompile LAMMPS. There is no limit on the number of atoms types.

## 19.167.6 Related commands

*pair\_coeff*, *pair\_style eam*, *pair\_style meam/spline*

## 19.167.7 Default

none

---

**(Baskes)** Baskes, Phys Rev B, 46, 2727-2742 (1992).

**(Gullet)** Gullet, Wagner, Slepoy, SANDIA Report 2003-8782 (2003). This report may be accessed on-line via [this link](#). **(Lee)** Lee, Baskes, Phys. Rev. B, 62, 8564-8567 (2000).

**(Lee2)** Lee, Baskes, Kim, Cho. Phys. Rev. B, 64, 184102 (2001).

**(Valone)** Valone, Baskes, Martin, Phys. Rev. B, 73, 214209 (2006).

**(Wang)** Wang, Van Hove, Ross, Baskes, J. Chem. Phys., 121, 5410 (2004).

**(ZBL)** J.F. Ziegler, J.P. Biersack, U. Littmark, "Stopping and Ranges of Ions in Matter", Vol 1, 1985, Pergamon Press.

## 19.168 pair\_style meam/spline command

Accelerator Variants: *meam/spline/omp*

### 19.168.1 Syntax

```
pair_style meam/spline
```

### 19.168.2 Examples

```
pair_style meam/spline
pair_coeff * * Ti.meam.spline Ti
pair_coeff * * Ti.meam.spline Ti 0
```

### 19.168.3 Description

The *meam/spline* style computes pairwise interactions for metals using a variant of modified embedded-atom method (MEAM) potentials (*Lenosky*). For a single species ("old-style") MEAM, the total energy  $E$  is given by

$$E = \sum_{i < j} \phi(r_{ij}) + \sum_i U(n_i)$$
$$n_i = \sum_j \rho(r_{ij}) + \sum_{\substack{j < k \\ j, k \neq i}} f(r_{ij})f(r_{ik})g[\cos(\theta_{jik})]$$

where  $\rho_i$  is the density at atom I,  $\theta_{jik}$  is the angle between atoms J, I, and K centered on atom I. The five functions  $\phi$ ,  $U$ ,  $\rho$ ,  $f$ , and  $g$  are represented by cubic splines.

The *meam/spline* style also supports a new style multicomponent modified embedded-atom method (MEAM) potential (Zhang), where the total energy  $E$  is given by

$$E = \sum_{i < j} \phi_{ij}(r_{ij}) + \sum_i U_i(n_i)$$

$$n_i = \sum_{j \neq i} \rho_j(r_{ij}) + \sum_{\substack{j < k, \\ j, k \neq i}} f_j(r_{ij}) f_k(r_{ik}) g_{jk}[\cos(\theta_{jik})]$$

where the five functions  $\phi$ ,  $U$ ,  $\rho$ ,  $f$ , and  $g$  depend on the chemistry of the atoms in the interaction. In particular, if there are  $N$  different chemistries, there are  $N$  different  $U$ ,  $\rho$ , and  $f$  functions, while there are  $N(N+1)/2$  different  $\phi$  and  $g$  functions. The new style multicomponent MEAM potential files are indicated by the second line in the file starts with “meam/spline” followed by the number of elements and the name of each element.

The cutoffs and the coefficients for these spline functions are listed in a parameter file which is specified by the *pair\_coeff* command. Parameter files for different elements are included in the “potentials” directory of the LAMMPS distribution and have a “.meam.spline” file suffix. All of these files are parameterized in terms of LAMMPS *metal units*.

Note that unlike for other potentials, cutoffs for spline-based MEAM potentials are not set in the *pair\_style* or *pair\_coeff* command; they are specified in the potential files themselves.

Unlike the EAM pair style, which retrieves the atomic mass from the potential file, the spline-based MEAM potentials do not include mass information; thus you need to use the *mass* command to specify it.

Only a single *pair\_coeff* command is used with the *meam/spline* style which specifies a potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying  $N$  additional arguments after the filename in the *pair\_coeff* command, where  $N$  is the number of LAMMPS atom types:

- filename
- $N$  element names = mapping of spline-based MEAM elements to atom types

See the *pair\_coeff* page for alternate ways to specify the path for the potential file.

As an example, imagine the *Ti.meam.spline* file has values for Ti (old style). In that case your LAMMPS simulation may only have one atom type which has to be mapped to the Ti element as follows:

```
pair_coeff * * Ti.meam.spline Ti
```

The first 2 arguments must be \* \* and there may be only one element following or NULL. Systems where there would be multiple atom types assigned to the same element are **not** supported by this pair style due to limitations in its implementation. If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *meam/spline* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

An example with a two component spline (new style) is *TiO.meam.spline*, where the command

```
pair_coeff * * TiO.meam.spline Ti O
```

will map the first atom type to Ti and the second atom type to O. Note in this case that the species names need to match exactly with the names of the elements in the *TiO.meam.spline* file; otherwise an error will be raised. This behavior is different than the old style MEAM files.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.168.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* shift, table, and tail options.

The *meam/spline* pair style does not write its information to *binary restart files*, since it is stored in an external potential parameter file. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

The *meam/spline* pair style can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

---

#### 19.168.5 Restrictions

This pair style requires the *newton* setting to be “on” for pair interactions.

This pair style does not support mapping multiple atom types to the same element.

This pair style is only enabled if LAMMPS was built with the MANYBODY package. See the *Build package* page for more info.

#### 19.168.6 Related commands

*pair\_coeff*, *pair\_style meam*

#### 19.168.7 Default

none

---

**(Lenosky)** Lenosky, Sadigh, Alonso, Bulatov, de la Rubia, Kim, Voter, Kress, Modelling Simulation Materials Science Engineering, 8, 825 (2000).

**(Zhang)** Zhang and Trinkle, Computational Materials Science, 124, 204-210 (2016).



## 19.169 pair\_style meam/sw/spline command

### 19.169.1 Syntax

```
pair_style meam/sw/spline
```

### 19.169.2 Examples

```
pair_style meam/sw/spline
pair_coeff * * Ti.meam.sw.spline Ti
pair_coeff * * Ti.meam.sw.spline Ti Ti Ti
```

### 19.169.3 Description

The *meam/sw/spline* style computes pairwise interactions for metals using a variant of modified embedded-atom method (MEAM) potentials (*Lenosky*) with an additional Stillinger-Weber (SW) term (*Stillinger*) in the energy. This form of the potential was first proposed by Nicklas, Fellingner, and Park (*Nicklas*). We refer to it as MEAM+SW. The total energy  $E$  is given by

$$\begin{aligned}
 E &= E_{MEAM} + E_{SW} \\
 E_{MEAM} &= \sum_{IJ} \phi(r_{IJ}) + \sum_I U(\rho_I) \\
 E_{SW} &= \sum_I \sum_{JK} F(r_{IJ}) F(r_{IK}) G(\cos(\theta_{JIK})) \\
 \rho_I &= \sum_J \rho(r_{IJ}) + \sum_{JK} f(r_{IJ}) f(r_{IK}) g(\cos(\theta_{JIK}))
 \end{aligned}$$

where  $\rho_I$  is the density at atom  $I$ ,  $\theta_{JIK}$  is the angle between atoms  $J$ ,  $I$ , and  $K$  centered on atom  $I$ . The seven functions  $\phi$ ,  $F$ ,  $G$ ,  $U$ ,  $\rho$ ,  $f$ , and  $g$  are represented by cubic splines.

The cutoffs and the coefficients for these spline functions are listed in a parameter file which is specified by the *pair\_coeff* command. Parameter files for different elements are included in the “potentials” directory of the LAMMPS distribution and have a “.meam.sw.spline” file suffix. All of these files are parameterized in terms of LAMMPS *metal units*.

Note that unlike for other potentials, cutoffs for spline-based MEAM+SW potentials are not set in the *pair\_style* or *pair\_coeff* command; they are specified in the potential files themselves.

Unlike the EAM pair style, which retrieves the atomic mass from the potential file, the spline-based MEAM+SW potentials do not include mass information; thus you need to use the *mass* command to specify it.

Only a single *pair\_coeff* command is used with the *meam/sw/spline* style which specifies a potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying  $N$  additional arguments after the filename in the *pair\_coeff* command, where  $N$  is the number of LAMMPS atom types:

- filename
- $N$  element names = mapping of spline-based MEAM+SW elements to atom types

See the *pair\_coeff* page for alternate ways to specify the path for the potential file.

As an example, imagine the *Ti.meam.sw.spline* file has values for *Ti*. If your LAMMPS simulation has 3 atoms types and they are all to be treated with this potential, you would use the following *pair\_coeff* command:

```
pair_coeff * * Ti meam.sw.spline Ti Ti Ti
```

The first 2 arguments must be \* \* so as to span all LAMMPS atom types. The three Ti arguments map LAMMPS atom types 1,2,3 to the Ti element in the potential file. If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *meam/sw/spline* potential is used as part of the hybrid pair style. The NULL values are placeholders for atom types that will be used with other potentials.

---

**Note:** The *meam/sw/spline* style currently supports only single-element MEAM+SW potentials. It may be extended for alloy systems in the future.

---

Example input scripts that use this pair style are provided in the `examples/PACKAGES/meam_sw_spline` directory.

---

### 19.169.4 Mixing, shift, table, tail correction, restart, rRESPA info

The pair style does not support multiple element types or mixing. It has been designed for pure elements only.

This pair style does not support the *pair\_modify* shift, table, and tail options.

The *meam/sw/spline* pair style does not write its information to *binary restart files*, since it is stored in an external potential parameter file. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

The *meam/sw/spline* pair style can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

---

### 19.169.5 Restrictions

This pair style requires the *newton* setting to be “on” for pair interactions.

This pair style is only enabled if LAMMPS was built with the MANYBODY package. See the *Build package* page for more info.

### 19.169.6 Related commands

*pair\_coeff*, *pair\_style meam*, *pair\_style meam/spline*

### 19.169.7 Default

none

---

**(Lenosky)** Lenosky, Sadigh, Alonso, Bulatov, de la Rubia, Kim, Voter, Kress, Modell. Simul. Mater. Sci. Eng. 8, 825 (2000).

**(Stillinger)** Stillinger, Weber, Phys. Rev. B 31, 5262 (1985).

**(Nicklas)** The spline-based MEAM+SW format was first devised and used to develop potentials for bcc transition metals by Jeremy Nicklas, Michael Fellinger, and Hyoungki Park at The Ohio State University.

---

## 19.170 pair\_style mesocnt command

## 19.171 pair\_style mesocnt/viscous command

### 19.171.1 Syntax

```
pair_style style neigh_cutoff mode neigh_mode
```

- style = *mesocnt* or *mesocnt/viscous*
- neigh\_cutoff = neighbor list cutoff (distance units)
- mode = *chain* or *segment* (optional)
- neigh\_mode = *id* or *topology* (optional)

### 19.171.2 Examples

```
pair_style mesocnt 30.0
pair_coeff * * C_10_10.mesocnt 2

pair_style mesocnt/viscous 60.0 chain topology
pair_coeff * * C_10_10.mesocnt 0.001 20.0 0.2 2 4
```

### 19.171.3 Description

Style *mesocnt* implements a mesoscopic potential for the interaction of carbon nanotubes (CNTs), or other quasi-1D objects such as other kinds of nanotubes or nanowires. In this potential, CNTs are modelled as chains of cylindrical segments in which each infinitesimal surface element interacts with all other CNT surface elements with the Lennard-Jones (LJ) term adopted from the *airebo* style. The interaction energy is then computed by integrating over the surfaces of all interacting CNTs.

In LAMMPS, cylindrical segments are represented by bonds. Each segment is defined by its two end points (“nodes”) which correspond to atoms in LAMMPS. For the exact functional form of the potential and implementation details, the reader is referred to the original papers (*Volkov1*) and (*Volkov2*).

Changed in version 15Sep2022.

The potential supports two modes, *segment* and *chain*. By default, *chain* mode is enabled. In *segment* mode, interactions are pair-wise between all neighboring segments based on a segment-segment approach (keyword *segment* in *pair\_style* command). In *chain* mode, interactions are calculated between each segment and infinitely or semi-infinitely long CNTs as described in (*Volkov1*). Chains of segments are converted to these (semi-)infinite CNTs bases on an approximate chain approach outlined in (*Volkov2*). Hence, interactions are calculated on a segment-chain basis (keyword *chain* in the *pair\_style* command). Using *chain* mode allows to simplify the computation of the interactions significantly and reduces the computational times to the same order of magnitude as for regular bead spring models where beads interact with the standard *pair\_lj/cut* potential. However, this method is only valid when the curvature of the CNTs in the system is small. When CNTs are buckled (see *angle\_mesocnt*), local curvature can be very high and the *pair\_style* automatically switches to *segment* mode for interactions involving buckled CNTs.

The potential further implements two different neighbor list construction modes. Mode *id* uses atom and mol IDs to construct neighbor lists while *topology* modes uses only the bond topology of the system. While *id* mode requires bonded atoms to have consecutive LAMMPS atom IDs and atoms in different CNTs to have different LAMMPS molecule IDs, *topology* mode has no such requirement. Using *id* mode is faster and is enabled by default.

**Note:** Neighbor *id* mode requires all CNTs in the system to have distinct LAMMPS molecule IDs and bonded atoms to have consecutive LAMMPS atom IDs. If this is not possible (e.g. in simulations of CNT rings), *topology* mode needs to be enabled in the `pair_style` command.

---

New in version 15Sep2022.

In addition to the LJ interactions described above, style *mesocnt/viscous* explicitly models friction between neighboring segments. Friction forces are a function of the relative velocity between a segment and its neighboring approximate chain (even in *segment* mode) and only act along the axes of the interacting segment and chain. In this potential, friction forces acting per unit length of a nanotube segment are modelled as a shifted logistic function:

$$F^{\text{FRICTION}}(v)/L = \frac{F^{\text{max}}}{1 + \exp(-k(v - v_0))} - \frac{F^{\text{max}}}{1 + \exp(kv_0)}$$

---

In the `pair_style` command, the modes described above can be toggled using the *segment* or *chain* keywords. The neighbor list cutoff defines the cutoff within which atoms are included in the neighbor list for constructing neighboring CNT chains. This is different from the potential cutoff, which is directly calculated from parameters specified in the potential file. We recommend using a neighbor list cutoff of at least 3 times the maximum segment length used in the simulation to ensure proper neighbor chain construction.

---

**Note:** CNT ends are treated differently by all *mesocnt* styles. Atoms on CNT ends need to be assigned different LAMMPS atom types than atoms not on CNT ends.

---

Style *mesocnt* requires tabulated data provided in a single ASCII text file, as well as a list of integers corresponding to all LAMMPS atom types representing CNT ends:

- filename
- *N* CNT end atom types

For example, if your LAMMPS simulation of (10, 10) nanotubes has 4 atom types where atom types 1 and 3 are assigned to 'inner' nodes and atom types 2 and 4 are assigned to CNT end nodes, the `pair_coeff` command would be:

```
pair_coeff * * C_10_10.mesocnt 2 4
```

---

Likewise, style *mesocnt/viscous* also requires the same information as style *mesocnt*, with the addition of 3 parameters for the viscous friction forces as listed above:

- filename
- $F^{\text{max}}$
- $k$
- $v_0$
- *N* CNT end atom types

Using the same example system as with style *mesocnt* with the addition of friction, the `pair_coeff` command is:

```
pair_coeff * * C_10_10.mesocnt 0.03 20.0 0.20 2 4
```

---

Potential files for CNTs can be readily generated using the freely available code provided on

<https://github.com/phankl/cntpot>

Using the same approach, it should also be possible to generate potential files for other 1D systems mentioned above.

---

**Note:** Because of their size, *mesocnt* style potential files are not bundled with LAMMPS. When compiling LAMMPS from source code, the file `C_10_10.mesocnt` should be downloaded separately from [https://download.lammps.org/potentials/C\\_10\\_10.mesocnt](https://download.lammps.org/potentials/C_10_10.mesocnt)

The first line of the potential file provides a time stamp and general information. The second line lists four integers giving the number of data points provided in the subsequent four data tables. The third line lists four floating point numbers: the CNT radius  $R$ , the LJ parameter  $\sigma$  and two numerical parameters  $\delta_1$  and  $\delta_2$ . These four parameters are given in Angstroms. This is followed by four data tables each separated by a single empty line. The first two tables have two columns and list the parameters `uInfParallel` and `Gamma` respectively. The last two tables have three columns giving data on a quadratic array and list the parameters `Phi` and `uSemiParallel` respectively. `uInfParallel` and `uSemiParallel` are given in eV/Angstrom, `Phi` is given in eV and `Gamma` is unitless.

If a simulation produces many warnings about segment-chain interactions falling outside the interpolation range, we recommend generating a potential file with lower values of  $\delta_1$  and  $\delta_2$ .

---

#### 19.171.4 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles does not support mixing.

These pair styles does not support the *pair\_modify* shift, table, and tail options.

These pair styles do not write their information to *binary restart files*, since it is stored in tabulated potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

---

#### 19.171.5 Restrictions

These styles are part of the MESONT package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These pair styles require the *newton* setting to be “on” for pair interactions.

These pair styles require all 3 *special\_bonds lj* settings to be non-zero for proper neighbor list construction.

Pair style *mesocnt/viscous* requires you to use the *comm\_modify vel yes* command so that velocities are stored by ghost atoms.

### 19.171.6 Related commands

*pair\_coeff*

### 19.171.7 Default

mode = chain, neigh\_mode = id

---

(Volkov1) Volkov and Zhigilei, J Phys Chem C, 114, 5513 (2010).

(Volkov2) Volkov, Simov and Zhigilei, APS Meeting Abstracts, Q31.013 (2008).

## 19.172 pair\_style edpd command

## 19.173 pair\_style mdpd command

## 19.174 pair\_style mdpd/rhosum command

## 19.175 pair\_style tdpd command

### 19.175.1 Syntax

<code>pair_style</code> style args
------------------------------------

- style = *edpd* or *mdp*d or *mdp/rhosum* or *tdpd*
- args = list of arguments for a particular style

*edpd* args = cutoff seed

cutoff = global cutoff for eDPD interactions (distance units)

seed = random # seed (integer) (if <= 0, eDPD will use current time as the seed)

*mdp*d args = T cutoff seed

T = temperature (temperature units)

cutoff = global cutoff for mDPD interactions (distance units)

seed = random # seed (integer) (if <= 0, mDPD will use current time as the seed)

*mdp*d/*rhosum* args =

*tdpd* args = T cutoff seed

T = temperature (temperature units)

cutoff = global cutoff for tDPD interactions (distance units)

seed = random # seed (integer) (if <= 0, tDPD will use current time as the seed)

## 19.175.2 Examples

```

pair_style edpd 1.58 9872598
pair_coeff * * 18.75 4.5 0.41 1.58 1.42E-5 2.0 1.58
pair_coeff 1 1 18.75 4.5 0.41 1.58 1.42E-5 2.0 1.58 power 10.54 -3.66 3.44 -4.10
pair_coeff 1 1 18.75 4.5 0.41 1.58 1.42E-5 2.0 1.58 power 10.54 -3.66 3.44 -4.10 kappa -
→0.44 -3.21 5.04 0.00

pair_style hybrid/overlay mdpd/rhosum mdpd 1.0 1.0 65689
pair_coeff 1 1 mdpd/rhosum 0.75
pair_coeff 1 1 mdpd -40.0 25.0 18.0 1.0 0.75

pair_style tdpd 1.0 1.58 935662
pair_coeff * * 18.75 4.5 0.41 1.58 1.58 1.0 1.0E-5 2.0
pair_coeff 1 1 18.75 4.5 0.41 1.58 1.58 1.0 1.0E-5 2.0 3.0 1.0E-5 2.0

```

## 19.175.3 Description

The *edpd* style computes the pairwise interactions and heat fluxes for eDPD particles following the formulations in (*Li2014\_JCP*) and *Li2015\_CC*. The time evolution of an eDPD particle is governed by the conservation of momentum and energy given by

$$\frac{d^2\mathbf{r}_i}{dt^2} = \frac{d\mathbf{v}_i}{dt} = \mathbf{F}_i = \sum_{i \neq j} (\mathbf{F}_{ij}^C + \mathbf{F}_{ij}^D + \mathbf{F}_{ij}^R)$$

$$C_v \frac{dT_i}{dt} = q_i = \sum_{i \neq j} (q_{ij}^C + q_{ij}^V + q_{ij}^R),$$

where the three components of  $F_i$  including the conservative force  $F_{ij}^C$ , dissipative force  $F_{ij}^D$  and random force  $F_{ij}^R$  are expressed as

$$\mathbf{F}_{ij}^C = \alpha_{ij} \omega_C(r_{ij}) \mathbf{e}_{ij}$$

$$\mathbf{F}_{ij}^D = -\gamma \omega_D(r_{ij}) (\mathbf{e}_{ij} \cdot \mathbf{v}_{ij}) \mathbf{e}_{ij}$$

$$\mathbf{F}_{ij}^R = \sigma \omega_R(r_{ij}) \xi_{ij} \Delta t^{-1/2} \mathbf{e}_{ij}$$

$$\omega_C(r) = 1 - r/r_c$$

$$\alpha_{ij} = A \cdot k_B (T_i + T_j) / 2$$

$$\omega_D(r) = \omega_R^2(r) = (1 - r/r_c)^s$$

$$\sigma_{ij}^2 = 4\gamma k_B T_i T_j / (T_i + T_j)$$

in which the exponent of the weighting function  $s$  can be defined as a temperature-dependent variable. The heat flux between particles accounting for the collisional heat flux  $q^C$ , viscous heat flux  $q^V$ , and random heat flux  $q^R$  are given

by

$$\begin{aligned}
 q_i^C &= \sum_{j \neq i} k_{ij} \omega_{CT}(r_{ij}) \left( \frac{1}{T_i} - \frac{1}{T_j} \right) \\
 q_i^V &= \frac{1}{2C_v} \sum_{j \neq i} \left\{ \omega_D(r_{ij}) \left[ \gamma_{ij} (\mathbf{e}_{ij} \cdot \mathbf{v}_{ij})^2 - \frac{(\sigma_{ij})^2}{m} \right] - \sigma_{ij} \omega_R(r_{ij}) (\mathbf{e}_{ij} \cdot \mathbf{v}_{ij}) \xi_{ij} \right\} \\
 q_i^R &= \sum_{j \neq i} \beta_{ij} \omega_{RT}(r_{ij}) dt^{-1/2} \xi_{ij}^e \\
 \omega_{CT}(r) &= \omega_{RT}^2(r) = (1 - r/r_{ct})^{s_T} \\
 k_{ij} &= C_v^2 \kappa (T_i + T_j)^2 / 4k_B \\
 \beta_{ij}^2 &= 2k_B k_{ij}
 \end{aligned}$$

where the mesoscopic heat friction  $\kappa$  is given by

$$\kappa = \frac{315k_B v}{2\pi \rho C_v r_{ct}^5 Pr},$$

with  $v$  being the kinematic viscosity. For more details, see Eq.(15) in (Li2014\_JCP).

The following coefficients must be defined in eDPD system for each pair of atom types via the *pair\_coeff* command as in the examples above.

- A (force units)
- $\gamma$  (force/velocity units)
- power\_f (positive real)
- cutoff (distance units)
- kappa (thermal conductivity units)
- power\_T (positive real)
- cutoff\_T (distance units)
- optional keyword = power or kappa

The keyword *power* or *kappa* is optional. Both “power” and “kappa” require 4 parameters  $c_1, c_2, c_3, c_4$  showing the temperature dependence of the exponent  $s(T) = \text{power}_f(1 + c_1(T - 1) + c_2(T - 1)^2 + c_3(T - 1)^3 + c_4(T - 1)^4)$  and of the mesoscopic heat friction  $s_T(T) = \kappa(1 + c_1(T - 1) + c_2(T - 1)^2 + c_3(T - 1)^3 + c_4(T - 1)^4)$ . If the keyword *power* or *kappa* is not specified, the eDPD system will use constant power\_f and  $\kappa$ , which is independent to temperature changes.

---

The *mdpd/rhosum* style computes the local particle mass density  $\rho$  for mDPD particles by kernel function interpolation.

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above.

- cutoff (distance units)
- 

The *mdpd* style computes the many-body interactions between mDPD particles following the formulations in (Li2013\_POF). The dissipative and random forces are in the form same as the classical DPD, but the conservative force is local density dependent, which are given by

$$\begin{aligned}
 \mathbf{F}_{ij}^C &= A w_c(r_{ij}) \mathbf{e}_{ij} + B(\rho_i + \rho_j) w_d(r_{ij}) \mathbf{e}_{ij} \\
 \mathbf{F}_{ij}^D &= -\gamma \omega_D(r_{ij}) (\mathbf{e}_{ij} \cdot \mathbf{v}_{ij}) \mathbf{e}_{ij} \\
 \mathbf{F}_{ij}^R &= \sigma \omega_R(r_{ij}) \xi_{ij} \Delta t^{-1/2} \mathbf{e}_{ij}
 \end{aligned}$$



where the first term in  $F_C$  with a negative coefficient  $A < 0$  stands for an attractive force within an interaction range  $r_c$ , and the second term with  $B > 0$  is the density-dependent repulsive force within an interaction range  $r_d$ .

The following coefficients must be defined for each pair of atom types via the `pair_coeff` command as in the examples above.

- A (force units)
- B (force units)
- $\gamma$  (force/velocity units)
- cutoff\_c (distance units)
- cutoff\_d (distance units)

The `tdpd` style computes the pairwise interactions and chemical concentration fluxes for tDPD particles following the formulations in ([Li2015\\_JCP](#)). The time evolution of a tDPD particle is governed by the conservation of momentum and concentration given by

$$\frac{d^2\mathbf{r}_i}{dt^2} = \frac{d\mathbf{v}_i}{dt} = \mathbf{F}_i = \sum_{i \neq j} (\mathbf{F}_{ij}^C + \mathbf{F}_{ij}^D + \mathbf{F}_{ij}^R)$$

$$\frac{dC_i}{dt} = Q_i = \sum_{i \neq j} (Q_{ij}^D + Q_{ij}^R) + Q_i^S$$

where the three components of  $F_i$  including the conservative force  $F_{ij}^C$ , dissipative force  $F_{ij}^D$  and random force  $F_{ij}^R$  are expressed as

$$\mathbf{F}_{ij}^C = A\omega_C(r_{ij})\mathbf{e}_{ij}$$

$$\mathbf{F}_{ij}^D = -\gamma\omega_D(r_{ij})(\mathbf{e}_{ij} \cdot \mathbf{v}_{ij})\mathbf{e}_{ij}$$

$$\mathbf{F}_{ij}^R = \sigma\omega_R(r_{ij})\xi_{ij}\Delta t^{-1/2}\mathbf{e}_{ij}$$

$$\omega_C(r) = 1 - r/r_c$$

$$\omega_D(r) = \omega_R^2(r) = (1 - r/r_c)^{\text{power}_f}$$

$$\sigma^2 = 2\gamma k_B T$$

The concentration flux between two tDPD particles includes the Fickian flux  $Q_{ij}^D$  and random flux  $Q_{ij}^R$ , which are given by

$$Q_{ij}^D = -\kappa_{ij}w_{DC}(r_{ij})(C_i - C_j)$$

$$Q_{ij}^R = \epsilon_{ij}(C_i + C_j)w_{RC}(r_{ij})\xi_{ij}$$

$$w_{DC}(r_{ij}) = w_{RC}^2(r_{ij}) = (1 - r/r_{cc})^{\text{power}_{cc}}$$

$$\epsilon_{ij}^2 = m_s^2\kappa_{ij}\rho$$

where the parameters kappa and epsilon determine the strength of the Fickian and random fluxes.  $m_s$  is the mass of a single solute molecule. In general,  $m_s$  is much smaller than the mass of a tDPD particle  $m$ . For more details, see ([Li2015\\_JCP](#)).

The following coefficients must be defined for each pair of atom types via the `pair_coeff` command as in the examples above.

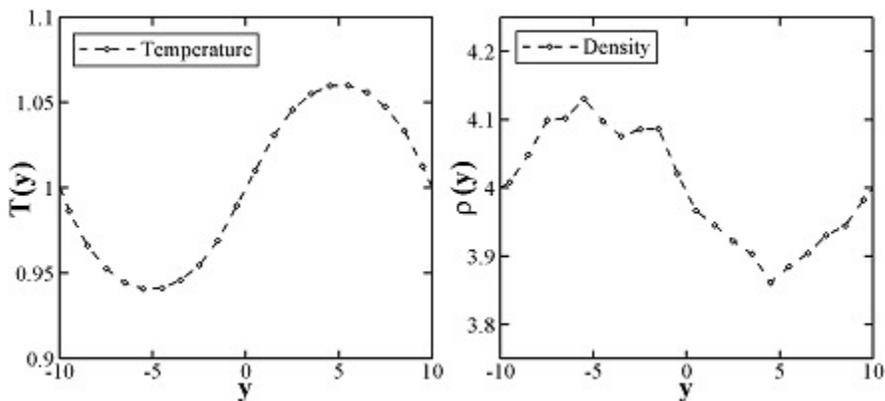
- A (force units)
- $\gamma$  (force/velocity units)
- power\_f (positive real)

- cutoff (distance units)
- cutoff\_CC (distance units)
- $\kappa_i$  (diffusivity units)
- $\epsilon_i$  (diffusivity units)
- power\_cc\_i (positive real)

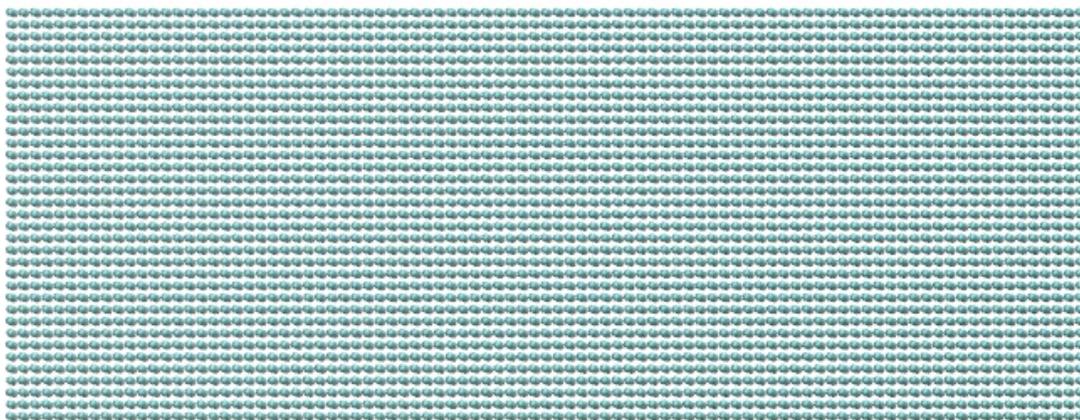
The last 3 values must be repeated Nspecies times, so that values for each of the Nspecies chemical species are specified, as indicated by the “I” suffix. In the first pair\_coeff example above for pair\_style tdpd, Nspecies = 1. In the second example, Nspecies = 2, so 3 additional coeffs are specified (for species 2).

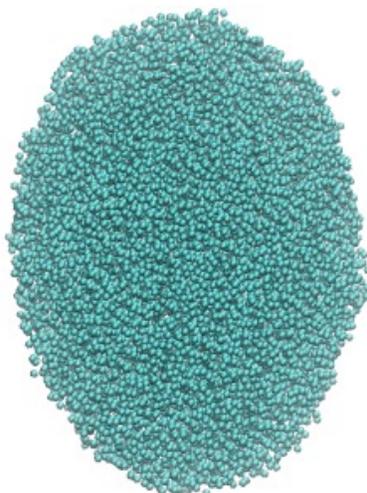
### 19.175.4 Example scripts

There are example scripts for using all these pair styles in examples/PACKAGES/mesodpd. The example for an eDPD simulation models heat conduction with source terms analog of periodic Poiseuille flow problem. The setup follows Fig.12 in (*Li2014\_JCP*). The output of the short eDPD simulation (about 2 minutes on a single core) gives a temperature and density profiles as



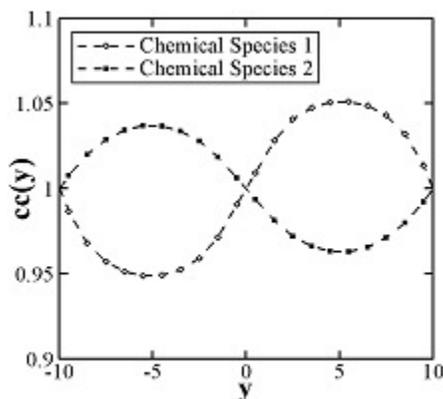
The example for a mDPD simulation models the oscillations of a liquid droplet started from a liquid film. The mDPD parameters are adopted from (*Li2013\_POF*). The short mDPD run (about 2 minutes on a single core) generates a particle trajectory which can be visualized as follows.





The first image is the initial state of the simulation. If you click it a GIF movie should play in your browser. The second image is the final state of the simulation.

The example for a tDPD simulation computes the effective diffusion coefficient of a tDPD system using a method analogous to the periodic Poiseuille flow. The tDPD system is specified with two chemical species, and the setup follows Fig.1 in (*Li2015\_JCP*). The output of the short tDPD simulation (about one and a half minutes on a single core) gives the concentration profiles of the two chemical species as



### 19.175.5 Mixing, shift, table, tail correction, restart, rRESPA info

The styles *edpd*, *mdpd*, *mdpd/rhosum* and *tdpd* do not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

The styles *edpd*, *mdpd*, *mdpd/rhosum* and *tdpd* do not support the *pair\_modify* shift, table, and tail options.

The styles *edpd*, *mdpd*, *mdpd/rhosum* and *tdpd* do not write information to *binary restart files*. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

## 19.175.6 Restrictions

The pair styles *edpd*, *mdpd*, *mdpd/rhosum* and *tdpd* are part of the DPD-MESO package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.175.7 Related commands

*pair\_coeff*, *fix mvv/dpd*, *fix mvv/edpd*, *fix mvv/tdpd*, *fix edpd/source*, *fix tdpd/source*, *compute edpd/temp/atom*, *compute tdpd/cc/atom*

## 19.175.8 Default

none

---

**(Li2014\_JCP)** Li, Tang, Lei, Caswell, Karniadakis, J Comput Phys, 265: 113-127 (2014). DOI: 10.1016/j.jcp.2014.02.003.

**(Li2015\_CC)** Li, Tang, Li, Karniadakis, Chem Commun, 51: 11038-11040 (2015). DOI: 10.1039/C5CC01684C.

**(Li2013\_POF)** Li, Hu, Wang, Ma, Zhou, Phys Fluids, 25: 072103 (2013). DOI: 10.1063/1.4812366.

**(Li2015\_JCP)** Li, Yazdani, Tartakovsky, Karniadakis, J Chem Phys, 143: 014101 (2015). DOI: 10.1063/1.4923254.

## 19.176 pair\_style mesont/tpm command

### 19.176.1 Syntax

```
pair_style mesont/tpm cut table_path BendingMode TPMTYPE
```

- cut = the cutoff distance
- table\_path = the path to the potential table
- BendingMode = the parameter defining the type of the bending potential for nanotubes: 0 - harmonic bending (*Srivastava*), 1 - anharmonic potential of bending and bending-buckling (*Zhigilei1*)
- TPMTYPE = the parameter determining the type of the inter-tube interaction term: 0 - segment-segment approach, 1 - segment-chain approach (*Zhigilei2*, *Zhigilei3*)

The segment-segment approach is approximately 5 times slower than segment-chain approximation. The parameter BendingMode also affects the calculation of the inter-tube interaction term when TPMTYPE = 1. In this case, when BendingMode = 1, each continuous chain of segments is additionally replaced by a number of sub-chains divided by bending buckling kinks.

## 19.176.2 Examples

```
pair_style mesont/tpm 30.0 MESONT-TABTP_10_10.xrs 0 0
```

## 19.176.3 Description

The tubular potential model (TPM) force field is designed for mesoscopic simulations of interacting flexible nanotubes. The force field is based on the mesoscopic computational model suggested in Ref. (*Srivastava*). In this model, each nanotube is represented by a chain of mesoscopic elements in the form of stretchable cylindrical segments, where each segment consists of multiple atoms. Each nanotube is divided into segments by a sequence of nodes placed on the nanotube centerline. This sequence of nodes determines the spatial position of the cylindrical segments and defines the configuration of the entire tube.

The potential force field that controls the dynamic behavior of a system of interacting nanotubes is given by the following equation defining the potential energy of the system:

$$U = U_{str} + U_{bnd} + U_{vdW}$$

where  $U_{str}$  is the harmonic potential describing the stretching of nanotube (*Srivastava*),  $U_{bnd}$  is the potential for nanotube bending (*Srivastava*) and bending-buckling (*Zhigilei1*), and  $U_{vdW}$  is the potential describing van-der Waals interaction between nanotubes (*Zhigilei2*, *Zhigilei3*). The stretching energy,  $U_{str}$ , is given by the sum of stretching energies of individual nanotube segments. The bending energy,  $U_{bnd}$ , is given by the sum of bending energies in all internal nanotube nodes. The tube-tube interaction energy,  $U_{vdW}$ , is calculated based on the tubular potential method suggested in Ref. (*Zhigilei2*). The tubular potential method is briefly described below.

The interaction between two straight nanotubes of arbitrary length and orientation is described by the approximate tubular potential developed in (*Zhigilei3*). This potential approximates the results of direct integration of carbon-carbon interatomic potential over the surfaces of the interacting nanotubes, with the force sources homogeneously distributed over the nanotube surfaces. The input data for calculation of tubular potentials are partially tabulated. For single-walled CNTs of arbitrary chirality, the tabulated potential data can be generated in the form of ASCII files TPMSSTP.xrs and TPMA.xrs by the stand-alone code TMDPotGen included in the tool directory of LAMMPS release. The potential provided with LAMMPS release, MESONT-TABTP\_10\_10.xrs, is tabulated for (10,10) nanotubes.

Calculations of the interaction between curved or bent nanotubes are performed on either segment-segment or segment-chain basis. In the first case, activated when parameter TPMTType is equal to 0, the tubular potential is calculated for each pair of interacting mesoscopic segments. In this case, however, small potential barriers for inter-tube sliding are introduced. While relatively small, these barriers are still larger than the ones that originate from the atomic-scale corrugation in atomistic modeling of inter-tube interaction. The latter are too weak to prevent room-temperature rearrangements of defect-free CNT, while the artificial mesoscopic barriers due to the segment-segment interaction can impede sliding of nanotubes with respect to each other and affect the kinetics of structural rearrangements in a system of nanotubes at moderate mesoscopic temperatures. In the second case, activated when parameter TPMTType is equal to 1, the inter-tube interaction term is calculated based on the segment-chain approach. In this case, for each NT segment, the list of its neighboring segments is divided into short continuous chains of segments belonging to individual nanotubes. For each pair of a segment and a chain, the curved chain is approximated by a straight equivalent nanotube based on the weighted approach suggested in Ref. (*Zhigilei2*). Finally, the interaction between the segment and straight equivalent chain is calculated based on the tubular potential. In this case, and in the absence of bending buckling (i.e., when parameter BendingMode is equal to 0), the tubular potential method ensures the absence of corrugation of the effective inter-tube interaction potential for curved nanotubes and eliminates any barriers for the inter-tube sliding. As a result, the tubular potential method can describe the spontaneous self-assembly of nanotubes into continuous networks of bundles (*Zhigilei1*, *Zhigilei3*).

The TMD force field has been used for generation of nanotube films, fibers, and vertically aligned forests of nanotubes. Mesoscopic dynamic simulations were used to prepare realistic structures of continuous networks of nanotube bundles

and to study their structural and mechanical properties (*Zhigilei1*, *Zhigilei3*, *Zhigilei4*, *Zhigilei5*, *Zhigilei6*). With additional models for heat transfer, this force field was also used to study the thermal transport properties of carbon nanotube films (*Zhigilei7*, *Zhigilei8*, *Zhigilei8*). The methods for modeling of the mechanical energy dissipation into heat (energy exchange between the dynamic degrees of freedom of the mesoscopic model and the energy of atomic vibrations that are not explicitly represented in the model) (*Zhigilei10*) and mesoscopic description of covalent cross-links between nanotubes (*Banna*) have also been developed but are not included in this first release of the LAMMPS implementation of the force field. Further details can be found in references provided below.

The MESONT package also provides TMDGen code designed to generate initial samples composed of straight and dispersed nanotubes of given chirality and length at a given material density, which is available in tools directory. In the generated samples, nanotubes are distributed with random positions and orientations. Both periodic and free boundary conditions are available along each axis of the system of coordinates. All parameters in the sample files generated with TMDGen are given in metal *units*.

### 19.176.4 Restrictions

This pair style is a part of the MSEONT package, and it is only enabled if LAMMPS is built with that package. See the *Build package* doc page for more information.

This pair potential requires use of *mesont atomic style*.

This pair potential requires the *newton* setting to be “on” for pair interactions.

The cutoff distance should be set to be at least  $\max \left[ 2L, \sqrt{L^2/2 + (2R + T_{cut})^2} \right]$ , where L is the maximum segment length, R is the maximum tube radius, and  $T_{cut} = 10.2$  Å is the maximum distance between the surfaces of interacting segments. Because of the use of extended chain concept at CNT ends, the recommended cutoff is 3L.

---

**Note:** Because of their size, *mesont* style potential files are not bundled with LAMMPS. When compiling LAMMPS from source code, the file `TABTP_10_10.mesont` should be downloaded transparently from [https://download.lammps.org/potentials/TABTP\\_10\\_10.mesont](https://download.lammps.org/potentials/TABTP_10_10.mesont)

---

The `TABTP_10_10.mesont` potential file is parameterized for metal *units*. You can use the carbon nanotube mesoscopic force field with any LAMMPS units, but you would need to create your own potential files with coefficients listed in appropriate units, if your simulation does not use “metal” units.

The chirality parameters set during system generation must match the values specified during generation of the potential tables.

### 19.176.5 Related commands

*pair\_coeff*

---

(**Srivastava**) Zhigilei, Wei, Srivastava, Phys. Rev. B 71, 165417 (2005).

(**Zhigilei1**) Volkov and Zhigilei, ACS Nano 4, 6187 (2010).

(**Zhigilei2**) Volkov, Simov, Zhigilei, ASME paper IMECE2008, 68021 (2008).

(**Zhigilei3**) Volkov, Zhigilei, J. Phys. Chem. C 114, 5513 (2010).

(**Zhigilei4**) Wittmaack, Banna, Volkov, Zhigilei, Carbon 130, 69 (2018).

(**Zhigilei5**) Wittmaack, Volkov, Zhigilei, Compos. Sci. Technol. 166, 66 (2018).

(**Zhigilei6**) Wittmaack, Volkov, Zhigilei, Carbon 143, 587 (2019).

(Zhgilei7) Volkov, Zhigilei, Phys. Rev. Lett. 104, 215902 (2010).

(Zhgilei8) Volkov, Shiga, Nicholson, Shiomi, Zhigilei, J. Appl. Phys. 111, 053501 (2012).

(Zhgilei9) Volkov, Zhigilei, Appl. Phys. Lett. 101, 043113 (2012).

(Zhgilei10) Jacobs, Nicholson, Zemer, Volkov, Zhigilei, Phys. Rev. B 86, 165414 (2012).

(Banna) Volkov, Banna, Comp. Mater. Sci. 176, 109410 (2020).

## 19.177 pair\_style mgpt command

### 19.177.1 Syntax

```
pair_style mgpt
```

### 19.177.2 Examples

```
pair_style mgpt
pair_coeff * * Ta6.8x.mgpt.parmin Ta6.8x.mgpt.potin Omega
cp ~/lammmps/potentials/Ta6.8x.mgpt.parmin parmin
cp ~/lammmps/potentials/Ta6.8x.mgpt.potin potin
pair_coeff * * parmin potin Omega volpress yes nbody 1234 precision double
pair_coeff * * parmin potin Omega volpress yes nbody 12
```

### 19.177.3 Description

Within DFT quantum mechanics, generalized pseudopotential theory (GPT) (*Moriarty1*) provides a first-principles approach to multi-ion interatomic potentials in d-band transition metals, with a volume-dependent, real-space total-energy functional for the N-ion elemental bulk material in the form

$$E_{\text{tot}}(\mathbf{R}_1 \dots \mathbf{R}_N) = NE_{\text{vol}}(\Omega) + \frac{1}{2} \sum'_{i,j} v_2(ij; \Omega) + \frac{1}{6} \sum'_{i,j,k} v_3(ijk; \Omega) + \frac{1}{24} \sum'_{i,j,k,l} v_4(ijkl; \Omega)$$

where the prime on each summation sign indicates the exclusion of all self-interaction terms from the summation. The leading volume term  $E_{\text{vol}}$  as well as the two-ion central-force pair potential  $v_2$  and the three- and four-ion angular-force potentials,  $v_3$  and  $v_4$ , depend explicitly on the atomic volume  $\Omega$ , but are structure independent and transferable to all bulk ion configurations, either ordered or disordered, and with or without the presence of point and line defects. The simplified model GPT or MGPT (*Moriarty2*, *Moriarty3*), which retains the form of  $E_{\text{tot}}$  and permits more efficient large-scale atomistic simulations, derives from the GPT through a series of systematic approximations applied to  $E_{\text{vol}}$  and the potentials  $v_n$  that are valid for mid-period transition metals with nearly half-filled d bands.

Both analytic (*Moriarty2*) and matrix (*Moriarty3*) representations of MGPT have been developed. In the more general matrix representation, which can also be applied to f-band actinide metals and permits both canonical and non-canonical d/f bands, the multi-ion potentials are evaluated on the fly during a simulation through d- or f-state matrix multiplication, and the forces that move the ions are determined analytically. Fast matrix-MGPT algorithms have been developed independently by Glosli (*Glosli*, *Moriarty3*) and by Opperstrup (*Opperstrup*)

The *mgpt* pair style calculates forces, energies, and the total energy per atom,  $E_{\text{tot}}/N$ , using the Opperstrup matrix-MGPT algorithm. Input potential and control data are entered through the *pair\_coeff* command. Each material treated requires input parmin and potin potential files, as shown in the above examples, as well as specification by the user of the initial atomic volume  $\Omega$  through *pair\_coeff*. At the beginning of a time step in any simulation, the total volume

of the simulation cell  $V$  should always be equal to  $\Omega \cdot N$ , where  $N$  is the number of metal ions present, taking into account the presence of any vacancies and/or interstitials in the case of a solid. In a constant-volume simulation, which is the normal mode of operation for the *mgpt* pair style,  $\Omega$ ,  $V$  and  $N$  all remain constant throughout the simulation and thus are equal to their initial values. In a constant-stress simulation, the cell volume  $V$  will change (slowly) as the simulation proceeds. After each time step, the atomic volume should be updated by the code as  $\Omega = V/N$ . In addition, the volume term  $E_{\text{vol}}$  and the potentials  $v_2$ ,  $v_3$  and  $v_4$  have to be removed at the end of the time step, and then respecified at the new value of  $\Omega$ . In all simulations,  $\Omega$  must remain within the defined volume range for  $E_{\text{vol}}$  and the potentials for the given material.

The default option `volpress yes` in the *pair\_coeff* command includes all volume derivatives of  $E_{\text{tot}}$  required to calculate the stress tensor and pressure correctly. The option `volpress no` disregards the pressure contribution resulting from the volume term  $E_{\text{vol}}$ , and can be used for testing and analysis purposes. The additional optional variable `nbody` controls the specific terms in  $E_{\text{tot}}$  that are calculated. The default option and the normal option for mid-period transition and actinide metals is `nbody 1234` for which all four terms in  $E_{\text{tot}}$  are retained. The option `nbody 12`, for example, retains only the volume term and the two-ion pair potential term and can be used for GPT series-end transition metals that can be well described without  $v_3$  and  $v_4$ . The `nbody` option can also be used to test or analyze the contribution of any of the four terms in  $E_{\text{tot}}$  to a given calculated property.

The *mgpt* pair style makes extensive use of matrix algebra and includes optimized kernels for the BlueGene/Q architecture and the Intel/AMD (x86) architectures. When compiled with the appropriate compiler and compiler switches (`-msse3` on x86, and using the IBM XL compiler on BG/Q), these optimized routines are used automatically. For BG/Q machines, building with the default Makefile for that architecture (e.g., “`make bgq`”) should enable the optimized algebra routines. For x-86 machines, there is a provided `Makefile.mgptfast` which enables the fast algebra routines, i.e. build LAMMPS with “`make mgptfast`”. The user will be informed in the output files of the matrix kernels in use. To further improve speed, on x86 the option `precision single` can be added to the *pair\_coeff* command line, which improves speed (up to a factor of two) at the cost of doing matrix calculations with 7 digit precision instead of the default 16. For consistency the default option can be specified explicitly by the option `precision double`.

All remaining potential and control data are contained with the `parmin` and `potin` files, including cutoffs, atomic mass, and other basic MGPT variables. Specific MGPT potential data for the transition metals tantalum (Ta4 and Ta6.8x potentials), molybdenum (Mo5.2 potentials), and vanadium (V6.1 potentials) are contained in the LAMMPS potentials directory. The stored files are, respectively, `Ta4.mgpt.parmin`, `Ta4.mgpt.potin`, `Ta6.8x.mgpt.parmin`, `Ta6.8x.mgpt.potin`, `Mo5.2.mgpt.parmin`, `Mo5.2.mgpt.potin`, `V6.1.mgpt.parmin`, and `V6.1.mgpt.potin`. Useful corresponding informational “README” files on the Ta4, Ta6.8x, Mo5.2 and V6.1 potentials are also included in the potentials directory. These latter files indicate the volume mesh and range for each potential and give appropriate references for the potentials. It is expected that MGPT potentials for additional materials will be added over time.

Useful example MGPT scripts are given in the `examples/PACKAGES/mgpt` directory. These scripts show the necessary steps to perform constant-volume calculations and simulations. It is strongly recommended that the user work through and understand these examples before proceeding to more complex simulations.

---

**Note:** For good performance, LAMMPS should be built with the compiler flags “`-O3 -msse3 -funroll-loops`” when including this pair style. The `src/MAKE/OPTIONS/Makefile.mgptfast` is an example machine Makefile with these options included as part of a standard MPI build. Note that it as provided, it will build with whatever low-level compiler (g++, icc, etc) is the default for your MPI installation.

---



### 19.177.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* mix, shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.177.5 Restrictions

This pair style is part of the MGPT package and is only enabled if LAMMPS is built with that package. See the *Build package* page for more info.

The MGPT potentials require the *newton* setting to be “on” for pair style interactions.

The stored parmin and potin potential files provided with LAMMPS in the “potentials” directory are written in Rydberg atomic units, with energies in Rydbergs and distances in Bohr radii. The *mgpt* pair style converts Rydbergs to Hartrees to make the potential files compatible with LAMMPS electron *units*.

The form of  $E_{\text{tot}}$  used in the *mgpt* pair style is only appropriate for elemental bulk solids and liquids. This includes solids with point and extended defects such as vacancies, interstitials, grain boundaries and dislocations. Alloys and free surfaces, however, require significant modifications, which are not included in the *mgpt* pair style. Likewise, the *hybrid* pair style is not allowed, where MGPT would be used for some atoms but not for others.

Electron-thermal effects are not included in the standard MGPT potentials provided in the “potentials” directory, where the potentials have been constructed at zero electron temperature. Physically, electron-thermal effects may be important in 3d (e.g., V) and 4d (e.g., Mo) transition metals at high temperatures near melt and above. It is expected that temperature-dependent MGPT potentials for such cases will be added over time.

### 19.177.6 Related commands

*pair\_coeff*

### 19.177.7 Default

The options defaults for the *pair\_coeff* command are volpress yes, nbody 1234, and precision double.

**(Moriarty1)** Moriarty, Physical Review B, 38, 3199 (1988).

**(Moriarty2)** Moriarty, Physical Review B, 42, 1609 (1990). Moriarty, Physical Review B 49, 12431 (1994).

**(Moriarty3)** Moriarty, Benedict, Glosli, Hood, Orlikowski, Patel, Soderlind, Streitz, Tang, and Yang, Journal of Materials Research, 21, 563 (2006).

**(Glosli)** Glosli, unpublished, 2005. Streitz, Glosli, Patel, Chan, Yates, de Supinski, Sexton and Gunnels, Journal of Physics: Conference Series, 46, 254 (2006).

**(Oppelstrup)** Oppelstrup, unpublished, 2015. Oppelstrup and Moriarty, to be published.

## 19.178 pair\_style mie/cut command

Accelerator Variants: *mie/cut/gpu*

### 19.178.1 Syntax

```
pair_style mie/cut cutoff
```

- cutoff = global cutoff for mie/cut interactions (distance units)

### 19.178.2 Examples

```
pair_style mie/cut 10.0
pair_coeff 1 1 0.72 3.40 23.00 6.66
pair_coeff 2 2 0.30 3.55 12.65 6.00
pair_coeff 1 2 0.46 3.32 16.90 6.31
```

### 19.178.3 Description

The *mie/cut* style computes the Mie potential, given by

$$E = C\epsilon \left[ \left( \frac{\sigma}{r} \right)^{\gamma_{rep}} - \left( \frac{\sigma}{r} \right)^{\gamma_{att}} \right] \quad r < r_c$$

$r_c$  is the cutoff and  $C$  is a function that depends on the repulsive and attractive exponents, given by:

$$C = \left( \frac{\gamma_{rep}}{\gamma_{rep} - \gamma_{att}} \right) \left( \frac{\gamma_{rep}}{\gamma_{att}} \right)^{\left( \frac{\gamma_{att}}{\gamma_{rep} - \gamma_{att}} \right)}$$

Note that for 12/6 exponents,  $C$  is equal to 4 and the formula is the same as the standard Lennard-Jones potential.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- epsilon (energy units)
- sigma (distance units)
- gammaR
- gammaA
- cutoff (distance units)

The last coefficient is optional. If not specified, the global cutoff specified in the *pair\_style* command is used.

---

## 19.178.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs  $I, J$  and  $I \neq J$ , the epsilon and sigma coefficients and cutoff distance for all of the mie/cut pair styles can be mixed. If not explicitly defined, both the repulsive and attractive gamma exponents for different atoms will be calculated following the same mixing rule defined for distances. The default mix value is *geometric*. See the “pair\_modify” command for details.

This pair style supports the *pair\_modify* shift option for the energy of the pair interaction.

This pair style supports the *pair\_modify* tail option for adding a long-range tail correction to the energy and pressure of the pair interaction.

This pair style writes its information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

This pair style supports the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command, meaning the pairwise forces can be partitioned by distance at different levels of the rRESPA hierarchy. See the *run\_style* command for details.

---

## 19.178.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.178.6 Related commands

*pair\_coeff*

## 19.178.7 Default

none

---

**(Mie)** G. Mie, Ann Phys, 316, 657 (1903).

**(Avendano)** C. Avendano, T. Lafitte, A. Galindo, C. S. Adjiman, G. Jackson, E. Muller, J Phys Chem B, 115, 11154 (2011).

## 19.179 pair\_style mliap command

### 19.179.1 Syntax

```
pair_style mliap ... keyword values ...
```

- two keyword/value pairs must be appended
- keyword = *model* or *descriptor*

```
model values = style filename
  style = linear or quadratic or nn or mliappy
  filename = name of file containing model definitions
descriptor values = style filename
  style = sna or so3
  filename = name of file containing descriptor definitions
```

## 19.179.2 Examples

```
pair_style mliap model linear InP.mliap.model descriptor sna InP.mliap.descriptor
pair_style mliap model quadratic W.mliap.model descriptor sna W.mliap.descriptor
pair_style mliap model nn Si.nn.mliap.model descriptor so3 Si.nn.mliap.descriptor
pair_coeff * * In P
```

## 19.179.3 Description

Pair style *mliap* provides a general interface to families of machine-learning interatomic potentials. It allows separate definitions of the interatomic potential functional form (*model*) and the geometric quantities that characterize the atomic positions (*descriptor*). By defining *model* and *descriptor* separately, it is possible to use many different models with a given descriptor, or many different descriptors with a given model. The pair style currently supports only *sna* and *so3* descriptor styles, but it is straightforward to add new descriptor styles.

The SNAP descriptor style *sna* is the same as that used by *pair\_style snap*, including the linear, quadratic, and chem variants. The available models are *linear*, *quadratic*, *nn*, and *mliappy*. The *mliappy* style can be used to couple python models, e.g. PyTorch neural network energy models, and requires building LAMMPS with the PYTHON package (see below). In order to train a model, it is useful to know the gradient or derivative of energy, force, and stress w.r.t. model parameters. This information can be accessed using the related *compute mliap* command.

The descriptor style *so3* is a descriptor that is derived from the the smooth SO(3) power spectrum with the explicit inclusion of a radial basis (*Bartok*) and (*Zagaceta*). The available models are *linear* and *nn*.

The *pair\_style mliap* command must be followed by two keywords *model* and *descriptor* in either order. A single *pair\_coeff* command is also required. The first 2 arguments must be \* \* so as to span all LAMMPS atom types. This is followed by a list of N arguments that specify the mapping of MLIAP element names to LAMMPS atom types, where N is the number of LAMMPS atom types.

The *model* keyword is followed by the model style. This is followed by a single argument specifying the model filename containing the parameters for a set of elements. The model filename usually ends in the *.mliap.model* extension. It may contain parameters for many elements. The only requirement is that it contain at least those element names appearing in the *pair\_coeff* command.

The top of the model file can contain any number of blank and comment lines (start with #), but follows a strict format after that. The first non-blank non-comment line must contain two integers:

- nelems = Number of elements
- nparams = Number of parameters

When the *model* keyword is *linear* or *quadratic*, this is followed by one block for each of the *nelem* elements. Each block consists of *nparams* parameters, one per line. Note that this format is similar, but not identical to that used for the *pair\_style snap* coefficient file. Specifically, the line containing the element weight and radius is omitted, since these are handled by the *descriptor*.

When the *model* keyword is *nn* (neural networks), the model file can contain blank and comment lines (start with #) anywhere. The second non-blank non-comment line must contain the string NET, followed by two integers:

- `ndescriptors` = Number of descriptors
- `nlayers` = Number of layers (including the hidden layers and the output layer)

and followed by a sequence of a string and an integer for each layer:

- Activation function (linear, sigmoid, tanh or relu)
- `nnodes` = Number of nodes

This is followed by one block for each of the *nelem* elements. Each block consists of *scale0* minimum value, *scale1* (maximum - minimum) value, in order to normalize the descriptors, followed by *nparams* parameters, including *bias* and *weights* of the model, starting with the first node of the first layer and so on, with a maximum of 30 values per line.

The detail of *nn* module implementation can be found at ([Yanxon](#)).

---

### Notes on `mliappy` models

When the *model* keyword is *mliappy*, the filename should end in `.pt`, `.pth` for pytorch models, or be a pickle file. To load a model from memory (i.e. an existing python object), specify the filename as “LATER”, and then call `lammmps.mliap.load_model(model)` from python before using the pair style. When using lammmps via the library mode, you will need to call `lammmps.mliappy.activate_mliappy(lmp)` on the active lammmps object before the pair style is defined. This call locates and loads the *mliap*-specific python module that is built into lammmps.

---

The *descriptor* keyword is followed by a descriptor style, and additional arguments. Currently two descriptor styles are available: *sna* and *so3*.

- *sna* indicates the bispectrum component descriptors used by the Spectral Neighbor Analysis Potential (SNAP) potentials of *pair\_style snap*. A single additional argument specifies the descriptor filename containing the parameters and setting used by the SNAP descriptor. The descriptor filename usually ends in the *.mliap.descriptor* extension.
- *so3* indicated the power spectrum component descriptors. A single additional argument specifies the descriptor filename containing the parameters and setting.

The SNAP descriptor file closely follows the format of the *pair\_style snap* parameter file. The file can contain blank and comment lines (start with #) anywhere. Each non-blank non-comment line must contain one keyword/value pair. The required keywords are *rcutfac* and *twojmax*. There are many optional keywords that are described on the *pair\_style snap* doc page. In addition, the SNAP descriptor file must contain the *nelems*, *elems*, *radelems*, and *welems* keywords. The *nelems* keyword specifies the number of elements provided in the other three keywords. The *elems* keyword is followed by a list of *nelems* element names that must include the element names appearing in the *pair\_coeff* command, but can contain other names too. Similarly, the *radelems* and *welems* keywords are followed by lists of *nelems* numbers giving the element radius and element weight of each element. Obviously, the order in which the elements are listed must be consistent for all three keywords.

The SO3 descriptor file is similar to the SNAP descriptor except that it contains a few more arguments (e.g., *nmax* and *alpha*). The preparation of SO3 descriptor and model files can be done with the `Pyxtal_FF` package.

See the *pair\_coeff* page for alternate ways to specify the path for these *model* and *descriptor* files.

---

**Note:** To significantly reduce SO3 descriptor/force calculation time, some properties are pre-computed and reused during the calculation. These can consume a significant amount of RAM for simulations of larger systems since their size depends on the total number of neighbors per MPI process.

---

### 19.179.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, mixing is performed by LAMMPS with user-specifiable parameters as described above. You never need to specify a `pair_coeff` command with I != J arguments for this style.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.179.5 Restrictions

This pair style is part of the ML-IAP package. It is only enabled if LAMMPS was built with that package. In addition, building LAMMPS with the ML-IAP package requires building LAMMPS with the ML-SNAP package. The *mliappy* model requires building LAMMPS with the PYTHON package. See the *Build package* page for more info.

### 19.179.6 Related commands

*pair\_style snap*, *compute mliap*

### 19.179.7 Default

none

---

(**Bartok2013**) Bartok, Kondor, Csanyi, Phys Rev B, 87, 184115 (2013).

(**Zagaceta2020**) Zagaceta, Yanxon, Zhu, J Appl Phys, 128, 045113 (2020).

(**Yanxon2020**) Yanxon, Zagaceta, Tang, Matteson, Zhu, Mach. Learn.: Sci. Technol. 2, 027001 (2020).

## 19.180 pair\_style momb command

### 19.180.1 Syntax

```
pair_style momb cutoff s6 d
```

- cutoff = global cutoff (distance units)
- s6 = global scaling factor of the exchange-correlation functional used (unitless)
- d = damping scaling factor of Grimme's method (unitless)

## 19.180.2 Examples

```
pair_style momb 12.0 0.75 20.0
pair_style hybrid/overlay eam/fs lj/charmm/coul/long 10.0 12.0 momb 12.0 0.75 20.0 morse_
→5.5

pair_coeff 1 2 momb 0.0 1.0 1.0 10.2847 2.361
```

## 19.180.3 Description

Style *momb* computes pairwise van der Waals (vdW) and short-range interactions using the Morse potential and (*Grimme*) method implemented in the Many-Body Metal-Organic (MOMB) force field described comprehensively in (*Fichthorn*) and (*Zhou*). Grimme's method is widely used to correct for dispersion in density functional theory calculations.

$$E = D_0[\exp^{-2\alpha(r-r_0)} - 2\exp^{-\alpha(r-r_0)}] - s_6 \frac{C_6}{r^6} f_{damp}(r, R_r)$$

$$f_{damp}(r, R_r) = \frac{1}{1 + \exp^{-d(r/R_r - 1)}}$$

For the *momb* pair style, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* as described below:

- $D_0$  (energy units)
- $\alpha$  (1/distance units)
- $r_0$  (distance units)
- $C_6$  (energy\*distance<sup>6</sup> units)
- $R_r$  (distance units, typically sum of atomic vdW radii)

## 19.180.4 Restrictions

This style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS is built with that package. See the *Build package* page on for more info.

## 19.180.5 Related commands

*pair\_coeff*, *pair\_style morse*

## 19.180.6 Default

none

**(Grimme)** Grimme, J Comput Chem, 27(15), 1787-1799 (2006).

**(Fichthorn)** Fichthorn, Balankura, Qi, CrystEngComm, 18(29), 5410-5417 (2016).

**(Zhou)** Zhou, Saidi, Fichthorn, J Phys Chem C, 118(6), 3366-3374 (2014).

## 19.181 pair\_style morse command

Accelerator Variants: *morse/gpu*, *morse/omp*, *morse/opt*, *morse/kk*

## 19.182 pair\_style morse/smooth/linear command

Accelerator Variants: *morse/smooth/linear/omp*

### 19.182.1 Syntax

```
pair_style style args
```

- style = *morse* or *morse/smooth/linear* or *morse/soft*
- args = list of arguments for a particular style

*morse* args = cutoff

cutoff = global cutoff for Morse interactions (distance units)

*morse/smooth/linear* args = cutoff

cutoff = global cutoff for Morse interactions (distance units)

### 19.182.2 Examples

```
pair_style morse 2.5
pair_style morse/smooth/linear 2.5
pair_coeff * * 100.0 2.0 1.5
pair_coeff 1 1 100.0 2.0 1.5 3.0
```

### 19.182.3 Description

Style *morse* computes pairwise interactions with the formula

$$E = D_0 \left[ e^{-2\alpha(r-r_0)} - 2e^{-\alpha(r-r_0)} \right] \quad r < r_c$$

$r_c$  is the cutoff.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $D_0$  (energy units)
- $\alpha$  (1/distance units)
- $r_0$  (distance units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global morse cutoff is used.

---



The *morse/smooth/linear* variant is similar to the *lj/smooth/linear* variant in that it adds to the potential a shift and a linear term so that both, potential energy and force, go to zero at the cut-off:

$$\begin{aligned}\phi(r) &= D_0 \left[ e^{-2\alpha(r-r_0)} - 2e^{-\alpha(r-r_0)} \right] & r < r_c \\ E(r) &= \phi(r) - \phi(R_c) - (r - R_c) \left. \frac{d\phi}{dr} \right|_{r=R_c} & r < R_c\end{aligned}$$

The syntax of the `pair_style` and `pair_coeff` commands are the same for the *morse* and *morse/smooth/linear* styles.

A version of the *morse* style with a soft core, *morse/soft*, suitable for use in free energy calculations, is part of the FEP package and is documented with the *pair\_style \*/soft* styles. The version with soft core is only available if LAMMPS was built with that package. See the *Build package* page for more info.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 19.182.4 Mixing, shift, table, tail correction, restart, rRESPA info

None of these pair styles support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

All of these pair styles support the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table options is not relevant for the Morse pair styles.

None of these pair styles support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

All of these pair styles write their information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

## 19.182.5 Restrictions

The *morse/smooth/linear* pair style is only enabled if LAMMPS was built with the EXTRA-PAIR package. See the *Build package* page for more info.

## 19.182.6 Related commands

*pair\_coeff*, *pair\_style \*/soft*

## 19.182.7 Default

none

## 19.183 pair\_style multi/lucy command

### 19.183.1 Syntax

```
pair_style multi/lucy style N keyword ...
```

- style = *lookup* or *linear* = method of interpolation
- N = use N values in *lookup*, *linear* tables

### 19.183.2 Examples

```
pair_style multi/lucy linear 1000  
pair_coeff * * multibody.table ENTRY1 7.0
```

### 19.183.3 Description

Style *multi/lucy* computes a density-dependent force following from the many-body form described in (*Moore*) and (*Warren*) as

$$F_i^{DD}(\rho_i, \rho_j, r_{ij}) = \frac{1}{2} \omega_{DD}(r_{ij}) [A(\rho_i) + A(\rho_j)] e_{ij}$$

which consists of a density-dependent function,  $A(\rho)$ , and a radial-dependent weight function,  $\omega_{DD}(r_{ij})$ . The radial-dependent weight function,  $\omega_{DD}(r_{ij})$ , is taken as the Lucy function:

$$\omega_{DD}(r_{ij}) = \left(1 + \frac{3r_{ij}}{r_{cut}}\right) \left(1 + \frac{r_{ij}}{r_{cut}}\right)^3$$

The density-dependent energy for a given particle is given by:

$$u_i^{DD}(\rho_i) = \frac{\pi r_{cut}^4}{84} \int_{\rho_0}^{\rho_i} A(\rho') d\rho'$$

See the supporting information of (*Brennan*) or the publication by (*Moore*) for more details on the functional form.

An interpolation table is used to evaluate the density-dependent energy ( $\int A(\rho')d\rho'$ ) and force ( $A(\rho')$ ). Note that the prefactor to the energy is computed after the interpolation, thus the  $\int A(\rho')d\rho'$  will have units of energy / length<sup>4</sup>.

The interpolation table is created as a pre-computation by fitting cubic splines to the file values and interpolating the density-dependent energy and force at each of  $N$  densities. During a simulation, the tables are used to interpolate the density-dependent energy and force as needed for each pair of particles separated by a distance  $R$ . The interpolation is done in one of 2 styles: *lookup* and *linear*.

For the *lookup* style, the density is used to find the nearest table entry, which is the density-dependent energy and force.

For the *linear* style, the density is used to find the 2 surrounding table values from which the density-dependent energy and force are computed by linear interpolation.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above.

- filename
- keyword
- cutoff (distance units)

The filename specifies a file containing the tabulated density-dependent energy and force. The keyword specifies a section of the file. The cutoff is an optional coefficient. If not specified, the outer cutoff in the table itself (see below) will be used to build an interpolation table that extend to the largest tabulated distance. If specified, only file values up to the cutoff are used to create the interpolation table. The format of this file is described below.

The format of a tabulated file is a series of one or more sections, defined as follows (without the parenthesized comments):

# Density-dependent function (one or more comment or blank lines)

```
DD-FUNCTION                (keyword is first text on line)
N 500 R 1.0 10.0          (N, R, RSQ parameters)
                          (blank)
1 1.0 25.5 102.34        (index, density, energy/r^4, force)
2 1.02 23.4 98.5
...
500 10.0 0.001 0.003
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The line can contain additional text, but the initial text must match the argument specified in the *pair\_coeff* command. The next line lists (in any order) one or more parameters for the table. Each parameter is a keyword followed by one or more numeric values.

The parameter “N” is required and its value is the number of table entries that follow. Note that this may be different than the  $N$  specified in the *pair\_style multi/lucy* command. Let  $N_{\text{table}} = N$  in the *pair\_style* command, and  $N_{\text{file}} = “N”$  in the tabulated file. What LAMMPS does is a preliminary interpolation by creating splines using the  $N_{\text{file}}$  tabulated values as nodal points. It uses these to interpolate the density-dependent energy and force at  $N_{\text{table}}$  different points. The resulting tables of length  $N_{\text{table}}$  are then used as described above, when computing the density-dependent energy and force. This means that if you want the interpolation tables of length  $N_{\text{table}}$  to match exactly what is in the tabulated file (with effectively no preliminary interpolation), you should set  $N_{\text{table}} = N_{\text{file}}$ , and use the “RSQ” parameter. This is because the internal table abscissa is always RSQ (separation distance squared), for efficient lookup.

All other parameters are optional. If “R” or “RSQ” does not appear, then the distances in each line of the table are used as-is to perform spline interpolation. In this case, the table values can be spaced in *density* uniformly or however you wish to position table values in regions of large gradients.

If used, the parameters “R” or “RSQ” are followed by 2 values *rlo* and *rhi*. If specified, the density associated with each density-dependent energy and force value is computed from these 2 values (at high accuracy), rather than using the (low-accuracy) value listed in each line of the table. The density values in the table file are ignored in this case. For “R”, distances uniformly spaced between *rlo* and *rhi* are computed; for “RSQ”, squared distances uniformly spaced between *rlo\*rlo* and *rhi\*rhi* are computed.

---

**Note:** If you use “R” or “RSQ”, the tabulated distance values in the file are effectively ignored, and replaced by new values as described in the previous paragraph. If the density value in the table is not very close to the new value (i.e. round-off difference), then you will be assigning density-dependent energy and force values to a different density, which is probably not what you want. LAMMPS will warn if this is occurring.

---

Following a blank line, the next N lines list the tabulated values. On each line, the first value is the index from 1 to N, the second value is *r* (in density units), the third value is the density-dependent function value (in energy units / length<sup>4</sup>), and the fourth is the force (in force units). The density values must increase from one line to the next.

Note that one file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified keyword.

---

#### 19.183.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

The *pair\_modify* shift, table, and tail options are not relevant for this pair style.

This pair style writes the settings for the “pair\_style multi/lucy” command to *binary restart files*, so a pair\_style command does not need to be specified in an input script that reads a restart file. However, the coefficient information is not stored in the restart file, since it is tabulated in the potential files. Thus, pair\_coeff commands do need to be specified in the restart input script.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.183.5 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 19.183.6 Related commands

*pair\_coeff*

## 19.183.7 Default

none

(Warren) Warren, Phys Rev E, 68, 066702 (2003).

(Brennan) Brennan, J Chem Phys Lett, 5, 2144-2149 (2014).

(Moore) Moore, J Chem Phys, 144, 104501 (2016).

## 19.184 pair\_style multi/lucy/rx command

Accelerator Variants: *multi/lucy/rx/kk*

### 19.184.1 Syntax

```
pair_style multi/lucy/rx style N keyword ...
```

- style = *lookup* or *linear* = method of interpolation
- N = use N values in *lookup*, *linear* tables
- weighting = fractional or molecular (optional)

### 19.184.2 Examples

```
pair_style multi/lucy/rx linear 1000
pair_style multi/lucy/rx linear 1000 fractional
pair_style multi/lucy/rx linear 1000 molecular
pair_coeff * * multibody.table ENTRY1 h2o h2o 7.0
pair_coeff * * multibody.table ENTRY1 h2o 1fluid 7.0
```

### 19.184.3 Description

Style *multi/lucy/rx* is used in reaction DPD simulations, where the coarse-grained (CG) particles are composed of  $m$  species whose reaction rate kinetics are determined from a set of  $n$  reaction rate equations through the *fix rx* command. The species of one CG particle can interact with a species in a neighboring CG particle through a site-site interaction potential model. Style *multi/lucy/rx* computes the site-site density-dependent force following from the many-body form described in (Moore) and (Warren) as

$$F_i^{DD}(\rho_i, \rho_j, r_{ij}) = \frac{1}{2} \omega_{DD}(r_{ij}) [A(\rho_i) + A(\rho_j)] e_{ij}$$

which consists of a density-dependent function,  $A(\rho)$ , and a radial-dependent weight function,  $\omega_{DD}(r_{ij})$ . The radial-dependent weight function,  $\omega_{DD}(r_{ij})$ , is taken as the Lucy function:

$$\omega_{DD}(r_{ij}) = \left(1 + \frac{3r_{ij}}{r_{cut}}\right) \left(1 + \frac{r_{ij}}{r_{cut}}\right)^3$$

The density-dependent energy for a given particle is given by:

$$u_i^{DD}(\rho_i) = \frac{\pi r_{cut}^4}{84} \int_{\rho_0}^{\rho_i} A(\rho') d\rho'$$

See the supporting information of (Brennan) or the publication by (Moore) for more details on the functional form.

An interpolation table is used to evaluate the density-dependent energy ( $\int A(\rho') d\rho'$ ) and force ( $A(\rho')$ ). Note that the prefactor to the energy is computed after the interpolation, thus the  $\int A(\rho') d\rho'$  will have units of energy / length<sup>4</sup>.

The interpolation table is created as a pre-computation by fitting cubic splines to the file values and interpolating the density-dependent energy and force at each of  $N$  densities. During a simulation, the tables are used to interpolate the density-dependent energy and force as needed for each pair of particles separated by a distance  $R$ . The interpolation is done in one of 2 styles: *lookup* and *linear*.

For the *lookup* style, the density is used to find the nearest table entry, which is the density-dependent energy and force.

For the *linear* style, the density is used to find the 2 surrounding table values from which the density-dependent energy and force are computed by linear interpolation.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above.

- filename
- keyword
- species1
- species2
- cutoff (distance units)

The filename specifies a file containing the tabulated density-dependent energy and force. The keyword specifies a section of the file. The cutoff is an optional coefficient. If not specified, the outer cutoff in the table itself (see below) will be used to build an interpolation table that extend to the largest tabulated distance. If specified, only file values up to the cutoff are used to create the interpolation table. The format of this file is described below.

The species tags define the site-site interaction potential between two species contained within two different particles. The species tags must either correspond to the species defined in the reaction kinetics files specified with the *fix rx* command or they must correspond to the tag “1fluid”, signifying interaction with a product species mixture determined through a one-fluid approximation. The interaction potential is weighted by the geometric average of either the mole fraction concentrations or the number of molecules associated with the interacting coarse-grained particles (see the *fractional* or *molecular* weighting pair style options). The coarse-grained potential is stored before and after the reaction kinetics solver is applied, where the difference is defined to be the internal chemical energy (uChem).

---

The format of a tabulated file is a series of one or more sections, defined as follows (without the parenthesized comments):

```
# Density-dependent function (one or more comment or blank lines)
```

```
DD-FUNCTION                (keyword is first text on line)
N 500 R 1.0 10.0           (N, R, RSQ parameters)
                            (blank)
1 1.0 25.5 102.34          (index, density, energy/r^4, force)
2 1.02 23.4 98.5
...
500 10.0 0.001 0.003
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The line can contain additional text, but the initial text must match the argument specified in the `pair_coeff` command. The next line lists (in any order) one or more parameters for the table. Each parameter is a keyword followed by one or more numeric values.

The parameter “N” is required and its value is the number of table entries that follow. Note that this may be different than the  $N$  specified in the `pair_style multi/lucy/rx` command. Let  $N_{\text{table}} = N$  in the `pair_style` command, and  $N_{\text{file}} = “N”$  in the tabulated file. What LAMMPS does is a preliminary interpolation by creating splines using the  $N_{\text{file}}$  tabulated values as nodal points. It uses these to interpolate the density-dependent energy and force at  $N_{\text{table}}$  different points. The resulting tables of length  $N_{\text{table}}$  are then used as described above, when computing the density-dependent energy and force. This means that if you want the interpolation tables of length  $N_{\text{table}}$  to match exactly what is in the tabulated file (with effectively no preliminary interpolation), you should set  $N_{\text{table}} = N_{\text{file}}$ , and use the “RSQ” parameter. This is because the internal table abscissa is always RSQ (separation distance squared), for efficient lookup.

All other parameters are optional. If “R” or “RSQ” does not appear, then the distances in each line of the table are used as-is to perform spline interpolation. In this case, the table values can be spaced in *density* uniformly or however you wish to position table values in regions of large gradients.

If used, the parameters “R” or “RSQ” are followed by 2 values *rlo* and *rhi*. If specified, the density associated with each density-dependent energy and force value is computed from these 2 values (at high accuracy), rather than using the (low-accuracy) value listed in each line of the table. The density values in the table file are ignored in this case. For “R”, distances uniformly spaced between *rlo* and *rhi* are computed; for “RSQ”, squared distances uniformly spaced between  $rlo*rlo$  and  $rhi*rhi$  are computed.

---

**Note:** If you use “R” or “RSQ”, the tabulated distance values in the file are effectively ignored, and replaced by new values as described in the previous paragraph. If the density value in the table is not very close to the new value (i.e. round-off difference), then you will be assigning density-dependent energy and force values to a different density, which is probably not what you want. LAMMPS will warn if this is occurring.

---

Following a blank line, the next  $N$  lines list the tabulated values. On each line, the first value is the index from 1 to  $N$ , the second value is  $r$  (in density units), the third value is the density-dependent function value (in energy units /  $\text{length}^4$ ), and the fourth is the force (in force units). The density values must increase from one line to the next.

Note that one file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified keyword.

---

#### 19.184.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

The `pair_modify` shift, table, and tail options are not relevant for this pair style.

This pair style writes the settings for the “`pair_style multi/lucy/rx`” command to *binary restart files*, so a `pair_style` command does not need to be specified in an input script that reads a restart file. However, the coefficient information is not stored in the restart file, since it is tabulated in the potential files. Thus, `pair_coeff` commands do need to be specified in the restart input script.

This pair style can only be used via the `pair` keyword of the `run_style respa` command. It does not support the *inner*, *middle*, *outer* keywords.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages*

page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 19.184.5 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.184.6 Related commands

*pair\_coeff*

## 19.184.7 Default

fractional weighting

---

(Warren) Warren, Phys Rev E, 68, 066702 (2003).

(Brennan) Brennan, J Chem Phys Lett, 5, 2144-2149 (2014).

(Moore) Moore, J Chem Phys, 144, 104501 (2016).

## 19.185 pair\_style nb3b/harmonic command

### 19.185.1 Syntax

```
pair_style nb3b/harmonic
```

### 19.185.2 Examples

```
pair_style nb3b/harmonic
pair_coeff * * MgOH.nb3bharmonic Mg O H
```



### 19.185.3 Description

This pair style computes a non-bonded 3-body harmonic potential for the energy  $E$  of a system of atoms as

$$E = K(\theta - \theta_0)^2$$

where  $\theta_0$  is the equilibrium value of the angle and  $K$  is a prefactor. Note that the usual 1/2 factor is included in  $K$ . The form of the potential is identical to that used in angle\_style *harmonic*, but in this case, the atoms do not need to be explicitly bonded.

Only a single pair\_coeff command is used with this style which specifies a potential file with parameters for specified elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the pair\_coeff command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the potential file.

As an example, imagine a file SiC.nb3b.harmonic has potential values for Si and C. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Si, and the fourth to be C, you would use the following pair\_coeff command:

```
pair_coeff * * SiC.nb3b.harmonic Si Si Si C
```

The first 2 arguments must be \* \* so as to span all LAMMPS atom types. The first three Si arguments map LAMMPS atom types 1,2,3 to the Si element in the potential file. The final C argument maps LAMMPS atom type 4 to the C element in the potential file. If a mapping value is specified as NULL, the mapping is not performed. This can be used when the potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials. An example of a pair\_coeff command for use with the *hybrid* pair style is:

```
pair_coeff * * nb3b/harmonic MgOH.nb3b.harmonic Mg O H
```

Three-body non-bonded harmonic files in the *potentials* directory of the LAMMPS distribution have a “.nb3b.harmonic” suffix. Lines that are not blank or comments (starting with #) define parameters for a triplet of elements.

Each entry has six arguments. The first three are atom types as referenced in the LAMMPS input file. The first argument specifies the central atom. The fourth argument indicates the  $K$  parameter. The fifth argument indicates  $\theta_0$ . The sixth argument indicates a separation cutoff in Angstroms.

For a given entry, if the second and third arguments are identical, then the entry is for a cutoff for the distance between types 1 and 2 (values for  $K$  and  $\theta_0$  are irrelevant in this case).

For a given entry, if the first three arguments are all different, then the entry is for the  $K$  and  $\theta_0$  parameters (the cutoff in this case is irrelevant).

It is required that the potential file contains entries for *all* permutations of the elements listed in the pair\_coeff command. If certain combinations are not parameterized the corresponding parameters should be set to zero. The potential file can also contain entries for additional elements which are not used in a particular simulation; LAMMPS ignores those entries.

### 19.185.4 Restrictions

This pair style can only be used if LAMMPS was built with the MANYBODY package. See the *Build package* page for more info.

### 19.185.5 Related commands

*pair\_coeff*

### 19.185.6 Default

none

## 19.186 pair\_style nm/cut command

Accelerator Variants: *nm/cut/omp*

## 19.187 pair\_style nm/cut/split command

## 19.188 pair\_style nm/cut/coul/cut command

Accelerator Variants: *nm/cut/coul/cut/omp*

## 19.189 pair\_style nm/cut/coul/long command

Accelerator Variants: *nm/cut/coul/long/omp*

### 19.189.1 Syntax

**pair\_style** style args

- style = *nm/cut* or *nm/cut/split* or *nm/cut/coul/cut* or *nm/cut/coul/long*
- args = list of arguments for a particular style

*nm/cut* args = cutoff

cutoff = global cutoff for Pair interactions (distance units)

*nm/cut/split* args = cutoff

cutoff = global cutoff for Pair interactions (distance units)

*nm/cut/coul/cut* args = cutoff (cutoff2)

cutoff = global cutoff for Pair (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

*nm/cut/coul/long* args = cutoff (cutoff2)

cutoff = global cutoff for Pair (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

## 19.189.2 Examples

```

pair_style nm/cut 12.0
pair_coeff * * 0.01 5.4 8.0 7.0
pair_coeff 1 1 0.01 4.4 7.0 6.0

pair_style nm/cut/split 1.12246
pair_coeff 1 1 1.0 1.1246 12 6
pair_coeff * * 1.0 1.1246 11 6

pair_style nm/cut/coul/cut 12.0 15.0
pair_coeff * * 0.01 5.4 8.0 7.0
pair_coeff 1 1 0.01 4.4 7.0 6.0

pair_style nm/cut/coul/long 12.0 15.0
pair_coeff * * 0.01 5.4 8.0 7.0
pair_coeff 1 1 0.01 4.4 7.0 6.0

```

## 19.189.3 Description

Style *nm* computes site-site interactions based on the N-M potential by *Clarke*, mainly used for ionic liquids. A site can represent a single atom or a united-atom site. The energy of an interaction has the following form:

$$E = \frac{E_0}{(n-m)} \left[ m \left( \frac{r_0}{r} \right)^n - n \left( \frac{r_0}{r} \right)^m \right] \quad r < r_c$$

where  $r_c$  is the cutoff and  $r_0$  is the minimum of the potential. Please note that this differs from the convention used for other Lennard-Jones potentials in LAMMPS where  $\sigma$  represents the location where the energy is zero.

Style *nm/cut/split* applies the standard LJ (12-6) potential above  $r_0 = 2^{\frac{1}{6}}\sigma$ . Style *nm/cut/split* is employed in polymer equilibration protocols that combine core-softening approaches with topology-changing moves *Dietz*.

Style *nm/cut/coul/cut* adds a Coulombic pairwise interaction given by

$$E = \frac{Cq_iq_j}{\epsilon r} \quad r < r_c$$

where  $C$  is an energy-conversion constant,  $q_i$  and  $q_j$  are the charges on the 2 atoms, and epsilon is the dielectric constant which can be set by the *dielectric* command. If one cutoff is specified in the *pair\_style* command, it is used for both the N-M and Coulombic terms. If two cutoffs are specified, they are used as cutoffs for the N-M and Coulombic terms respectively.

Styles *nm/cut/coul/long* compute the same Coulombic interactions as style *nm/cut/coul/cut* except that an additional damping factor is applied to the Coulombic term so it can be used in conjunction with the *kspace\_style* command and its *ewald* or *pppm* option. The Coulombic cutoff specified for this style means that pairwise interactions within this distance are computed directly; interactions outside that distance are computed in reciprocal space.

For all of the *nm* pair styles, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands.

- $E_0$  (energy units)
- $r_0$  (distance units)
- $n$  (unitless)
- $m$  (unitless)

- cutoff1 (distance units)
- cutoff2 (distance units)

The latter 2 coefficients are optional. If not specified, the global N-M and Coulombic cutoffs specified in the `pair_style` command are used. If only one cutoff is specified, it is used as the cutoff for both N-M and Coulombic interactions for this type pair. If both coefficients are specified, they are used as the N-M and Coulombic cutoffs for this type pair. You cannot specify 2 cutoffs for style `nm`, since it has no Coulombic terms.

For `nm/cut/coul/long` only the N-M cutoff can be specified since a Coulombic cutoff cannot be specified for an individual I,J type pair. All type pairs use the same global Coulombic cutoff specified in the `pair_style` command.

---

#### 19.189.4 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles do not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

All of the `nm` pair styles supports the `pair_modify` shift option for the energy of the pair interaction.

The `nm/cut/coul/long` pair styles support the `pair_modify` table option since they can tabulate the short-range portion of the long-range Coulombic interaction.

All of the `nm` pair styles support the `pair_modify` tail option for adding a long-range tail correction to the energy and pressure for the N-M portion of the pair interaction.

All of the `nm` pair styles write their information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

All of the `nm` pair styles can only be used via the `pair` keyword of the `run_style respa` command. They do not support the `inner`, `middle`, `outer` keywords.

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 19.189.5 Restrictions

These pair styles are part of the EXTRA-PAIR package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.189.6 Related commands

*pair\_coeff*, *pair style lj/cut*, *bond style fene/nm*

## 19.189.7 Default

none

---

(**Clarke**) Clarke and Smith, J Chem Phys, 84, 2290 (1986).

(**Dietz**) Dietz and Hoy, J. Chem Phys, 156, 014103 (2022).

## 19.190 pair\_style none command

### 19.190.1 Syntax

```
pair_style none
```

### 19.190.2 Examples

```
pair_style none
```

### 19.190.3 Description

Using a pair style of *none* means that any previous pair style setting will be deleted and pairwise forces and energies are not computed.

As a consequence there will be a pairwise force cutoff of 0.0, which has implications for the default setting of the neighbor list and the communication cutoff. Those are the sum of the largest pairwise cutoff and the neighbor skin distance (see the documentation of the *neighbor* command and the *comm\_modify* command). When you have bonds, angles, dihedrals, or impropers defined at the same time, you must set the communication cutoff so that communication cutoff distance is large enough to acquire and communicate sufficient ghost atoms from neighboring sub-domains as needed for computing bonds, angles, etc.

A pair style of *none* will also not request a pairwise neighbor list. However if the *neighbor* style is *bin*, data structures for binning are still allocated. If the neighbor list cutoff is small, then these data structures can consume a large amount of memory. So you should either set the neighbor style to *nsq* or set the skin distance to a larger value.

See the *pair\_style zero* for a way to set a pairwise cutoff and thus trigger the building of a neighbor lists and setting a corresponding communication cutoff, but compute no pairwise interactions.

## 19.190.4 Restrictions

You must not use a *pair\_coeff* command with this pair style. Since there is no interaction computed, you cannot set any coefficients for it.

## 19.190.5 Related commands

*pair\_style zero*

## 19.190.6 Default

none

## 19.191 pair\_style oxdna/excv command

## 19.192 pair\_style oxdna/stk command

## 19.193 pair\_style oxdna/hbond command

## 19.194 pair\_style oxdna/xstk command

## 19.195 pair\_style oxdna/coaxstk command

### 19.195.1 Syntax

```
pair_style style1
pair_coeff * * style2 args
```

- style1 = *hybrid/overlay oxdna/excv oxdna/stk oxdna/hbond oxdna/xstk oxdna/coaxstk*
- style2 = *oxdna/excv* or *oxdna/stk* or *oxdna/hbond* or *oxdna/xstk* or *oxdna/coaxstk*
- args = list of arguments for these particular styles

```
oxdna/stk args = seq T xi kappa 6.0 0.4 0.9 0.32 0.75 1.3 0 0.8 0.9 0 0.95 0.9 0 0.95 2.
↪ 0 0.65 2.0 0.65
```

seq = seqav (for average sequence stacking strength) or seqdep (for sequence-dependent stacking strength)

T = temperature (oxDNA units, 0.1 = 300 K)

xi = 1.3448 (temperature-independent coefficient in stacking strength)

kappa = 2.6568 (coefficient of linear temperature dependence in stacking strength)

```
oxdna/hbond args = seq eps 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.5 0 0.7 0.46 3.
↪ 141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966 0.45
```

seq = seqav (for average sequence base-pairing strength) or seqdep (for sequence-dependent base-pairing strength)

eps = 1.077 (between base pairs A-T and C-G) or 0 (all other pairs)

## 19.195.2 Examples

```

pair_style hybrid/overlay oxdna/excv oxdna/stk oxdna/hbond oxdna/xstk oxdna/coaxstk
pair_coeff * * oxdna/excv      2.0 0.7 0.675 2.0 0.515 0.5 2.0 0.33 0.32
pair_coeff * * oxdna/stk      seqdep 0.1 1.3448 2.6568 6.0 0.4 0.9 0.32 0.75 1.3 0 0.8 0.
↪9 0 0.95 0.9 0 0.95 2.0 0.65 2.0 0.65
pair_coeff * * oxdna/hbond    seqdep 0.0 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.5 0.
↪0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966 0.45
pair_coeff 1 4 oxdna/hbond    seqdep 1.077 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.5.
↪0 0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966 0.
↪45
pair_coeff 2 3 oxdna/hbond    seqdep 1.077 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.5.
↪0 0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966 0.
↪45
pair_coeff * * oxdna/xstk     47.5 0.575 0.675 0.495 0.655 2.25 0.791592653589793 0.58 1.
↪7 1.0 0.68 1.7 1.0 0.68 1.5 0 0.65 1.7 0.875 0.68 1.7 0.875 0.68
pair_coeff * * oxdna/coaxstk 46.0 0.4 0.6 0.22 0.58 2.0 2.541592653589793 0.65 1.3 0 0.8.
↪0.9 0 0.95 0.9 0 0.95 2.0 -0.65 2.0 -0.65

```

## 19.195.3 Description

The *oxdna* pair styles compute the pairwise-additive parts of the oxDNA force field for coarse-grained modelling of DNA. The effective interaction between the nucleotides consists of potentials for the excluded volume interaction *oxdna/excv*, the stacking *oxdna/stk*, cross-stacking *oxdna/xstk* and coaxial stacking interaction *oxdna/coaxstk* as well as the hydrogen-bonding interaction *oxdna/hbond* between complementary pairs of nucleotides on opposite strands. Average sequence or sequence-dependent stacking and base-pairing strengths are supported (*Sulc*). Quasi-unique base-pairing between nucleotides can be achieved by using more complementary pairs of atom types like 5-8 and 6-7, 9-12 and 10-11, 13-16 and 14-15, etc. This prevents the hybridization of in principle complementary bases within Ntypes/4 bases up and down along the backbone.

The exact functional form of the pair styles is rather complex. The individual potentials consist of products of modulation factors, which themselves are constructed from a number of more basic potentials (Morse, Lennard-Jones, harmonic angle and distance) as well as quadratic smoothing and modulation terms. We refer to (*Ouldridge-DPhil*) and (*Ouldridge*) for a detailed description of the oxDNA force field.

---

**Note:** These pair styles have to be used together with the related oxDNA bond style *oxdnaffene* for the connectivity of the phosphate backbone (see also documentation of *bond\_style oxdnaffene*). Most of the coefficients in the above example have to be kept fixed and cannot be changed without reparameterizing the entire model. Exceptions are the first four coefficients after *oxdna/stk* (seq=seqdep, T=0.1, xi=1.3448 and kappa=2.6568 in the above example) and the first coefficient after *oxdna/hbond* (seq=seqdep in the above example). When using a Langevin thermostat, e.g. through *fix langevin* or *fix nve/dotc/langevin* the temperature coefficients have to be matched to the one used in the fix.

---



---

**Note:** These pair styles have to be used with the *atom\_style hybrid bond ellipsoid oxdna* (see documentation of *atom\_style*). The *atom\_style oxdna* stores the 3'-to-5' polarity of the nucleotide strand, which is set through the bond topology in the data file. The first (second) atom in a bond definition is understood to point towards the 3'-end (5'-end) of the strand.

---

Example input and data files for DNA duplexes can be found in examples/PACKAGES/cgdna/examples/oxDNA/ and /oxDNA2/. A simple python setup tool which creates single straight or helical DNA strands, DNA duplexes or arrays of DNA duplexes can be found in examples/PACKAGES/cgdna/util/.

Please cite (*Henrich*) in any publication that uses this implementation. An updated documentation that contains general information on the model, its implementation and performance as well as the structure of the data and input file can be found [here](#).

Please cite also the relevant oxDNA publications (*Ouldridge*), (*Ouldridge-DPhil*) and (*Sulc*).

---

## 19.195.4 Restrictions

These pair styles can only be used if LAMMPS was built with the CG-DNA package and the MOLECULE and ASPHERE package. See the *Build package* page for more info.

## 19.195.5 Related commands

*bond\_style oxdna/fene, pair\_coeff, bond\_style oxdna2/fene, pair\_style oxdna2/excv, bond\_style oxrna2/fene, pair\_style oxrna2/excv, atom\_style oxdna, fix nve/dotc/langevin*

## 19.195.6 Default

none

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**(Henrich)** O. Henrich, Y. A. Gutierrez-Fosado, T. Curk, T. E. Ouldridge, Eur. Phys. J. E 41, 57 (2018).

**(Ouldridge-DPhil)** T.E. Ouldridge, Coarse-grained modelling of DNA and DNA self-assembly, DPhil. University of Oxford (2011).

**(Ouldridge)** T.E. Ouldridge, A.A. Louis, J.P.K. Doye, J. Chem. Phys. 134, 085101 (2011).

**(Sulc)** P. Sulc, F. Romano, T.E. Ouldridge, L. Rovigatti, J.P.K. Doye, A.A. Louis, J. Chem. Phys. 137, 135101 (2012).

## 19.196 pair\_style oxdna2/excv command

## 19.197 pair\_style oxdna2/stk command

## 19.198 pair\_style oxdna2/hbond command

## 19.199 pair\_style oxdna2/xstk command

## 19.200 pair\_style oxdna2/coaxstk command

## 19.201 pair\_style oxdna2/dh command

### 19.201.1 Syntax



```
pair_style style1
```

```
pair_coeff * * style2 args
```

- style1 = *hybrid/overlay oxdna2/excv oxdna2/stk oxdna2/hbond oxdna2/xstk oxdna2/coaxstk oxdna2/dh*
- style2 = *oxdna2/excv* or *oxdna2/stk* or *oxdna2/hbond* or *oxdna2/xstk* or *oxdna2/coaxstk* or *oxdna2/dh*
- args = list of arguments for these particular styles

```
oxdna2/stk args = seq T xi kappa 6.0 0.4 0.9 0.32 0.75 1.3 0 0.8 0.9 0 0.95 0.9 0 0.95 2.
↳ 0 0.65 2.0 0.65
```

```
seq = seqav (for average sequence stacking strength) or seqdep (for sequence-dependent
↳ stacking strength)
```

```
T = temperature (oxDNA units, 0.1 = 300 K)
```

```
xi = 1.3523 (temperature-independent coefficient in stacking strength)
```

```
kappa = 2.6717 (coefficient of linear temperature dependence in stacking strength)
```

```
oxdna2/hbond args = seq eps 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.5 0 0.7 0.46 3.
↳ 141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966 0.45
```

```
seq = seqav (for average sequence base-pairing strength) or seqdep (for
↳ sequence-dependent base-pairing strength)
```

```
eps = 1.0678 (between base pairs A-T and C-G) or 0 (all other pairs)
```

```
oxdna2/dh args = T rhos qeff
```

```
T = temperature (oxDNA units, 0.1 = 300 K)
```

```
rhos = salt concentration (mole per litre)
```

```
qeff = 0.815 (effective charge in elementary charges)
```

## 19.201.2 Examples

```
pair_style hybrid/overlay oxdna2/excv oxdna2/stk oxdna2/hbond oxdna2/xstk oxdna2/coaxstk
↳ oxdna2/dh
pair_coeff * * oxdna2/excv 2.0 0.7 0.675 2.0 0.515 0.5 2.0 0.33 0.32
pair_coeff * * oxdna2/stk seqdep 0.1 1.3523 2.6717 6.0 0.4 0.9 0.32 0.75 1.3 0 0.8 0.
↳ 9 0 0.95 0.9 0 0.95 2.0 0.65 2.0 0.65
pair_coeff * * oxdna2/hbond seqdep 0.0 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.5 0
↳ 0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966 0.45
pair_coeff 1 4 oxdna2/hbond seqdep 1.0678 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.
↳ 5 0 0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966
↳ 0.45
pair_coeff 2 3 oxdna2/hbond seqdep 1.0678 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.
↳ 5 0 0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966
↳ 0.45
pair_coeff * * oxdna2/xstk 47.5 0.575 0.675 0.495 0.655 2.25 0.791592653589793 0.58 1.
↳ 7 1.0 0.68 1.7 1.0 0.68 1.5 0 0.65 1.7 0.875 0.68 1.7 0.875 0.68
pair_coeff * * oxdna2/coaxstk 58.5 0.4 0.6 0.22 0.58 2.0 2.891592653589793 0.65 1.3 0 0.
↳ 8 0.9 0 0.95 0.9 0 0.95 40.0 3.116592653589793
pair_coeff * * oxdna2/dh 0.1 0.5 0.815
```

### 19.201.3 Description

The *oxdna2* pair styles compute the pairwise-additive parts of the oxDNA force field for coarse-grained modelling of DNA. The effective interaction between the nucleotides consists of potentials for the excluded volume interaction *oxdna2/excv*, the stacking *oxdna2/stk*, cross-stacking *oxdna2/xstk* and coaxial stacking interaction *oxdna2/coaxstk*, electrostatic Debye-Hueckel interaction *oxdna2/dh* as well as the hydrogen-bonding interaction *oxdna2/hbond* between complementary pairs of nucleotides on opposite strands. Average sequence or sequence-dependent stacking and base-pairing strengths are supported (*Sulc*). Quasi-unique base-pairing between nucleotides can be achieved by using more complementary pairs of atom types like 5-8 and 6-7, 9-12 and 10-11, 13-16 and 14-15, etc. This prevents the hybridization of in principle complementary bases within Ntypes/4 bases up and down along the backbone.

The exact functional form of the pair styles is rather complex. The individual potentials consist of products of modulation factors, which themselves are constructed from a number of more basic potentials (Morse, Lennard-Jones, harmonic angle and distance) as well as quadratic smoothing and modulation terms. We refer to (*Snodin*) and the original oxDNA publications (*Ouldridge-DPhil*) and (*Ouldridge*) for a detailed description of the oxDNA2 force field.

---

**Note:** These pair styles have to be used together with the related oxDNA2 bond style *oxdna2/fene* for the connectivity of the phosphate backbone (see also documentation of *bond\_style oxdna2/fene*). Most of the coefficients in the above example have to be kept fixed and cannot be changed without reparameterizing the entire model. Exceptions are the first four coefficients after *oxdna2/stk* (seq=seqdep, T=0.1, xi=1.3523 and kappa=2.6717 in the above example), the first coefficient after *oxdna2/hbond* (seq=seqdep in the above example) and the three coefficients after *oxdna2/dh* (T=0.1, rhos=0.5, qeff=0.815 in the above example). When using a Langevin thermostat e.g. through *fix langevin* or *fix nve/dotc/langevin* the temperature coefficients have to be matched to the one used in the fix.

---

---

**Note:** These pair styles have to be used with the *atom\_style hybrid bond ellipsoid oxdna* (see documentation of *atom\_style*). The *atom\_style oxdna* stores the 3'-to-5' polarity of the nucleotide strand, which is set through the bond topology in the data file. The first (second) atom in a bond definition is understood to point towards the 3'-end (5'-end) of the strand.

---

Example input and data files for DNA duplexes can be found in examples/PACKAGES/cgdna/examples/oxDNA/ and /oxDNA2/. A simple python setup tool which creates single straight or helical DNA strands, DNA duplexes or arrays of DNA duplexes can be found in examples/PACKAGES/cgdna/util/.

Please cite (*Henrich*) in any publication that uses this implementation. An updated documentation that contains general information on the model, its implementation and performance as well as the structure of the data and input file can be found [here](#).

Please cite also the relevant oxDNA2 publications (*Snodin*) and (*Sulc*).

---

### 19.201.4 Restrictions

These pair styles can only be used if LAMMPS was built with the CG-DNA package and the MOLECULE and AS-PHERE package. See the *Build package* page for more info.

## 19.201.5 Related commands

*bond\_style oxdna2/fene, pair\_coeff, bond\_style oxdna/fene, pair\_style oxdna/excv, bond\_style oxrna2/fene, pair\_style oxrna2/excv, atom\_style oxdna, fix nve/dotc/langevin*

## 19.201.6 Default

none

**(Henrich)** O. Henrich, Y. A. Gutierrez-Fosado, T. Curk, T. E. Ouldridge, Eur. Phys. J. E 41, 57 (2018).

**(Snodin)** B.E. Snodin, F. Randisi, M. Mosayebi, et al., J. Chem. Phys. 142, 234901 (2015).

**(Sulc)** P. Sulc, F. Romano, T.E. Ouldridge, L. Rovigatti, J.P.K. Doye, A.A. Louis, J. Chem. Phys. 137, 135101 (2012).

**(Ouldridge-DPhil)** T.E. Ouldridge, Coarse-grained modelling of DNA and DNA self-assembly, DPhil. University of Oxford (2011).

**(Ouldridge)** T.E. Ouldridge, A.A. Louis, J.P.K. Doye, J. Chem. Phys. 134, 085101 (2011).

## 19.202 pair\_style oxrna2/excv command

## 19.203 pair\_style oxrna2/stk command

## 19.204 pair\_style oxrna2/hbond command

## 19.205 pair\_style oxrna2/xstk command

## 19.206 pair\_style oxrna2/coaxstk command

## 19.207 pair\_style oxrna2/dh command

### 19.207.1 Syntax

```
pair_style style1
```

```
pair_coeff * * style2 args
```

- style1 = *hybrid/overlay oxrna2/excv oxrna2/stk oxrna2/hbond oxrna2/xstk oxrna2/coaxstk oxrna2/dh*
- style2 = *oxrna2/excv* or *oxrna2/stk* or *oxrna2/hbond* or *oxrna2/xstk* or *oxrna2/coaxstk* or *oxrna2/dh*
- args = list of arguments for these particular styles

```
oxrna2/stk args = seq T xi kappa 6.0 0.43 0.93 0.35 0.78 0.9 0 0.95 0.9 0 0.95 1.3 0 0.8
↪ 1.3 0 0.8 2.0 0.65 2.0 0.65
```

seq = seqav (for average sequence stacking strength) or seqdep (for sequence-dependent stacking strength)

T = temperature (oxDNA units, 0.1 = 300 K)

```

xi = 1.40206 (temperature-independent coefficient in stacking strength)
kappa = 2.77 (coefficient of linear temperature dependence in stacking strength)
oxrna2/hbond args = seq eps 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.5 0 0.7 0.46 3.
↳141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966 0.45
seq = seqav (for average sequence base-pairing strength) or seqdep (for
↳sequence-dependent base-pairing strength)
eps = 0.870439 (between base pairs A-T, C-G and G-T) or 0 (all other pairs)
oxrna2/dh args = T rhos qeff
T = temperature (oxDNA units, 0.1 = 300 K)
rhos = salt concentration (mole per litre)
qeff = 1.02455 (effective charge in elementary charges)

```

## 19.207.2 Examples

```

pair_style hybrid/overlay oxrna2/excv oxrna2/stk oxrna2/hbond oxrna2/xstk oxrna2/coaxstk
↳oxrna2/dh
pair_coeff * * oxrna2/excv      2.0 0.7 0.675 2.0 0.515 0.5 2.0 0.33 0.32
pair_coeff * * oxrna2/stk      seqdep 0.1 1.40206 2.77 6.0 0.43 0.93 0.35 0.78 0.9 0 0.95
↳0.9 0 0.95 1.3 0 0.8 1.3 0 0.8 2.0 0.65 2.0 0.65
pair_coeff * * oxrna2/hbond    seqdep 0.0 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7 1.5 0
↳0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.5707963267948966 0.45
pair_coeff 1 4 oxrna2/hbond    seqdep 0.870439 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7
↳1.5 0 0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.
↳5707963267948966 0.45
pair_coeff 2 3 oxrna2/hbond    seqdep 0.870439 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7
↳1.5 0 0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.
↳5707963267948966 0.45
pair_coeff 3 4 oxrna2/hbond    seqdep 0.870439 8.0 0.4 0.75 0.34 0.7 1.5 0 0.7 1.5 0 0.7
↳1.5 0 0.7 0.46 3.141592653589793 0.7 4.0 1.5707963267948966 0.45 4.0 1.
↳5707963267948966 0.45
pair_coeff * * oxrna2/xstk     59.9626 0.5 0.6 0.42 0.58 2.25 0.505 0.58 1.7 1.266 0.68 1.
↳7 1.266 0.68 1.7 0.309 0.68 1.7 0.309 0.68
pair_coeff * * oxrna2/coaxstk  80 0.5 0.6 0.42 0.58 2.0 2.592 0.65 1.3 0.151 0.8 0.9 0.
↳685 0.95 0.9 0.685 0.95 2.0 -0.65 2.0 -0.65
pair_coeff * * oxrna2/dh      0.1 0.5 1.02455

```

## 19.207.3 Description

The *oxrna2* pair styles compute the pairwise-additive parts of the oxDNA force field for coarse-grained modelling of DNA. The effective interaction between the nucleotides consists of potentials for the excluded volume interaction *oxrna2/excv*, the stacking *oxrna2/stk*, cross-stacking *oxrna2/xstk* and coaxial stacking interaction *oxrna2/coaxstk*, electrostatic Debye-Hueckel interaction *oxrna2/dh* as well as the hydrogen-bonding interaction *oxrna2/hbond* between complementary pairs of nucleotides on opposite strands. Average sequence or sequence-dependent stacking and base-pairing strengths are supported (*Sulc2*). Quasi-unique base-pairing between nucleotides can be achieved by using more complementary pairs of atom types like 5-8 and 6-7, 9-12 and 10-11, 13-16 and 14-15, etc. This prevents the hybridization of in principle complementary bases within  $N_{\text{types}}/4$  bases up and down along the backbone.

The exact functional form of the pair styles is rather complex. The individual potentials consist of products of modulation factors, which themselves are constructed from a number of more basic potentials (Morse, Lennard-Jones, harmonic angle and distance) as well as quadratic smoothing and modulation terms. We refer to (*Sulc1*) and the original oxDNA publications (*Ouldrige-DPhil*) and (*Ouldrige*) for a detailed description of the oxRNA2 force field.

---

**Note:** These pair styles have to be used together with the related oxDNA2 bond style *oxrna2/fene* for the connectivity of the phosphate backbone (see also documentation of *bond\_style oxrna2/fene*). Most of the coefficients in the above example have to be kept fixed and cannot be changed without reparameterizing the entire model. Exceptions are the first four coefficients after *oxrna2/stk* (*seq=seqdep*,  $T=0.1$ ,  $\xi=1.40206$  and  $\kappa=2.77$  in the above example), the first coefficient after *oxrna2/hbond* (*seq=seqdep* in the above example) and the three coefficients after *oxrna2/dh* ( $T=0.1$ ,  $\rho_{\text{hos}}=0.5$ ,  $q_{\text{eff}}=1.02455$  in the above example). When using a Langevin thermostat e.g. through *fix langevin* or *fix nve/dotc/langevin* the temperature coefficients have to be matched to the one used in the fix.

---

**Note:** These pair styles have to be used with the *atom\_style hybrid bond ellipsoid oxdna* (see documentation of *atom\_style*). The *atom\_style oxdna* stores the 3'-to-5' polarity of the nucleotide strand, which is set through the bond topology in the data file. The first (second) atom in a bond definition is understood to point towards the 3'-end (5'-end) of the strand.

---

Example input and data files for DNA duplexes can be found in `examples/PACKAGES/cgdna/examples/oxDNA/` and `/oxDNA2/`. A simple python setup tool which creates single straight or helical DNA strands, DNA duplexes or arrays of DNA duplexes can be found in `examples/PACKAGES/cgdna/util/`.

Please cite (*Henrich*) in any publication that uses this implementation. The article contains general information on the model, its implementation and performance as well as the structure of the data and input file. The preprint version of the article can be found [here](#). Please cite also the relevant oxRNA2 publications (*Sulc1*) and (*Sulc2*).

---

## 19.207.4 Restrictions

These pair styles can only be used if LAMMPS was built with the CG-DNA package and the MOLECULE and ASPHERE package. See the *Build package* page for more info.

## 19.207.5 Related commands

*bond\_style oxrna2/fene, pair\_coeff, bond\_style oxdna/fene, pair\_style oxdna/excv, bond\_style oxdna2/fene, pair\_style oxdna2/excv, atom\_style oxdna, fix nve/dotc/langevin*

## 19.207.6 Default

none

---

**(Henrich)** O. Henrich, Y. A. Gutierrez-Fosado, T. Curk, T. E. Ouldridge, Eur. Phys. J. E 41, 57 (2018).

**(Sulc1)** P. Sulc, F. Romano, T. E. Ouldridge, et al., J. Chem. Phys. 140, 235102 (2014).

**(Sulc2)** P. Sulc, F. Romano, T.E. Ouldridge, L. Rovigatti, J.P.K. Doye, A.A. Louis, J. Chem. Phys. 137, 135101 (2012).

**(Ouldridge-DPhil)** T.E. Ouldridge, Coarse-grained modelling of DNA and DNA self-assembly, DPhil. University of Oxford (2011).

**(Ouldridge)** T.E. Ouldridge, A.A. Louis, J.P.K. Doye, J. Chem. Phys. 134, 085101 (2011).

## 19.208 pair\_style pace command

Accelerator Variants: *pace/kk*

### 19.208.1 Syntax

```
pair_style pace ... keyword values ...
```

- one or more keyword/value pairs may be appended

keyword = *product* or *recursive* or *chunksize*

*product* = use product algorithm for basis functions

*recursive* = use recursive algorithm for basis functions

*chunksize* value = number of atoms in each pass

### 19.208.2 Examples

```
pair_style pace
pair_style pace product chunksize 2048
pair_coeff * * Cu-PBE-core-rep.ace Cu
```

### 19.208.3 Description

Pair style *pace* computes interactions using the Atomic Cluster Expansion (ACE), which is a general expansion of the atomic energy in multi-body basis functions. (*Drautz*). The *pace* pair style provides an efficient implementation that is described in this paper (*Lysogorskiy*).

In ACE, the total energy is decomposed into a sum over atomic energies. The energy of atom *i* is expressed as a linear or non-linear function of one or more density functions. By projecting the density onto a local atomic base, the lowest order contributions to the energy can be expressed as a set of scalar polynomials in basis function contributions summed over neighbor atoms.

Only a single *pair\_coeff* command is used with the *pace* style which specifies an ACE coefficient file followed by N additional arguments specifying the mapping of ACE elements to LAMMPS atom types, where N is the number of LAMMPS atom types:

- ACE coefficient file
- N element names = mapping of ACE elements to atom types

Only a single *pair\_coeff* command is used with the *pace* style which specifies an ACE file that fully defines the potential. Note that unlike for other potentials, cutoffs are not set in the *pair\_style* or *pair\_coeff* command; they are specified in the ACE file.

The *pair\_style pace* command may be followed by the optional keyword *product* or *recursive*, which determines which of two algorithms is used for the calculation of basis functions and derivatives. The default is *recursive*.

The keyword *chunksize* is only applicable when using the pair style *pace* with the KOKKOS package on GPUs and is ignored otherwise. This keyword controls the number of atoms in each pass used to compute the atomic cluster expansion and is used to avoid running out of memory. For example if there are 8192 atoms in the simulation and the *chunksize* is set to 4096, the ACE calculation will be broken up into two passes (running on a single GPU).

See the *pair\_coeff* page for alternate ways to specify the path for the ACE coefficient file.

### 19.208.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, mixing is performed by LAMMPS with user-specifiable parameters as described above. You never need to specify a `pair_coeff` command with I != J arguments for this style.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.208.5 Restrictions

This pair style is part of the ML-PACE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.208.6 Related commands

*pair\_style snap*

### 19.208.7 Default

recursive, chunksize = 4096

**(Drautz)** Drautz, Phys Rev B, 99, 014104 (2019).

**(Lysogorskiy)** Lysogorskiy, van der Oord, Bochkarev, Menon, Rinaldi, Hammerschmidt, Mrovec, Thompson, Csanyi, Ortner, Drautz, TBD (2021).

## 19.209 pair\_style peri/pmb command

Accelerator Variants: *peri/pmb/omp*

## 19.210 pair\_style peri/lps command

Accelerator Variants: *peri/lps/omp*

## 19.211 pair\_style peri/ves command

## 19.212 pair\_style peri/eps command

### 19.212.1 Syntax

```
pair_style style
```

- style = *peri/pmb* or *peri/lps* or *peri/ves* or *peri/eps*

### 19.212.2 Examples

```
pair_style peri/pmb
pair_coeff * * 1.6863e22 0.0015001 0.0005 0.25

pair_style peri/lps
pair_coeff * * 14.9e9 14.9e9 0.0015001 0.0005 0.25

pair_style peri/ves
pair_coeff * * 14.9e9 14.9e9 0.0015001 0.0005 0.25 0.5 0.001

pair_style peri/eps
pair_coeff * * 14.9e9 14.9e9 0.0015001 0.0005 0.25 118.43
```

### 19.212.3 Description

The peridynamic pair styles implement material models that can be used at the mesoscopic and macroscopic scales. See [this document](#) for an overview of LAMMPS commands for Peridynamics modeling.

Style *peri/pmb* implements the Peridynamic bond-based prototype microelastic brittle (PMB) model.

Style *peri/lps* implements the Peridynamic state-based linear peridynamic solid (LPS) model.

Style *peri/ves* implements the Peridynamic state-based linear peridynamic viscoelastic solid (VES) model.

Style *peri/eps* implements the Peridynamic state-based elastic-plastic solid (EPS) model.

The canonical papers on Peridynamics are (*Silling 2000*) and (*Silling 2007*). The implementation of Peridynamics in LAMMPS is described in (*Parks*). Also see the [PDLAMMPS user guide](#) for more details about its implementation.



The peridynamic VES and EPS models in PDLAMMPS were implemented by R. Rahman and J. T. Foster at University of Texas at San Antonio. The original VES formulation is described in “(Mitchell2011)” and the original EPS formulation is in “(Mitchell2011a)”. Additional PDF docs that describe the VES and EPS implementations are include in the LAMMPS distribution in [doc/PDF/PDLammmps\\_VES.pdf](#) and [doc/PDF/PDLammmps\\_EPS.pdf](#). For questions regarding the VES and EPS models in LAMMPS you can contact R. Rahman ([rezwanur.rahman@utsa.edu](mailto:rezwanur.rahman@utsa.edu)).

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below.

For the *peri/pmb* style:

- *c* (energy/distance/volume<sup>2</sup> units)
- horizon (distance units)
- *s00* (unitless)
- $\alpha$  (unitless)

*C* is the effectively a spring constant for Peridynamic bonds, the horizon is a cutoff distance for truncating interactions, and *s00* and  $\alpha$  are used as a bond breaking criteria. The units of *c* are such that  $c/\text{distance} = \text{stiffness}/\text{volume}^2$ , where stiffness is energy/distance<sup>2</sup> and volume is distance<sup>3</sup>. See the users guide for more details.

For the *peri/lps* style:

- *K* (force/area units)
- *G* (force/area units)
- horizon (distance units)
- *s00* (unitless)
- $\alpha$  (unitless)

*K* is the bulk modulus and *G* is the shear modulus. The horizon is a cutoff distance for truncating interactions, and *s00* and  $\alpha$  are used as a bond breaking criteria. See the users guide for more details.

For the *peri/ves* style:

- *K* (force/area units)
- *G* (force/area units)
- horizon (distance units)
- *s00* (unitless)
- $\alpha$  (unitless)
- *m\_lambdai* (unitless)
- *m\_taubi* (unitless)

*K* is the bulk modulus and *G* is the shear modulus. The horizon is a cutoff distance for truncating interactions, and *s00* and  $\alpha$  are used as a bond breaking criteria. *m\_lambdai* and *m\_taubi* are the viscoelastic relaxation parameter and time constant, respectively. *m\_lambdai* varies within zero to one. For very small values of *m\_lambdai* the viscoelastic model responds very similar to a linear elastic model. For details please see the description in “(Mitchell2011)”.

For the *peri/eps* style:

- *K* (force/area units)
- *G* (force/area units)
- horizon (distance units)

- $s00$  (unitless)
- $\alpha$  (unitless)
- $m\_yield\_stress$  (force/area units)

$K$  is the bulk modulus and  $G$  is the shear modulus. The horizon is a cutoff distance and  $s00$  and  $\alpha$  are used as a bond breaking criteria.  $m\_yield\_stress$  is the yield stress of the material. For details please see the description in “(Mitchell2011a)”.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.212.4 Mixing, shift, table, tail correction, restart, rRESPA info

These pair styles do not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

These pair styles do not support the *pair\_modify* shift option.

The *pair\_modify* table and tail options are not relevant for these pair styles.

These pair styles write their information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

These pair styles can only be used via the *pair* keyword of the *run\_style respa* command. They do not support the *inner*, *middle*, *outer* keywords.

---

#### 19.212.5 Restrictions

All of these styles are part of the PERI package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

#### 19.212.6 Related commands

*pair\_coeff*

## 19.212.7 Default

none

**(Parks)** Parks, Lehoucq, Plimpton, Silling, Comp Phys Comm, 179(11), 777-783 (2008).

**(Silling 2000)** Silling, J Mech Phys Solids, 48, 175-209 (2000).

**(Silling 2007)** Silling, Epton, Weckner, Xu, Askari, J Elasticity, 88, 151-184 (2007).

**(Mitchell2011)** Mitchell. A non-local, ordinary-state-based viscoelasticity model for peridynamics. Sandia National Lab Report, 8064:1-28 (2011).

**(Mitchell2011a)** Mitchell. A Nonlocal, Ordinary, State-Based Plasticity Model for Peridynamics. Sandia National Lab Report, 3166:1-34 (2011).

## 19.213 pair\_style polymorphic command

### 19.213.1 Syntax

```
pair_style polymorphic
```

style = *polymorphic*

### 19.213.2 Examples

```
pair_style polymorphic
pair_coeff * * FeCH_BOP_I.poly Fe C H
pair_coeff * * TlBr_msw.poly Tl Br
pair_coeff * * CuTa_eam.poly Cu Ta
pair_coeff * * GaN_tersoff.poly Ga N
pair_coeff * * GaN_sw.poly Ga N
```

### 19.213.3 Description

The *polymorphic* pair style computes a 3-body free-form potential (*Zhou3*) for the energy  $E$  of a system of atoms as

$$E = \frac{1}{2} \sum_{i=1}^{i=N} \sum_{j=1}^{j=N} [(1 - \delta_{ij}) \cdot U_{IJ}(r_{ij}) - (1 - \eta_{ij}) \cdot F_{IJ}(X_{ij}) \cdot V_{IJ}(r_{ij})]$$

$$X_{ij} = \sum_{k=i_1, k \neq j}^{i_N} W_{IK}(r_{ik}) \cdot G_{JIK}(\cos \theta_{jik}) \cdot P_{JIK}(\Delta r_{jik})$$

$$\Delta r_{jik} = r_{ij} - \xi_{IJ} \cdot r_{ik}$$

where I, J, K represent species of atoms i, j, and k,  $i_1, \dots, i_N$  represents a list of  $i$ 's neighbors,  $\delta_{ij}$  is a Dirac constant (i.e.,  $\delta_{ij} = 1$  when  $i = j$ , and  $\delta_{ij} = 0$  otherwise),  $\eta_{ij}$  is similar constant that can be set either to  $\eta_{ij} = \delta_{ij}$  or  $\eta_{ij} = 1 - \delta_{ij}$  depending on the potential type,  $U_{IJ}(r_{ij})$ ,  $V_{IJ}(r_{ij})$ ,  $W_{IK}(r_{ik})$  are pair functions,  $G_{JIK}(\cos \theta_{jik})$  is an angular function,  $P_{JIK}(\Delta r_{jik})$  is a function of atomic spacing differential  $\Delta r_{jik} = r_{ij} - \xi_{IJ} \cdot r_{ik}$  with  $\xi_{IJ}$  being a pair-dependent parameter, and  $F_{IJ}(X_{ij})$  is a function of the local environment variable  $X_{ij}$ . This generic potential

is fully defined once the constants  $\eta_{ij}$  and  $\xi_{IJ}$ , and the six functions  $U_{IJ}(r_{ij})$ ,  $V_{IJ}(r_{ij})$ ,  $W_{IK}(r_{ik})$ ,  $G_{JIK}(\cos \theta_{jik})$ ,  $P_{JIK}(\Delta r_{jik})$ , and  $F_{IJ}(X_{ij})$  are given. Here LAMMPS uses a global parameter  $\eta$  to represent  $\eta_{ij}$ . When  $\eta = 1$ ,  $\eta_{ij} = 1 - \delta_{ij}$ , otherwise  $\eta_{ij} = \delta_{ij}$ . Additionally,  $\eta = 3$  indicates that the function  $P_{JIK}(\Delta r)$  depends on species I, J and K, otherwise  $P_{JIK}(\Delta r) = P_{IK}(\Delta r)$  only depends on species I and K. Note that these six functions are all one dimensional, and hence can be provided in a tabular form. This allows users to design different potentials solely based on a manipulation of these functions. For instance, the potential reduces to a Stillinger-Weber potential (*SW*) if we set

$$\begin{aligned}\eta_{ij} &= \delta_{ij} (\eta = 2 \text{ or } \eta = 0), \xi_{IJ} = 0 \\ U_{IJ}(r) &= A_{IJ} \cdot \epsilon_{IJ} \cdot \left(\frac{\sigma_{IJ}}{r}\right)^q \cdot \left[ B_{IJ} \cdot \left(\frac{\sigma_{IJ}}{r}\right)^{p-q} - 1 \right] \cdot \exp\left(\frac{\sigma_{IJ}}{r - a_{IJ} \cdot \sigma_{IJ}}\right) \\ V_{IJ}(r) &= \sqrt{\lambda_{IJ} \cdot \epsilon_{IJ}} \cdot \exp\left(\frac{\gamma_{IJ} \cdot \sigma_{IJ}}{r - a_{IJ} \cdot \sigma_{IJ}}\right) \\ F_{IJ}(X) &= -X \\ P_{JIK}(\Delta r) &= P_{IK}(\Delta r) = 1 \\ W_{IJ}(r) &= \sqrt{\lambda_{IJ} \cdot \epsilon_{IJ}} \cdot \exp\left(\frac{\gamma_{IJ} \cdot \sigma_{IJ}}{r - a_{IJ} \cdot \sigma_{IJ}}\right) \\ G_{JIK}(\cos \theta) &= \left(\cos \theta + \frac{1}{3}\right)^2\end{aligned}$$

The potential reduces to a Tersoff potential (*Tersoff* or *Albe*) if we set

$$\begin{aligned}\eta_{ij} &= \delta_{ij} (\eta = 2 \text{ or } \eta = 0), \xi_{IJ} = 1 \\ U_{IJ}(r) &= \frac{D_{e,IJ}}{S_{IJ} - 1} \cdot \exp\left[-\beta_{IJ} \sqrt{2S_{IJ}} (r - r_{e,IJ})\right] \cdot f_{c,IJ}(r) \\ V_{IJ}(r) &= \frac{S_{IJ} \cdot D_{e,IJ}}{S_{IJ} - 1} \cdot \exp\left[-\beta_{IJ} \sqrt{\frac{2}{S_{IJ}}} (r - r_{e,IJ})\right] \cdot f_{c,IJ}(r) \\ F_{IJ}(X) &= (1 + X)^{-\frac{1}{2}} \\ P_{JIK}(\Delta r) &= P_{IK}(\Delta r) = \exp(2\mu_{IK} \cdot \Delta r) \\ W_{IJ}(r) &= f_{c,IJ}(r) \\ G_{JIK}(\cos \theta) &= \gamma_{IK} \left[ 1 + \frac{c_{IK}^2}{d_{IK}^2} - \frac{c_{IK}^2}{d_{IK}^2 + (h_{IK} + \cos \theta)^2} \right]\end{aligned}$$

where

$$f_{c,IJ}(r) = \begin{cases} 1, & r \leq R_{IJ} - D_{IJ} \\ \frac{1}{2} + \frac{1}{2} \cos\left[\frac{\pi(r + D_{IJ} - R_{IJ})}{2D_{IJ}}\right], & R_{IJ} - D_{IJ} < r < R_{IJ} + D_{IJ} \\ 0, & r \geq R_{IJ} + D_{IJ} \end{cases}$$

The potential reduces to a modified Stillinger-Weber potential (*Zhou3*) if we set

$$\begin{aligned}\eta_{ij} &= \delta_{ij} (\eta = 2 \text{ or } \eta = 0), \xi_{IJ} = 0 \\ U_{IJ}(r) &= \varphi_{R,IJ}(r) - \varphi_{A,IJ}(r) \\ V_{IJ}(r) &= u_{IJ}(r) \\ F_{IJ}(X) &= -X \\ P_{JIK}(\Delta r) &= P_{IK}(\Delta r) = 1 \\ W_{IJ}(r) &= u_{IJ}(r) \\ G_{JIK}(\cos \theta) &= g_{JIK}(\cos \theta)\end{aligned}$$

The potential reduces to a Rockett-Tersoff potential (*Wang3*) if we set

$$\begin{aligned}
 \eta_{ij} &= \delta_{ij} (\eta = 2 \text{ or } \eta = 0), \xi_{IJ} = 1 \\
 U_{IJ}(r) &= A_{IJ} \exp(-\lambda_{1,IJ} \cdot r) f_{c,IJ}(r) f_{ca,IJ}(r) \\
 V_{IJ}(r) &= \left\{ \begin{array}{l} B_{IJ} \exp(-\lambda_{2,IJ} \cdot r) f_{c,IJ}(r) + \\ A_{IJ} \exp(-\lambda_{1,IJ} \cdot r) f_{c,IJ}(r) [1 - f_{ca,IJ}(r)] \end{array} \right\} \\
 F_{IJ}(X) &= [1 + (\beta_{IJ} X)^{n_{IJ}}]^{-\frac{1}{2n_{IJ}}} \\
 P_{JIK}(\Delta r) &= P_{IK}(\Delta r) = \exp(\lambda_{3,IK} \cdot \Delta r^3) \\
 W_{IJ}(r) &= f_{c,IJ}(r) \\
 G_{JIK}(\cos \theta) &= 1 + \frac{c_{IK}^2}{d_{IK}^2} - \frac{c_{IK}^2}{d_{IK}^2 + (h_{IK} + \cos \theta)^2}
 \end{aligned}$$

where  $f_{ca,IJ}(r)$  is similar to the  $f_{c,IJ}(r)$  defined above:

$$f_{ca,IJ}(r) = \begin{cases} 1, & r \leq R_{a,IJ} - D_{a,IJ} \\ \frac{1}{2} + \frac{1}{2} \cos \left[ \frac{\pi(r + D_{a,IJ} - R_{a,IJ})}{2D_{a,IJ}} \right], & R_{a,IJ} - D_{a,IJ} < r < R_{a,IJ} + D_{a,IJ} \\ 0, & r \geq R_{a,IJ} + D_{a,IJ} \end{cases}$$

The potential becomes the embedded atom method (*Daw*) if we set

$$\begin{aligned}
 \eta_{ij} &= 1 - \delta_{ij} (\eta = 1), \xi_{IJ} = 0 \\
 U_{IJ}(r) &= \phi_{IJ}(r) \\
 V_{IJ}(r) &= 1 \\
 F_{IJ}(X) &= -2F_I(X) \\
 P_{JIK}(\Delta r) &= P_{IK}(\Delta r) = 1 \\
 W_{IJ}(r) &= f_J(r) \\
 G_{JIK}(\cos \theta) &= 1
 \end{aligned}$$

In the embedded atom method case,  $\phi_{IJ}(r)$  is the pair energy,  $F_I(X)$  is the embedding energy,  $X$  is the local electron density, and  $f_J(r)$  is the atomic electron density function.

The potential reduces to another type of Tersoff potential (*Zhou4*) if we set

$$\begin{aligned}
 \eta_{ij} &= \delta_{ij} (\eta = 3), \xi_{IJ} = 1 \\
 U_{IJ}(r) &= \frac{D_{e,IJ}}{S_{IJ} - 1} \cdot \exp \left[ -\beta_{IJ} \sqrt{2S_{IJ}} (r - r_{e,IJ}) \right] \cdot f_{c,IJ}(r) \cdot T_{IJ}(r) + V_{ZBL,IJ}(r) [1 - T_{IJ}(r)] \\
 V_{IJ}(r) &= \frac{S_{IJ} \cdot D_{e,IJ}}{S_{IJ} - 1} \cdot \exp \left[ -\beta_{IJ} \sqrt{\frac{2}{S_{IJ}}} (r - r_{e,IJ}) \right] \cdot f_{c,IJ}(r) \cdot T_{IJ}(r) \\
 F_{IJ}(X) &= (1 + X)^{-\frac{1}{2}} \\
 P_{JIK}(\Delta r) &= \omega_{JIK} \cdot \exp(\alpha_{JIK} \cdot \Delta r) \\
 W_{IJ}(r) &= f_{c,IJ}(r) \\
 G_{JIK}(\cos \theta) &= \gamma_{JIK} \left[ 1 + \frac{c_{JIK}^2}{d_{JIK}^2} - \frac{c_{JIK}^2}{d_{JIK}^2 + (h_{JIK} + \cos \theta)^2} \right] \\
 T_{IJ}(r) &= \frac{1}{1 + \exp[-b_{f,IJ}(r - r_{f,IJ})]} \\
 V_{ZBL,IJ}(r) &= 14.4 \cdot \frac{Z_I \cdot Z_J}{r} \sum_{k=1}^4 \mu_k \cdot \exp[-\nu_k (Z_I^{0.23} + Z_J^{0.23}) r]
 \end{aligned}$$

where  $f_{c,IJ}(r)$  is the same as defined above. This Tersoff potential differs from the one above because the  $P_{JIK}(\Delta r)$  function is now dependent on all three species I, J, and K.

If the tabulated functions are created using the parameters of Stillinger-Weber, Tersoff, and EAM potentials, the polymorphic pair style will produce the same global properties (energies and stresses) and the same forces as the *sw*, *tersoff*, and *eam* pair styles. The polymorphic pair style also produces the same per-atom properties (energies and stresses) as the corresponding *tersoff* and *eam* pair styles. However, due to a different partitioning of global properties to per-atom properties, the polymorphic pair style will produce different per-atom properties (energies and stresses) as the *sw* pair style. This does not mean that polymorphic pair style is different from the *sw* pair style. It just means that the definitions of the atom energies and atom stresses are different.

Only a single `pair_coeff` command is used with the polymorphic pair style which specifies a potential file for all needed elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of polymorphic potential elements to atom types

See the `pair_coeff` page for alternate ways to specify the path for the potential file. Several files for polymorphic potentials are included in the potentials directory of the LAMMPS distribution. They have a “poly” suffix.

As an example, imagine the `GaN_tersoff.poly` file has tabulated functions for Ga-N tersoff potential. If your LAMMPS simulation has 4 atom types and you want the first 3 to be Ga, and the fourth to be N, you would use the following `pair_coeff` command:

```
pair_coeff * * GaN_tersoff.poly Ga Ga Ga N
```

The first two arguments must be `**` to span all pairs of LAMMPS atom types. The first three `Ga` arguments map LAMMPS atom types 1,2,3 to the `Ga` element in the polymorphic file. The final `N` argument maps LAMMPS atom type 4 to the `N` element in the polymorphic file. If a mapping value is specified as `NULL`, the mapping is not performed. This can be used when an polymorphic potential is used as part of the hybrid pair style. The `NULL` values are placeholders for atom types that will be used with other potentials.

Potential files in the potentials directory of the LAMMPS distribution have a “.poly” suffix. At the beginning of the files, an unlimited number of lines starting with `#` are used to describe the potential and are ignored by LAMMPS. The next line lists two numbers:

```
nypes eta
```

Here *nypes* represent total number of species defined in the potential file,  $\eta = 1$  reduces to embedded atom method,  $\eta = 3$  assumes a three species dependent  $P_{JIK}(\Delta r)$  function, and all other values of  $\eta$  assume a two species dependent  $P_{JK}(\Delta r)$  function. The value of *nypes* must equal the total number of different species defined in the `pair_coeff` command. The next *nypes* lines each lists two numbers and a character string representing atomic number, atomic mass, and name of the species of the *nypes* elements:

```
atomic-number atomic-mass element-name(1)
atomic-number atomic-mass element-name(2)
...
atomic-number atomic-mass element-name(nypes)
```

The next line contains four numbers:

```
nr ntheta nx xmax
```

Here *nr* is total number of tabular points for radial functions *U*, *V*, *W*, *P*, *ntheta* is total number of tabular points for the angular function *G*, *nx* is total number of tabular points for the function *F*, *xmax* is a maximum value of the argument of function *F*. Note that the pair functions  $U_{IJ}(r)$ ,  $V_{IJ}(r)$ ,  $W_{IJ}(r)$  are uniformly tabulated between 0 and cutoff distance of the *IJ* pair,  $G_{JIK}(\cos \theta)$  is uniformly tabulated between -1 and 1,  $P_{JIK}(\Delta r)$  is uniformly tabulated between -*rcmax* and *rcmax* where *rcmax* is the maximum cutoff distance of all pairs, and  $F_{IJ}(X)$  is uniformly tabulated between 0 and *xmax*. Linear extrapolation is assumed if actual simulations exceed these ranges.

The next  $n_{\text{types}}*(n_{\text{types}}+1)/2$  lines contain two numbers:

```
cut xi(1)
cut xi(2)
...
cut xi(ntypes*(ntypes+1)/2)
```

Here cut means the cutoff distance of the pair functions, “xi” is  $\xi$  as defined in the potential functions above. The  $n_{\text{types}}*(n_{\text{types}}+1)/2$  lines are related to the pairs according to the sequence of first ii (self) pairs,  $i = 1, 2, \dots, n_{\text{types}}$ , and then ij (cross) pairs,  $i = 1, 2, \dots, n_{\text{types}}-1$ , and  $j = i+1, i+2, \dots, n_{\text{types}}$  (i.e., the sequence of the ij pairs follows 11, 22, ..., 12, 13, 14, ..., 23, 24, ...).

In the final blocks of the potential file, U, V, W, P, G, and F functions are listed sequentially. First, U functions are given for each of the  $n_{\text{types}}*(n_{\text{types}}+1)/2$  pairs according to the sequence described above. For each of the pairs, nr values are listed. Next, similar arrays are given for V and W functions. If P functions depend only on pair species, i.e.,  $\eta \neq 3$ , then P functions are also listed the same way the next. If P functions depend on three species, i.e.,  $\eta = 3$ , then P functions are listed for all the  $n_{\text{types}}*n_{\text{types}}*n_{\text{types}}$  IJK triplets in a natural sequence I from 1 to  $n_{\text{types}}$ , J from 1 to  $n_{\text{types}}$ , and K from 1 to  $n_{\text{types}}$  (i.e., IJK = 111, 112, 113, ..., 121, 122, 123 ..., 211, 212, ...). Next, G functions are listed for all the  $n_{\text{types}}*n_{\text{types}}*n_{\text{types}}$  IJK triplets similarly. For each of the G functions, ntheta values are listed. Finally, F functions are listed for all the  $n_{\text{types}}*(n_{\text{types}}+1)/2$  pairs in the same sequence as described above. For each of the F functions, nx values are listed.

#### 19.213.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write their information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

#### 19.213.5 Restrictions

If using *create\_atoms* command, atomic masses must be defined in the input script. If using *read\_data*, atomic masses must be defined in the atomic structure data file.

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair potential requires the *newton* setting to be “on” for pair interactions.

The potential files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use any LAMMPS units, but you would need to create your own potential files.

#### 19.213.6 Related commands

*pair\_coeff*

(Zhou3) X. W. Zhou, M. E. Foster, R. E. Jones, P. Yang, H. Fan, and F. P. Doty, J. Mater. Sci. Res., 4, 15 (2015).

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## 19.214 pair\_style python command

### 19.214.1 Syntax

```
pair_style python cutoff
```

cutoff = global cutoff for interactions in python potential classes

### 19.214.2 Examples

```
pair_style python 2.5
pair_coeff * * py_pot.LJCutMelt lj

pair_style python 10.0
pair_coeff * * py_pot.HarmonicCut A B

pair_style hybrid/overlay coul/long 12.0 python 12.0
pair_coeff * * coul/long
pair_coeff * * python py_pot.LJCutSPCE OW NULL
```

### 19.214.3 Description

The *python* pair style provides a way to define pairwise additive potential functions as python script code that is loaded into LAMMPS from a python file which must contain specific python class definitions. This allows to rapidly evaluate different potential functions without having to modify and re-compile LAMMPS. Due to python being an interpreted language, however, the performance of this pair style is going to be significantly slower (often between 20x and 100x) than corresponding compiled code. This penalty can be significantly reduced through generating tabulations from the python code through the *pair\_write* command, which is supported by this style.

Only a single *pair\_coeff* command is used with the *python* pair style which specifies a python class inside a python module or a file that LAMMPS will look up in the current directory, a folder pointed to by the LAMMPS\_POTENTIALS environment variable or somewhere in your python path. A single python module can hold multiple python pair class definitions. The class definitions itself have to follow specific rules that are explained below.

Atom types in the python class are specified through symbolic constants, typically strings. These are mapped to LAMMPS atom types by specifying N additional arguments after the class name in the *pair\_coeff* command, where N must be the number of currently defined atom types:

As an example, imagine a file *py\_pot.py* has a python potential class names *LJCutMelt* with parameters and potential functions for a two Lennard-Jones atom types labeled as ‘LJ1’ and ‘LJ2’. In your LAMMPS input and you would have defined 3 atom types, out of which the first two are supposed to be using the ‘LJ1’ parameters and the third the ‘LJ2’ parameters, then you would use the following *pair\_coeff* command:

```
pair_coeff * * py_pot.LJCutMelt LJ1 LJ1 LJ2
```



The first two arguments **must** be `**` so as to span all LAMMPS atom types. The first two LJ1 arguments map LAMMPS atom types 1 and 2 to the LJ1 atom type in the `LJCutMelt` class of the `py_pot.py` file. The final LJ2 argument maps LAMMPS atom type 3 to the LJ2 atom type the python file. If a mapping value is specified as `NULL`, the mapping is not performed, any pair interaction with this atom type will be skipped. This can be used when a *python* potential is used as part of the *hybrid* or *hybrid/overlay* pair style. The `NULL` values are then placeholders for atom types that will be used with other potentials.

The python potential file has to start with the following code:

```
from __future__ import print_function

class LAMMSPairPotential(object):
    def __init__(self):
        self.pmap=dict()
        self.units='lj'
    def map_coeff(self,name,ltype):
        self.pmap[ltype]=name
    def check_units(self,units):
        if (units != self.units):
            raise Exception("Conflicting units: %s vs. %s" % (self.units,units))
```

Any classes with definitions of specific potentials have to be derived from this class and should be initialize in a similar fashion to the example given below.

**Note:** The class constructor has to set up a data structure containing the potential parameters supported by this class. It should also define a variable `self.units` containing a string matching one of the options of LAMMPS' `units` command, which is used to verify, that the potential definition in the python class and in the LAMMPS input match.

Here is an example for a single type Lennard-Jones potential class `LJCutMelt` in reduced units, which defines an atom type `lj` for which the parameters epsilon and sigma are both 1.0:

```
class LJCutMelt(LAMMSPairPotential):
    def __init__(self):
        super(LJCutMelt,self).__init__()
        # set coeffs: 48*eps*sig**12, 24*eps*sig**6,
        #              4*eps*sig**12, 4*eps*sig**6
        self.units = 'lj'
        self.coeff = {'lj' : {'lj' : (48.0,24.0,4.0,4.0)}}
```

The class also has to provide two methods for the computation of the potential energy and forces, which have been named `compute_force`, and `compute_energy`, which both take 3 numerical arguments:

- `rsq` = the square of the distance between a pair of atoms (float)
- `itype` = the (numerical) type of the first atom
- `jtype` = the (numerical) type of the second atom

These functions need to compute the (scaled) force and the energy, respectively, and use the result as return value. The functions need to use the `pmap` dictionary to convert the LAMMPS atom type number to the symbolic value of the internal potential parameter data structure. Following the `LJCutMelt` example, here are the two functions:

```

def compute_force(self,rsq,itype,jtype):
    coeff = self.coeff[self.pmap[itype]][self.pmap[jtype]]
    r2inv = 1.0/rsq
    r6inv = r2inv*r2inv*r2inv
    lj1 = coeff[0]
    lj2 = coeff[1]
    return (r6inv * (lj1*r6inv - lj2))*r2inv

def compute_energy(self,rsq,itype,jtype):
    coeff = self.coeff[self.pmap[itype]][self.pmap[jtype]]
    r2inv = 1.0/rsq
    r6inv = r2inv*r2inv*r2inv
    lj3 = coeff[2]
    lj4 = coeff[3]
    return (r6inv * (lj3*r6inv - lj4))

```

**Note:** for consistency with the C++ pair styles in LAMMPS, the `compute_force` function follows the conventions of the `Pair::single()` methods and does not return the pairwise force directly, but the force divided by the distance between the two atoms, so this value only needs to be multiplied by delta x, delta y, and delta z to conveniently obtain the three components of the force vector between these two atoms.

Below is a more complex example using *real* units and defines an interaction equivalent to:

```

units real
pair_style harmonic/cut
pair_coeff 1 1 0.2 9.0
pair_coeff 2 2 0.4 9.0

```

This uses the default geometric mixing. The equivalent input with pair style *python* is:

```

units real
pair_style python 10.0
pair_coeff * * py_pot.Harmonic A B

```

Note that while for pair style *harmonic/cut* the cutoff is implicitly set to the minimum of the harmonic potential, for pair style *python* a global cutoff must be set and it must be equal or larger to the implicit cutoff of the potential in *python*, which has to explicitly return zero force and energy beyond the cutoff. Also, the mixed parameters have to be explicitly provided. The corresponding *python* code is:

```

class Harmonic(LAMMPSPairPotential):
    def __init__(self):
        super(Harmonic,self).__init__()
        self.units = 'real'
        # set coeffs: K, r0
        self.coeff = {'A' : {'A' : (0.2,9.0),
                             'B' : (math.sqrt(0.2*0.4),9.0)},
                     'B' : {'A' : (math.sqrt(0.2*0.4),9.0),
                             'B' : (0.4,9.0)}}

    def compute_force(self,rsq,itype,jtype):
        coeff = self.coeff[self.pmap[itype]][self.pmap[jtype]]
        r = math.sqrt(rsq)

```

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```

delta = coeff[1]-r
if (r < coeff[1]):
    return 2.0*delta*coeff[0]/r
else:
    return 0.0

def compute_energy(self,rsq,itpe,jtpe):
    coeff = self.coeff[self.pmap[itpe]][self.pmap[jtpe]]
    r = math.sqrt(rsq)
    delta = coeff[1]-r
    if (r < coeff[1]):
        return delta*delta*coeff[0]
    else:
        return 0.0

```

---

### Performance Impact

The evaluation of scripted python code will slow down the computation of pairwise interactions quite significantly. However, this performance penalty can be worked around through using the python pair style not for the actual simulation, but to generate tabulated potentials using the *pair\_write* command. This will also enable GPU or multi-thread acceleration through the GPU, KOKKOS, or OPENMP package versions of the *table* pair style. Please see below for a LAMMPS input example demonstrating how to build a table file:

```

pair_style python 2.5
pair_coeff * * py_pot.LJCutMelt lj
shell rm -f lj.table
pair_write 1 1 2000 rsq 0.01 2.5 lj.table lj

```

Note that it is strongly recommended to try to **delete** the potential table file before generating it. Since the *pair\_write* command will always **append** to a table file, while pair style table will use the **first match**. Thus when changing the potential function in the python class, the table pair style will still read the old variant unless the table file is first deleted.

After switching the pair style to *table*, the potential tables need to be assigned to the LAMMPS atom types like this:

```

pair_style      table linear 2000
pair_coeff      1 1  lj.table lj

```

This can also be done for more complex systems. Please see the *examples/python* folders for a few more examples.

### 19.214.4 Mixing, shift, table, tail correction, restart, rRESPA info

Mixing of potential parameters has to be handled inside the provided python module. The python pair style simply assumes that force and energy computation can be correctly performed for all pairs of atom types as they are mapped to the atom type labels inside the python potential class.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.214.5 Restrictions

This pair style is part of the PYTHON package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.214.6 Related commands

*pair\_coeff*, *pair\_write*, *pair style table*

### 19.214.7 Default

none

## 19.215 pair\_style quip command

### 19.215.1 Syntax

```
pair_style quip
```

### 19.215.2 Examples

```
pair_style      quip
pair_coeff      * * gap_example.xml "Potential xml_label=GAP_2014_5_8_60_17_10_38_466" 14
pair_coeff      * * sw_example.xml "IP SW" 14
```

### 19.215.3 Description

Style *quip* provides an interface for calling potential routines from the QUIP package. QUIP is built separately, and then linked to LAMMPS. The most recent version of the QUIP package can be downloaded from GitHub: <https://github.com/libAtoms/QUIP>. The interface is chiefly intended to be used to run Gaussian Approximation Potentials (GAP), which are described in the following publications: (*Bartok et al*) and (*PhD thesis of Bartok*).

Only a single *pair\_coeff* command is used with the *quip* style that specifies a QUIP potential file containing the parameters of the potential for all needed elements in XML format. This is followed by a QUIP initialization string. Finally, the QUIP elements are mapped to LAMMPS atom types by specifying N atomic numbers, where N is the number of LAMMPS atom types:

- QUIP filename
- QUIP initialization string
- N atomic numbers = mapping of QUIP elements to atom types

See the *pair\_coeff* page for alternate ways to specify the path for the potential file.

A QUIP potential is fully specified by the filename which contains the parameters of the potential in XML format, the initialization string, and the map of atomic numbers.

GAP potentials can be obtained from the Data repository section of <http://www.libatoms.org>, where the appropriate initialization strings are also advised. The list of atomic numbers must be matched to the LAMMPS atom types specified in the LAMMPS data file or elsewhere.

Two examples input scripts are provided in the examples/PACKAGES/quip directory.

#### 19.215.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* mix, shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

#### 19.215.5 Restrictions

This pair style is part of the ML-QUIP package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

QUIP potentials are parameterized in electron-volts and Angstroms and therefore should be used with LAMMPS metal *units*.

QUIP potentials are generally not designed to work with the scaling factors set by the *special\_bonds* command. The recommended setting in molecular systems is to include all interactions, i.e. to use *special\_bonds lj/coul 1.0 1.0 1.0*. Scaling factors > 0.0 will be ignored and treated as 1.0. The only exception to this rule is if you know that your QUIP potential needs to exclude bonded, 1-3, or 1-4 interactions and does not already do this exclusion within QUIP. Then a factor 0.0 needs to be used which will remove such pairs from the neighbor list. This needs to be very carefully tested, because it may remove pairs from the neighbor list that are still required.

#### 19.215.6 Related commands

*pair\_coeff*

---

**(Bartok\_2010)** AP Bartok, MC Payne, R Kondor, and G Csanyi, Physical Review Letters 104, 136403 (2010).

**(Bartok\_PhD)** A Bartok-Partay, PhD Thesis, University of Cambridge, (2010).

### 19.216 pair\_style rann command

#### 19.216.1 Syntax

```
pair_style rann
pair_coeff file Type1_element Type2_element Type3_element...
```

## 19.216.2 Examples

```
pair_style rann
pair_coeff * * Mg.rann Mg
pair_coeff * * MgAlalloy.rann Mg Mg Al Mg
```

## 19.216.3 Description

Pair style *rann* computes pairwise interactions for a variety of materials using rapid atomistic neural network (RANN) potentials (*Dickel*, *Nitol*). Neural network potentials work by first generating a series of symmetry functions i.e. structural fingerprints from the neighbor list and then using these values as the input layer of a neural network. There is a single output neuron in the final layer which is the energy. Atomic forces are found by analytical derivatives computed via back-propagation. For alloy systems, each element has a unique network.

## 19.216.4 Potential file syntax

The RANN potential is defined by a single text file which contains all the fitting parameters for the alloy system. The potential file also defines the active fingerprints, network architecture, activation functions, etc. The potential file is divided into several sections which are identified by one of the following keywords:

- atomtypes
- mass
- fingerprintsperelement
- fingerprints
- fingerprintconstants
- screening (optional)
- networklayers
- layersize
- weight
- bias
- activationfunctions
- calibrationparameters (ignored)

The '#' character is treated as a comment marker, similar to LAMMPS input scripts. Sections are not required to follow a rigid ordering, but do require previous definition of prerequisite information. E.g., fingerprintconstants for a particular fingerprint must follow the fingerprints definition; layersize for a particular layer must follow the declaration of network layers.

*atomtypes* are defined as follows using element keywords separated by spaces.

```
atomtypes:
Fe Mg Al etc.
```

*mass* must be specified for each element keyword as follows:

```

mass:Mg:
24.305
mass:Fe:
55.847
mass:Al:
26.982

```

*fingerprintsperelement* specifies how many fingerprints are active for computing the energy of a given atom. This number must be specified for each element keyword. Active elements for each fingerprint depend upon the type of the central atom and the neighboring atoms. Pairwise fingerprints may be defined for a Mg atom based exclusively on its Al neighbors, for example. Bond fingerprints may use two neighbor lists of different element types. In computing fingerprintsperelement from all defined fingerprints, only the fingerprints defined for atoms of a particular element should be considered, regardless of the elements used in its neighbor list. In the following code, for example, some fingerprints may compute pairwise fingerprints summing contributions about Fe atoms based on a neighbor list of exclusively Al atoms, but if there are no fingerprints summing contributions of all neighbors about a central Al atom, then fingerprintsperelement of Al is zero:

```

fingerprintsperelement:Mg:
5
fingerprintsperelement:Fe:
2
fingerprintsperelement:Al:
0

```

*fingerprints* specifies the active fingerprints for a certain element combination. Pair fingerprints are specified for two elements, while bond fingerprints are specified for three elements. Only one fingerprints header should be used for an individual combination of elements. The ordering of the fingerprints in the network input layer is determined by the order of element combinations specified by subsequent *fingerprints* lines, and the order of the fingerprints defined for each element combination. Multiple fingerprints of the same style or different ones may be specified. If the same style and element combination is used for multiple fingerprints, they should have different id numbers. The first element specifies the atoms for which this fingerprint is computed while the other(s) specify which atoms to use in the neighbor lists for the computation. Switching the second and third element type in bond fingerprints has no effect on the computation:

```

fingerprints:Mg_Mg:
radial_0 radialscreened_0 radial_1
fingerprints:Mg_Al_Fe:
bond_0 bondspin_0
fingerprints:Mg_Al:
radial_0 radialscreened_0

```

The following fingerprint styles are currently defined. See the *formulation section* below for their definitions:

- radial
- radialscreened
- radials핀
- radialscreenedspin
- bond
- bondscreened
- bondspin
- bondscreenedspin

*fingerprintconstants* specifies the meta-parameters for a defined fingerprint. For all radial styles, *re*, *rc*, *alpha*, *dr*, *o*, and *n* must be specified. *re* should usually be the stable interatomic distance, *rc* is the cutoff radius, *dr* is the cutoff smoothing distance, *o* is the lowest radial power term (which may be negative), and *n* is the highest power term. The total length of the fingerprint vector is  $(n-o+1)$ . *alpha* is a list of decay parameters used for exponential decay of radial contributions. It may be set proportionally to the bulk modulus similarly to MEAM potentials, but other values may be provided better fitting in special cases. Bond style fingerprints require specification of *re*, *rc*, *alphak*, *dr*, *k*, and *m*. Here *m* is the power of the bond cosines and *k* is the number of decay parameters. Cosine powers go from 0 to  $m-1$  and are each computed for all values of *alphak*. Thus the total length of the fingerprint vector is  $m*k$ .

```
fingerprintconstants:Mg_Mg:radialscreened_0:re:
3.193592
fingerprintconstants:Mg_Mg:radialscreened_0:rc:
6.000000
fingerprintconstants:Mg_Mg:radialscreened_0:alpha:
5.520000 5.520000 5.520000 5.520000 5.520000
fingerprintconstants:Mg_Mg:radialscreened_0:dr:
2.806408
fingerprintconstants:Mg_Mg:radialscreened_0:o:
-1
fingerprintconstants:Mg_Mg:radialscreened_0:n:
3
```

*screening* specifies the *Cmax* and *Cmin* values used in the screening fingerprints. Contributions from neighbors to the fingerprint are omitted if they are blocked by a closer neighbor, and reduced if they are partially blocked. Larger values of *Cmin* correspond to neighbors being blocked more easily. *Cmax* cannot be greater than 3, and *Cmin* cannot be greater than *Cmax* or less than zero. Screening may be omitted in which case the default values  $Cmax = 2.8$ ,  $Cmin = 0.8$  are used. Since screening is a bond computation, it is specified separately for each combination of three elements in which the latter two may be interchanged with no effect.

```
screening:Mg_Mg_Mg:Cmax:
2.700000
screening:Mg_Mg_Mg:Cmin:
0.400000
```

*networklayers* specifies the size of the neural network for each atom. It counts both the input and output layer and so is  $2 + \langle \text{hidden layers} \rangle$ .

```
networklayers:Mg:
3
```

*layersize* specifies the length of each layer, including the input layer and output layer. The input layer is layer 0. The size of the input layer size must match the summed length of all the fingerprints for that element, and the output layer size must be 1:

```
layersize:Mg:0:
14
layersize:Mg:1:
20
layersize:Mg:2:
1
```

*weight* specifies the weight for a given element and layer. Weight cannot be specified for the output layer. The weight of layer *i* is a  $m \times n$  matrix where *m* is the layer size of *i* and *n* is the layer size of *i+1*:



```
weight:Mg:0:
w11 w12 w13 ...
w21 w22 w23 ...
...
```

*bias* specifies the bias for a given element and layer. Bias cannot be specified for the output layer. The bias of layer *i* is a  $n \times 1$  vector where *n* is the layer size of *i*+1:

```
bias:Mg:0:
b1
b2
b3
...
```

*activationfunctions* specifies the activation function for a given element and layer. Activation functions cannot be specified for the output layer:

```
activationfunctions:Mg:0:
sigI
activationfunctions:Mg:1:
linear
```

The following activation styles are currently specified. See the [formulation section](#) below for their definitions.

- sigI
- linear

*calibrationparameters* specifies a number of parameters used to calibrate the potential. These are ignored by LAMMPS.

### 19.216.5 Formulation

In the RANN formulation, the total energy of a system of atoms is given by:

$$E = \sum_{\alpha} E^{\alpha}$$

$$E^{\alpha} = N A^{\alpha}$$

$${}^{n+1}A_i^{\alpha} = {}^nF ({}^nW_{ij} {}^nA_j^{\alpha} + {}^nB_i)$$

$${}^0A_i^{\alpha} = \begin{bmatrix} {}^1Sf^{\alpha} \\ {}^2Sf^{\alpha} \\ \dots \end{bmatrix}$$

Here  $E^{\alpha}$  is the energy of atom  $\alpha$ ,  ${}^nF()$ ,  ${}^nW_{ij}$  and  ${}^nB_i$  are the activation function, weight matrix and bias vector of the *n*-th layer respectively. The inputs to the first layer are a collection of structural fingerprints which are collected and reshaped into a single long vector. The individual fingerprints may be defined in any order and have various shapes and sizes. Multiple fingerprints of the same type and varying parameters may also be defined in the input layer.

Eight types of structural fingerprints are currently defined. In the following,  $\beta$  and  $\gamma$  span the full neighbor list of atom  $\alpha$ .  $\delta_i$  are decay meta-parameters, and  $r_e$  is a meta-parameter roughly proportional to the first neighbor distance.  $r_c$  and  $dr$  are the neighbor cutoff distance and cutoff smoothing distance respectively.  $S^{\alpha\beta}$  is the MEAM screening function

(*Baskes*),  $s_i^\alpha$  and  $s_i^\beta$  are the atom spin vectors (*Tranchida*).  $r^{\alpha\beta}$  is the distance from atom  $\alpha$  to atom  $\beta$ , and  $\theta^{\alpha\beta\gamma}$  is the bond angle:

$$\cos(\theta^{\alpha\beta\gamma}) = \frac{\mathbf{r}^{\alpha\beta} \cdot \mathbf{r}^{\alpha\gamma}}{r^{\alpha\beta} r^{\alpha\gamma}}$$

$S^{\alpha\beta}$  is defined as (*Baskes*):

$$X^{\gamma\beta} = \left(\frac{r^{\gamma\beta}}{r^{\alpha\beta}}\right)^2$$

$$X^{\alpha\gamma} = \left(\frac{r^{\alpha\gamma}}{r^{\alpha\beta}}\right)^2$$

$$C = \frac{2(X^{\alpha\gamma} + X^{\gamma\beta}) - (X^{\alpha\gamma} - X^{\gamma\beta})^2 - 1}{1 - (X^{\alpha\gamma} - X^{\gamma\beta})^2}$$

$$f_c(x) = \begin{cases} 1 & x \geq 1 \\ (1 - (1 - x)^4)^2 & 0 < x < 1 \\ 0 & x \leq 0 \end{cases}$$

$$S^{\alpha\beta\gamma} = f_c\left(\frac{C - C_{min}}{C_{max} - C_{min}}\right)$$

$$S^{\alpha\beta} = \prod_{\gamma} S^{\alpha\beta\gamma}$$

The structural fingerprints are computed as follows:

- **radial**

$$rSf_i^\alpha = \sum_{\beta} \left(\frac{r^{\alpha\beta}}{r_e}\right)^i e^{-\delta_i \frac{r^{\alpha\beta}}{r_e}} f_c\left(\frac{r_c - r^{\alpha\beta}}{dr}\right)$$

- **bond**

$$bSf_{ij}^\alpha = \sum_{\beta} \sum_{\gamma} (\cos(\theta_{\alpha\beta\gamma}))^i e^{-\delta_j \frac{r^{\alpha\beta}}{r_e}} e^{-\delta_j \frac{r^{\alpha\gamma}}{r_e}} f_c\left(\frac{r_c - r^{\alpha\beta}}{dr}\right) f_c\left(\frac{r_c - r^{\alpha\gamma}}{dr}\right)$$

- **radialscreened**

$$rscSf_i^\alpha = \sum_{\beta} \left(\frac{r^{\alpha\beta}}{r_e}\right)^i e^{-\delta_i \frac{r^{\alpha\beta}}{r_e}} S^{\alpha\beta} f_c\left(\frac{r_c - r^{\alpha\beta}}{dr}\right)$$

- **bondscreened**

$$bscSf_{ij}^\alpha = \sum_{\beta} \sum_{\gamma} (\cos(\theta_{\alpha\beta\gamma}))^i e^{-\delta_j \frac{r^{\alpha\beta}}{r_e}} e^{-\delta_j \frac{r^{\alpha\gamma}}{r_e}} S^{\alpha\beta} S^{\alpha\gamma} f_c\left(\frac{r_c - r^{\alpha\beta}}{dr}\right) f_c\left(\frac{r_c - r^{\alpha\gamma}}{dr}\right)$$

- **radialspin**

$$rspSf_i^\alpha = \sum_{\beta} \left(\frac{r^{\alpha\beta}}{r_e}\right)^i e^{-\delta_i \frac{r^{\alpha\beta}}{r_e}} (\mathbf{s}^\alpha \cdot \mathbf{s}^\beta) f_c\left(\frac{r_c - r^{\alpha\beta}}{dr}\right)$$

- **bondspin**

$$bspSf_{ij}^{\alpha} = \sum_{\beta} \sum_{\gamma} (\cos(\theta_{\alpha\beta\gamma}))^i e^{-\delta_j \frac{r^{\alpha\beta}}{r_e}} e^{-\delta_j \frac{r^{\alpha\gamma}}{r_e}} (\mathbf{s}^{\alpha} \cdot \mathbf{s}^{\beta}) (\mathbf{s}^{\alpha} \cdot \mathbf{s}^{\gamma}) f_c \left( \frac{r_c - r^{\alpha\beta}}{dr} \right) f_c \left( \frac{r_c - r^{\alpha\gamma}}{dr} \right)$$

- **radialscreenedspin**

$$rscspSf_i^{\alpha} = \sum_{\beta} \left( \frac{r^{\alpha\beta}}{r_e} \right)^i e^{-\delta_i \frac{r^{\alpha\beta}}{r_e}} S^{\alpha\beta} (\mathbf{s}^{\alpha} \cdot \mathbf{s}^{\beta}) f_c \left( \frac{r_c - r^{\alpha\beta}}{dr} \right)$$

- **bondscreenedspin**

$$bscspSf_{ij}^{\alpha} = \sum_{\beta} \sum_{\gamma} (\cos(\theta_{\alpha\beta\gamma}))^i e^{-\delta_j \frac{r^{\alpha\beta}}{r_e}} e^{-\delta_j \frac{r^{\alpha\gamma}}{r_e}} S^{\alpha\beta} S^{\alpha\gamma} (\mathbf{s}^{\alpha} \cdot \mathbf{s}^{\beta}) (\mathbf{s}^{\alpha} \cdot \mathbf{s}^{\gamma}) f_c \left( \frac{r_c - r^{\alpha\beta}}{dr} \right) f_c \left( \frac{r_c - r^{\alpha\gamma}}{dr} \right)$$

The activation functions are computed as follows:

- **sigI**

$$F^{sigI}(x) = 0.1x + 0.9 \ln(e^x + 1)$$

- **linear**

$$F^{linear}(x) = x$$

## 19.216.6 Restrictions

Pair style *rann* is part of the ML-RANN package. It is only enabled if LAMMPS was built with that package. Additionally, if any spin fingerprint styles are used LAMMPS must be built with the SPIN package as well.

Pair style *rann* does not support computing per-atom stress or using *pair\_modify nofdotr*.

## 19.216.7 Defaults

Cmin = 0.8, Cmax = 2.8.

**(Baskes)** Baskes, Materials Chemistry and Physics, 50(2), 152-158, (1997).

**(Dickel)** Dickel, Francis, and Barrett, Computational Materials Science 171 (2020): 109157.

**(Nitol)** Nitol, Dickel, and Barrett, Computational Materials Science 188 (2021): 110207.

**(Tranchida)** Tranchida, Plimpton, Thibaudau and Thompson, Journal of Computational Physics, 372, 406-425, (2018).

## 19.217 pair\_style reaxff command

Accelerator Variants: *reaxff/kk*, *reaxff/omp*

## 19.217.1 Syntax

```
pair_style reaxff cfile keyword value
```

- cfile = NULL or name of a control file
- zero or more keyword/value pairs may be appended

```
keyword = checkqeq or lgvdw or safezone or mincap or minhbonds or tabulate or list/
↳blocking
  checkqeq value = yes or no = whether or not to require qeq/reaxff or acks2/reaxff.
↳fix
  enobonds value = yes or no = whether or not to tally energy of atoms with no bonds
  lgvdw value = yes or no = whether or not to use a low gradient vdW correction
  safezone = factor used for array allocation
  mincap = minimum size for array allocation
  minhbonds = minimum size use for storing hydrogen bonds
  tabulate value = size of interpolation table for Lennard-Jones and Coulomb.
↳interactions
  list/blocking value = yes or no = whether or not to use "blocking" scheme for.
↳bond list build
```

## 19.217.2 Examples

```
pair_style reaxff NULL
pair_style reaxff controlfile checkqeq no
pair_style reaxff NULL lgvdw yes
pair_style reaxff NULL safezone 1.6 mincap 100
pair_coeff * *ffield.reax C H O N
```

## 19.217.3 Description

Style *reaxff* computes the ReaxFF potential of van Duin, Goddard and co-workers. ReaxFF uses distance-dependent bond-order functions to represent the contributions of chemical bonding to the potential energy. There is more than one version of ReaxFF. The version implemented in LAMMPS uses the functional forms documented in the supplemental information of the following paper: (*Chenoweth et al., 2008*). The version integrated into LAMMPS matches the version of ReaxFF From Summer 2010. For more technical details about the pair reaxff implementation of ReaxFF, see the (*Aktulga*) paper. The *reaxff* style was initially implemented as a stand-alone C code and is now converted to C++ and integrated into LAMMPS as a package.

The *reaxff/kk* style is a Kokkos version of the ReaxFF potential that is derived from the *reaxff* style. The Kokkos version can run on GPUs and can also use OpenMP multithreading. For more information about the Kokkos package, see *Packages details* and *Speed kokkos* doc pages. One important consideration when using the *reaxff/kk* style is the choice of either half or full neighbor lists. This setting can be changed using the Kokkos *package* command.

The *reaxff* style differs from the (obsolete) “pair\_style reax” command in the implementation details. The *reax* style was a Fortran library, linked to LAMMPS. The *reax* style has been removed from LAMMPS after the 12 December 2018 version.

LAMMPS provides several different versions of *ffield.reax* in its potentials dir, each called potentials/*ffield.reax.label*. These are documented in potentials/README.reax. The default *ffield.reax* contains parameterizations for the following elements: C, H, O, N.

The format of these files is identical to that used originally by van Duin. We have tested the accuracy of *pair\_style reaxff* potential against the original ReaxFF code for the systems mentioned above. You can use other ffield files for specific chemical systems that may be available elsewhere (but note that their accuracy may not have been tested).

---

**Note:** We do not distribute a wide variety of ReaxFF force field files with LAMMPS. Adri van Duin's group at PSU is the central repository for this kind of data as they are continuously deriving and updating parameterizations for different classes of materials. You can submit a contact request at the Materials Computation Center (MCC) website <https://www.mri.psu.edu/materials-computation-center/connect-mcc>, describing the material(s) you are interested in modeling with ReaxFF. They can tell you what is currently available or what it would take to create a suitable ReaxFF parameterization.

---

The *cfile* setting can be specified as NULL, in which case default settings are used. A control file can be specified which defines values of control variables. Some control variables are global parameters for the ReaxFF potential. Others define certain performance and output settings. Each line in the control file specifies the value for a control variable. The format of the control file is described below.

---

**Note:** The LAMMPS default values for the ReaxFF global parameters correspond to those used by Adri van Duin's stand-alone serial code. If these are changed by setting control variables in the control file, the results from LAMMPS and the serial code will not agree.

---

Examples using *pair\_style reaxff* are provided in the examples/reax sub-directory.

Use of this pair style requires that a charge be defined for every atom. See the *atom\_style* and *read\_data* commands for details on how to specify charges.

The ReaxFF parameter files provided were created using a charge equilibration (QEq) model for handling the electrostatic interactions. Therefore, by default, LAMMPS requires that either the *fix qeq/reaxff* or the *fix qeq/shielded* or *fix acks2/reaxff* command be used with *pair\_style reaxff* when simulating a ReaxFF model, to equilibrate the charges each timestep.

Using the keyword *checkqeq* with the value *no* turns off the check for the QEq fixes, allowing a simulation to be run without charge equilibration. In this case, the static charges you assign to each atom will be used for computing the electrostatic interactions in the system. See the *fix qeq/reaxff* or *fix qeq/shielded* or *fix acks2/reaxff* command documentation for more details.

Using the optional keyword *lgvdw* with the value *yes* turns on the low-gradient correction of ReaxFF for long-range London Dispersion, as described in the (Liu) paper. The bundled force field file *ffield.reax.lg* is designed for this correction, and is trained for several energetic materials (see "Liu"). When using *lgvdw yes*, the recommended value for parameter *thb* is 0.01, which can be set in the control file. Note: Force field files are different for the original or lg corrected pair styles, using the wrong ffield file generates an error.

Using the optional keyword *enobonds* with the value *yes*, the energy of atoms with no bonds (i.e. isolated atoms) is included in the total potential energy and the per-atom energy of that atom. If the value *no* is specified then the energy of atoms with no bonds is set to zero. The latter behavior is usual not desired, as it causes discontinuities in the potential energy when the bonding of an atom drops to zero.

Optional keywords *safezone*, *mincap*, and *minhbonds* are used for allocating reaxff arrays. Increasing these values can avoid memory problems, such as segmentation faults and bondchk failed errors, that could occur under certain conditions. These keywords are not used by the Kokkos version, which instead uses a more robust memory allocation scheme that checks if the sizes of the arrays have been exceeded and automatically allocates more memory.

The keyword *tabulate* controls the size of interpolation table for Lennard-Jones and Coulomb interactions. Tabulation may also be set in the control file (see below). If tabulation is set in both the input script and the control file, the value in the control file will be ignored. A size of 10000 is typically used for the interpolation table. A value of 0 means no tabulation will be used.

The keyword *list/blocking* is only supported by the Kokkos version of ReaxFF and ignored otherwise. Setting the value to *yes* enables the “blocking” scheme (dynamically building interaction lists) for the ReaxFF bond neighbor list. This reduces the number of empty interactions and can improve performance in some cases (e.g. large number of atoms/GPU on AMD hardware). It is also enabled by default when running the CPU with Kokkos.

The thermo variable *evdwl* stores the sum of all the ReaxFF potential energy contributions, with the exception of the Coulombic and charge equilibration contributions which are stored in the thermo variable *ecoul*. The output of these quantities is controlled by the *thermo* command.

This pair style tallies a breakdown of the total ReaxFF potential energy into sub-categories, which can be accessed via the *compute pair* command as a vector of values of length 14. The 14 values correspond to the following sub-categories (the variable names in italics match those used in the original FORTRAN ReaxFF code):

1. *eb* = bond energy
2. *ea* = atom energy
3. *elp* = lone-pair energy
4. *emol* = molecule energy (always 0.0)
5. *ev* = valence angle energy
6. *epen* = double-bond valence angle penalty
7. *ecoa* = valence angle conjugation energy
8. *ehb* = hydrogen bond energy
9. *et* = torsion energy
10. *eco* = conjugation energy
11. *ew* = van der Waals energy
12. *ep* = Coulomb energy
13. *efi* = electric field energy (always 0.0)
14. *eqeq* = charge equilibration energy

To print these quantities to the log file (with descriptive column headings) the following commands could be included in an input script:

```
compute reax all pair reaxff
variable eb      equal c_reax[1]
variable ea      equal c_reax[2]
[...]
variable eqeq    equal c_reax[14]
thermo_style custom step temp epair v_eb v_ea [...] v_eqeq
```

Only a single *pair\_coeff* command is used with the *reaxff* style which specifies a ReaxFF potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the *pair\_coeff* command, where N is the number of LAMMPS atom types:

- filename
- N indices = ReaxFF elements

The filename is the ReaxFF potential file.

In the ReaxFF potential file, near the top, after the general parameters, is the atomic parameters section that contains element names, each with a couple dozen numeric parameters. If there are M elements specified in the *ffield* file, think of these as numbered 1 to M. Each of the N indices you specify for the N atom types of LAMMPS atoms must be an integer from 1 to M. Atoms with LAMMPS type 1 will be mapped to whatever element you specify as the first index

value, etc. If a mapping value is specified as NULL, the mapping is not performed. This can be used when the *reaxff* style is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

As an example, say your LAMMPS simulation has 4 atom types and the elements are ordered as C, H, O, N in the *ffield* file. If you want the LAMMPS atom type 1 and 2 to be C, type 3 to be N, and type 4 to be H, you would use the following *pair\_coeff* command:

```
pair_coeff * * ffield.reax C C N H
```

## 19.217.4 Control file

The format of a line in the control file is as follows:

```
variable_name value
```

and it may be followed by an “!” character and a trailing comment.

If the value of a control variable is not specified, then default values are used. What follows is the list of variables along with a brief description of their use and default values.

### *simulation\_name*

Output files produced by *pair\_style reaxff* carry this name + extensions specific to their contents. Partial energies are reported with a “.pot” extension, while the trajectory file has “.trj” extension.

### *tabulate\_long\_range*

To improve performance, long range interactions can optionally be tabulated (0 means no tabulation). Value of this variable denotes the size of the long range interaction table. The range from 0 to long range cutoff (defined in the *ffield* file) is divided into *tabulate\_long\_range* points. Then at the start of simulation, we fill in the entries of the long range interaction table by computing the energies and forces resulting from van der Waals and Coulomb interactions between every possible atom type pairs present in the input system. During the simulation we consult to the long range interaction table to estimate the energy and forces between a pair of atoms. Linear interpolation is used for estimation. (default value = 0)

### *energy\_update\_freq*

Denotes the frequency (in number of steps) of writes into the partial energies file. (default value = 0)

### *nbrhood\_cutoff*

Denotes the near neighbors cutoff (in Angstroms) regarding the bonded interactions. (default value = 5.0)

### *hbond\_cutoff*

Denotes the cutoff distance (in Angstroms) for hydrogen bond interactions.(default value = 7.5. A value of 0.0 turns off hydrogen bonds)

### *bond\_graph\_cutoff*

is the threshold used in determining what is a physical bond, what is not. Bonds and angles reported in the trajectory file rely on this cutoff. (default value = 0.3)

### *thb\_cutoff*

cutoff value for the strength of bonds to be considered in three body interactions. (default value = 0.001)

### *thb\_cutoff\_sq*

cutoff value for the strength of bond order products to be considered in three body interactions. (default value = 0.00001)

### *write\_freq*

Frequency of writes into the trajectory file. (default value = 0)

***traj\_title***

Title of the trajectory - not the name of the trajectory file.

***atom\_info***

1 means print only atomic positions + charge (default = 0)

***atom\_forces***

1 adds net forces to atom lines in the trajectory file (default = 0)

***atom\_velocities***

1 adds atomic velocities to atoms line (default = 0)

***bond\_info***

1 prints bonds in the trajectory file (default = 0)

***angle\_info***

1 prints angles in the trajectory file (default = 0)

---

### 19.217.5 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* mix, shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.217.6 Restrictions

This pair style is part of the REAXFF package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

The ReaxFF potential files provided with LAMMPS in the potentials directory are parameterized for *real units*. You can use the ReaxFF pair style with any LAMMPS units, but you would need to create your own potential file with coefficients listed in the appropriate units if your simulation does not use “real” units.



## 19.217.7 Related commands

*pair\_coeff*, *fix qeq/reaxff*, *fix acks2/reaxff*, *fix reaxff/bonds*, *fix reaxff/species*

## 19.217.8 Default

The keyword defaults are `checkqeq = yes`, `enobonds = yes`, `lgvdw = no`, `safezone = 1.2`, `mincap = 50`, `minhbonds = 25`, `tabulate = 0`, `list/blocking = yes` on CPU, no on GPU.

(**Chenoweth\_2008**) Chenoweth, van Duin and Goddard, Journal of Physical Chemistry A, 112, 1040-1053 (2008).

(Aktulga) Aktulga, Fogarty, Pandit, Grama, Parallel Computing, 38, 245-259 (2012).

(**Liu**) L. Liu, Y. Liu, S. V. Zybin, H. Sun and W. A. Goddard, Journal of Physical Chemistry A, 115, 11016-11022 (2011).

## 19.218 pair\_style resquared command

Accelerator Variants: *resquared/gpu*, *resquared/omp*

### 19.218.1 Syntax

```
pair_style resquared cutoff
```

- `cutoff` = global cutoff for interactions (distance units)

### 19.218.2 Examples

```
pair_style resquared 10.0
pair_coeff * * 1.0 1.0 1.7 3.4 3.4 1.0 1.0 1.0
```

### 19.218.3 Description

Style *resquared* computes the RE-squared anisotropic interaction (*Everaers*), (*Babadi*) between pairs of ellipsoidal and/or spherical Lennard-Jones particles. For ellipsoidal interactions, the potential considers the ellipsoid as being comprised of small spheres of size  $\sigma$ . LJ particles are a single sphere of size  $\sigma$ . The distinction is made to allow the pair style to make efficient calculations of ellipsoid/solvent interactions.

Details for the equations used are given in the references below and in [this supplementary document](#).

Use of this pair style requires the NVE, NVT, or NPT fixes with the *asphere* extension (e.g. *fix nve/asphere*) in order to integrate particle rotation. Additionally, *atom\_style ellipsoid* should be used since it defines the rotational state and the size and shape of each ellipsoidal particle.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $A_{12}$  = Energy Prefactor/Hamaker constant (energy units)
- $\sigma$  = atomic interaction diameter (distance units)

- $\epsilon_{i,a}$  = relative well depth of type I for side-to-side interactions
- $\epsilon_{i,b}$  = relative well depth of type I for face-to-face interactions
- $\epsilon_{i,c}$  = relative well depth of type I for end-to-end interactions
- $\epsilon_{j,a}$  = relative well depth of type J for side-to-side interactions
- $\epsilon_{j,b}$  = relative well depth of type J for face-to-face interactions
- $\epsilon_{j,c}$  = relative well depth of type J for end-to-end interactions
- cutoff (distance units)

The parameters used depend on the type of the interacting particles, i.e. ellipsoids or LJ spheres. The type of a particle is determined by the diameters specified for its 3 shape parameters. If all 3 shape parameters = 0.0, then the particle is treated as an LJ sphere. The  $\epsilon_{i,*}$  or  $\epsilon_{j,*}$  parameters are ignored for LJ spheres. If the 3 shape parameters are > 0.0, then the particle is treated as an ellipsoid (even if the 3 parameters are equal to each other).

A12 specifies the energy prefactor which depends on the types of the two interacting particles.

For ellipsoid/ellipsoid interactions, the interaction is computed by the formulas in the supplementary document referenced above. A12 is the Hamaker constant as described in (*Everaers*). In LJ units:

$$A_{12} = 4\pi^2 \epsilon_{LJ} (\rho\sigma^3)^2$$

where  $\rho$  gives the number density of the spherical particles composing the ellipsoids and  $\epsilon_{LJ}$  determines the interaction strength of the spherical particles.

For ellipsoid/LJ sphere interactions, the interaction is also computed by the formulas in the supplementary document referenced above. A12 has a modified form (see [here](#) for details):

$$A_{12} = 4\pi^2 \epsilon_{LJ} (\rho\sigma^3)$$

For ellipsoid/LJ sphere interactions, a correction to the distance- of-closest approach equation has been implemented to reduce the error from two particles of disparate sizes; see [this supplementary document](#).

For LJ sphere/LJ sphere interactions, the interaction is computed using the standard Lennard-Jones formula, which is much cheaper to compute than the ellipsoidal formulas. A12 is used as epsilon in the standard LJ formula:

$$A_{12} = \epsilon_{LJ}$$

and the specified  $\sigma$  is used as the  $\sigma$  in the standard LJ formula.

When one of both of the interacting particles are ellipsoids, then  $\sigma$  specifies the diameter of the continuous distribution of constituent particles within each ellipsoid used to model the RE-squared potential. Note that this is a different meaning for  $\sigma$  than the *pair\_style gayberne* potential uses.

The  $\epsilon_i$  and  $\epsilon_j$  coefficients are defined for atom types, not for pairs of atom types. Thus, in a series of pair\_coeff commands, they only need to be specified once for each atom type.

Specifically, if any of  $\epsilon_{i,a}$ ,  $\epsilon_{i,b}$ ,  $\epsilon_{i,c}$  are non-zero, the three values are assigned to atom type I. If all the  $\epsilon_i$  values are zero, they are ignored. If any of  $\epsilon_{j,a}$ ,  $\epsilon_{j,b}$ ,  $\epsilon_{j,c}$  are non-zero, the three values are assigned to atom type J. If all three  $\epsilon_i$  values are zero, they are ignored. Thus the typical way to define the  $\epsilon_i$  and  $\epsilon_j$  coefficients is to list their values in “pair\_coeff I J” commands when I = J, but set them to 0.0 when I != J. If you do list them when I != J, you should insure they are consistent with their values in other pair\_coeff commands.

Note that if this potential is being used as a sub-style of *pair\_style hybrid*, and there is no “pair\_coeff I I” setting made for RE-squared for a particular type I (because I-I interactions are computed by another hybrid pair potential), then you still need to insure the epsilon a,b,c coefficients are assigned to that type in a “pair\_coeff I J” command.

For large uniform molecules it has been shown that the  $\epsilon_{*,*}$  energy parameters are approximately representable in terms of local contact curvatures (*Everaers*):

$$\epsilon_a = \sigma \cdot \frac{a}{b \cdot c}; \epsilon_b = \sigma \cdot \frac{b}{a \cdot c}; \epsilon_c = \sigma \cdot \frac{c}{a \cdot b}$$

where a, b, and c give the particle diameters.

The last coefficient is optional. If not specified, the global cutoff specified in the `pair_style` command is used.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 19.218.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance can be mixed, but only for sphere pairs. The default mix value is *geometric*. See the “pair\_modify” command for details. Other type pairs cannot be mixed, due to the different meanings of the energy prefactors used to calculate the interactions and the implicit dependence of the ellipsoid-sphere interaction on the equation for the Hamaker constant presented here. Mixing of sigma and epsilon followed by calculation of the energy prefactors using the equations above is recommended.

This pair style supports the *pair\_modify* shift option for the energy of the Lennard-Jones portion of the pair interaction, but only for sphere-sphere interactions. There is no shifting performed for ellipsoidal interactions due to the anisotropic dependence of the interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords of the *run\_style command*.

#### 19.218.5 Restrictions

This style is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires that atoms be ellipsoids as defined by the *atom\_style ellipsoid* command.

Particles acted on by the potential can be finite-size aspherical or spherical particles, or point particles. Spherical particles have all 3 of their shape parameters equal to each other. Point particles have all 3 of their shape parameters equal to 0.0.

The distance-of-closest-approach approximation used by LAMMPS becomes less accurate when high-aspect ratio ellipsoids are used.

### 19.218.6 Related commands

*pair\_coeff*, *fix nve/asphere*, *compute temp/asphere*, *pair\_style gayberne*

### 19.218.7 Default

none

---

**(Everaers)** Everaers and Ejtehadi, Phys Rev E, 67, 041710 (2003).

**(Babadi)** Babadi, Ejtehadi, Everaers, J Comp Phys, 219, 770-779 (2006).

## 19.219 pair\_style saip/metal command

Accelerator Variant: *saip/metal/opt*

### 19.219.1 Syntax

```
pair_style [hybrid/overlay ...] saip/metal cutoff tap_flag
```

- cutoff = global cutoff (distance units)
- tap\_flag = 0/1 to turn off/on the taper function

### 19.219.2 Examples

```
pair_style hybrid/overlay saip/metal 16.0 1
pair_coeff * * saip/metal CHAu.ILP Au C H

pair_style hybrid/overlay eam rebo saip/metal 16.0
pair_coeff 1 1 eam Au_u3.eam Au NULL NULL
pair_coeff * * rebo CH.rebo NULL C H
pair_coeff * * saip/metal CHAu.ILP Au C H
```

### 19.219.3 Description

The *saip/metal* style computes the registry-dependent interlayer potential (ILP) potential for hetero-junctions formed with hexagonal 2D materials and metal surfaces, as described in (Ouyang6).

$$E = \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij}$$

$$V_{ij} = \text{Tap}(r_{ij}) \left\{ e^{-\alpha(r_{ij}/\beta-1)} [\epsilon + f(\rho_{ij}) + f(\rho_{ji})] - \frac{1}{1 + e^{-d[(r_{ij}/(s_R \cdot r^{eff})) - 1]}} \cdot \frac{C_6}{r_{ij}^6} \right\}$$

$$\rho_{ij}^2 = r_{ij}^2 - (\mathbf{r}_{ij} \cdot \mathbf{n}_i)^2$$

$$\rho_{ji}^2 = r_{ij}^2 - (\mathbf{r}_{ij} \cdot \mathbf{n}_j)^2$$

$$f(\rho) = C e^{-(\rho/\delta)^2}$$

$$\text{Tap}(r_{ij}) = 20 \left( \frac{r_{ij}}{R_{cut}} \right)^7 - 70 \left( \frac{r_{ij}}{R_{cut}} \right)^6 + 84 \left( \frac{r_{ij}}{R_{cut}} \right)^5 - 35 \left( \frac{r_{ij}}{R_{cut}} \right)^4 + 1$$

Where  $\text{Tap}(r_{ij})$  is the taper function which provides a continuous cutoff (up to third derivative) for interatomic separations larger than  $r_c$  *pair\_style ilp\_graphene\_hbn*.

It is important to include all the pairs to build the neighbor list for calculating the normals.

---

**Note:** To account for the isotropic nature of the isolated gold atom electron cloud, their corresponding normal vectors (*bf n\_i*) are assumed to lie along the interatomic vector *bf r\_ij*. Notably, this assumption is suitable for many bulk material surfaces, for example, for systems possessing s-type valence orbitals or metallic surfaces, whose valence electrons are mostly delocalized, such that their Pauli repulsion with the electrons of adjacent surfaces are isotropic. Caution should be used in the case of very small gold contacts, for example, nano-clusters, where edge effects may become relevant.

---

The parameter file (e.g. CHAu.ILP), is intended for use with *metal units*, with energies in meV. Two additional parameters, *S*, and *rcut* are included in the parameter file. *S* is designed to facilitate scaling of energies. *rcut* is designed to build the neighbor list for calculating the normals for each atom pair.

---

**Note:** The parameters presented in the parameter file (e.g. BNCH.ILP), are fitted with taper function by setting the cutoff equal to 16.0 Angstrom. Using different cutoff or taper function should be careful.

---

This potential must be used in combination with hybrid/overlay. Other interactions can be set to zero using *pair\_style none*.

This pair style tallies a breakdown of the total interlayer potential energy into sub-categories, which can be accessed via the *compute pair* command as a vector of values of length 2. The 2 values correspond to the following sub-categories:

1.  $E_{vdW}$  = vdW (attractive) energy
2.  $E_{Rep}$  = Repulsive energy

To print these quantities to the log file (with descriptive column headings) the following commands could be included in an input script:

```
compute 0 all pair saip/metal
variable Evdw equal c_0[1]
variable Erep equal c_0[2]
thermo_style custom step temp epair v_Erep v_Evdw
```

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.219.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the `pair_modify` `mix`, `shift`, `table`, and `tail` options.

This pair style does not write their information to binary restart files, since it is stored in potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

#### 19.219.5 Restrictions

This pair style is part of the INTERLAYER package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This pair style requires the `newton` setting to be *on* for pair interactions.

The CHAu.ILP potential file provided with LAMMPS (see the potentials directory) are parameterized for *metal* units. You can use this potential with any LAMMPS units, but you would need to create your BNCH.ILP potential file with coefficients listed in the appropriate units, if your simulation does not use *metal* units.

#### 19.219.6 Related commands

*pair\_coeff*, *pair\_none*, *pair\_style hybrid/overlay*, *pair\_style drip*, *pair\_style ilp\_tmd*, *pair\_style ilp\_graphene\_hbn*, *pair\_style pair\_kolmogorov\_crespi\_z*, *pair\_style pair\_kolmogorov\_crespi\_full*, *pair\_style pair\_lebedeva\_z*, *pair\_style pair\_coul\_shield*.

#### 19.219.7 Default

`tap_flag = 1`

---

(Ouyang<sup>6</sup>) W. Ouyang, O. Hod, and R. Guerra, J. Chem. Theory Comput. 17, 7215 (2021).

## 19.220 pair\_style sdpd/taitwater/isothermal command

### 19.220.1 Syntax

```
pair_style sdpd/taitwater/isothermal temperature viscosity seed
```

- temperature = temperature of the fluid (temperature units)
- viscosity = dynamic viscosity of the fluid (mass\*distance/time units)
- seed = random number generator seed (positive integer, optional)

### 19.220.2 Examples

```
pair_style sdpd/taitwater/isothermal 300. 1. 28681
pair_coeff * * 1000.0 1430.0 2.4
```

### 19.220.3 Description

The sdpd/taitwater/isothermal style computes forces between mesoscopic particles according to the Smoothed Dissipative Particle Dynamics model described in this paper by (*Espanol and Revenga*) under the following assumptions:

1. The temperature is constant and uniform.
2. The shear viscosity is constant and uniform.
3. The volume viscosity is negligible before the shear viscosity.
4. The Boltzmann constant is negligible before the heat capacity of a single mesoscopic particle of fluid.

The third assumption is true for water in nearly incompressible flows. The fourth holds true for water for any reasonable size one can imagine for a mesoscopic particle.

The pressure forces between particles will be computed according to Tait's equation of state:

$$p = B \left[ \left( \frac{\rho}{\rho_0} \right)^\gamma - 1 \right]$$

where  $\gamma = 7$  and  $B = c_0^2 \rho_0 / \gamma$ , with  $\rho_0$  being the reference density and  $c_0$  the reference speed of sound.

The laminar viscosity and the random forces will be computed according to formulas described in (*Espanol and Revenga*).

**Warning:** Similar to *brownian* and *dpd* styles, the *newton* setting for pairwise interactions needs to be on when running LAMMPS in parallel if you want to ensure linear momentum conservation. Otherwise random forces generated for pairs straddling processor boundary will not be equal and opposite.

**Note:** The actual random seed used will be a mix of what you specify and other parameters like the MPI ranks. This is to ensure that different MPI tasks have distinct seeds.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above.

- $\rho_0$  reference density (mass/volume units)
  - $c_0$  reference soundspeed (distance/time units)
  - h kernel function cutoff (distance units)
- 

#### 19.220.4 Mixing, shift, table, tail correction, restart, rRESPA info

This style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This style does not support the *pair\_modify* shift, table, and tail options.

This style does not write information to *binary restart files*. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

#### 19.220.5 Restrictions

This pair style is part of the DPD-SMOOTH package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 19.220.6 Related commands

*pair coeff*, *pair sph/rhosum*, *pair sph/taitwater*

#### 19.220.7 Default

The default seed is 0 (before mixing).

---

(Espanol and Revenga) Espanol, Revenga, Physical Review E, 67, 026705 (2003).

### 19.221 *pair\_style smatb* command

### 19.222 *pair\_style smatb/single* command

#### 19.222.1 Syntax

```
pair_style style args
```

- style = *smatb* or *smatb/single*
- args = none



## 19.222.2 Examples

```
pair_style smatb
pair_coeff 1 1 2.88 10.35 4.178 0.210 1.818 4.07293506 4.9883063257983666
```

```
pair_style smatb/single
pair_coeff 1 1 2.88 10.35 4.178 0.210 1.818 4.07293506 4.9883063257983666
```

## 19.222.3 Description

New in version 4May2022.

The *smatb* and *smatb/single* styles compute the Second Moment Approximation to the Tight Binding (*Cyrot*), (*Gupta*), (*Rosato*), given by

$$E_i = \sum_{j, R_{ij} \leq R_c} \alpha(R_{ij}) - \sqrt{\sum_{j, R_{ij} \leq R_c} \Xi^2(R_{ij})}$$

$R_{ij}$  is the distance between the atom  $i$  and  $j$ . And the two functions  $\alpha(r)$  and  $\Xi(r)$  are:

$$\alpha(r) = \begin{cases} A e^{-p\left(\frac{r}{R_0}-1\right)} & r < R_{sc} \\ a_3 (r - R_c)^3 + a_4 (r - R_c)^4 + a_5 (r - R_c)^5 & R_{sc} < r < R_c \end{cases}$$

$$\Xi(r) = \begin{cases} \xi e^{-q\left(\frac{r}{R_0}-1\right)} & r < R_{sc} \\ x_3 (r - R_c)^3 + x_4 (r - R_c)^4 + x_5 (r - R_c)^5 & R_{sc} < r < R_c \end{cases}$$

The polynomial coefficients  $a_3$ ,  $a_4$ ,  $a_5$ ,  $x_3$ ,  $x_4$ ,  $x_5$  are computed by LAMMPS: the two exponential terms and their first and second derivatives are smoothly reduced to zero, from the inner cutoff  $R_{sc}$  to the outer cutoff  $R_c$ .

The *smatb/single* style is an optimization when using only a single atom type.

## 19.222.4 Coefficients

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $R_0$  (distance units)
- $p$  (dimensionless)
- $q$  (dimensionless)
- $A$  (energy units)
- $\xi$  (energy units)
- $R_{cs}$  (distance units)
- $R_c$  (distance units)

Note that:  $R_0$  is the nearest neighbor distance, usually coincides with the diameter of the atoms

See the *run\_style* command for details.

### 19.222.5 Mixing info

For atom type pairs I,J and I != J the coefficients are not automatically mixed.

---

### 19.222.6 Restrictions

These pair styles are part of the SMTBQ package and are only enabled if LAMMPS is built with that package. See the *Build package* page for more info.

These pair styles require the *newton* setting to be “on” for pair interactions.

### 19.222.7 Related commands

- *pair\_coeff*

### 19.222.8 Default

none

---

**(Cyrot)** Cyrot-Lackmann and Ducastelle, Phys Rev. B, 4, 2406-2412 (1971).

**(Gupta)** Gupta, Phys Rev. B, 23, 6265-6270 (1981).

**(Rosato)** Rosato and Guillope and Legrand, Philosophical Magazine A, 59.2, 321-336 (1989).

## 19.223 pair\_style smd/hertz command

### 19.223.1 Syntax

```
pair_style smd/hertz scale_factor
```

---

### 19.223.2 Examples

```
pair_style smd/hertz 1.0  
pair_coeff 1 1 <contact_stiffness>
```

---

### 19.223.3 Description

The *smd/hertz* style calculates contact forces between SPH particles belonging to different physical bodies.

The contact forces are calculated using a Hertz potential, which evaluates the overlap between two particles (whose spatial extents are defined via its contact radius). The effect is that a particles cannot penetrate into each other. The parameter <contact\_stiffness> has units of pressure and should equal roughly one half of the Young’s modulus (or bulk modulus in the case of fluids) of the material model associated with the SPH particles.

The parameter *scale\_factor* can be used to scale the particles' contact radii. This can be useful to control how close particles can approach each other. Usually, *scale\_factor* =1.0.

#### 19.223.4 Mixing, shift, table, tail correction, restart, rRESPA info

No mixing is performed automatically. Currently, no part of MACHDYN supports restarting nor minimization. rRESPA does not apply to this pair style.

#### 19.223.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 19.223.6 Related commands

*pair\_coeff*

#### 19.223.7 Default

none

### 19.224 pair\_style smd/tlsph command

#### 19.224.1 Syntax

```
pair_style smd/tlsph args
```

#### 19.224.2 Examples

```
pair_style smd/tlsph
```

#### 19.224.3 Description

The *smd/tlsph* style computes particle interactions according to continuum mechanics constitutive laws and a Total-Lagrangian Smooth-Particle Hydrodynamics algorithm.

This pair style is invoked with the following command:

```
pair_style smd/tlsph
pair_coeff i j *COMMON rho0 E nu Q1 Q2 hg Cp &
          *END
```

Here,  $i$  and  $j$  denote the *LAMMPS* particle types for which this pair style is defined. Note that  $i$  and  $j$  must be equal, i.e., no *tlsph* cross interactions between different particle types are allowed. In contrast to the usual *LAMMPS pair coeff* definitions, which are given solely a number of floats and integers, the *tlsph pair coeff* definition is organized using keywords. These keywords mark the beginning of different sets of parameters for particle properties, material constitutive models, and damage models. The *pair coeff* line must be terminated with the *\*END* keyword. The use of the line continuation operator *&* is recommended. A typical invocation of the *tlsph* for a solid body would consist of an equation of state for computing the pressure (the diagonal components of the stress tensor), and a material model to compute shear stresses (the off-diagonal components of the stress tensor). Damage and failure models can also be added.

Please see the [SMD user guide](#) for a complete listing of the possible keywords and material models.

---

#### 19.224.4 Mixing, shift, table, tail correction, restart, rRESPA info

No mixing is performed automatically. Currently, no part of MACHDYN supports restarting nor minimization. rRESPA does not apply to this pair style.

---

#### 19.224.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

#### 19.224.6 Related commands

*pair\_coeff*

#### 19.224.7 Default

none

### 19.225 pair\_style smd/tri\_surface command

#### 19.225.1 Syntax

```
pair_style smd/tri_surface scale_factor
```

## 19.225.2 Examples

```
pair_style smd/tri_surface 1.0
pair_coeff 1 1 <contact_stiffness>
```

## 19.225.3 Description

The *smd/tri\_surface* style calculates contact forces between SPH particles and a rigid wall boundary defined via the *smd/wall\_surface* fix.

The contact forces are calculated using a Hertz potential, which evaluates the overlap between a particle (whose spatial extents are defined via its contact radius) and the triangle. The effect is that a particle cannot penetrate into the triangular surface. The parameter <contact\_stiffness> has units of pressure and should equal roughly one half of the Young's modulus (or bulk modulus in the case of fluids) of the material model associated with the SPH particle

The parameter *scale\_factor* can be used to scale the particles' contact radii. This can be useful to control how close particles can approach the triangulated surface. Usually, *scale\_factor* =1.0.

## 19.225.4 Mixing, shift, table, tail correction, restart, rRESPA info

No mixing is performed automatically. Currently, no part of MACHDYN supports restarting nor minimization. rRESPA does not apply to this pair style.

## 19.225.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 19.225.6 Related commands

*pair\_coeff*

## 19.225.7 Default

none

## 19.226 pair\_style smd/ulsph command

### 19.226.1 Syntax

```
pair_style smd/ulsph args
```

- these keywords must be given

keyword = \*DENSITY\_SUMMATION or \*DENSITY\_CONTINUITY and \*VELOCITY\_GRADIENT or \*NO\_↪VELOCITY\_GRADIENT and \*GRADIENT\_CORRECTION or \*NO\_GRADIENT\_CORRECTION

## 19.226.2 Examples

```
pair_style smd/ulsph *DENSITY_CONTINUITY *VELOCITY_GRADIENT *NO_GRADIENT_CORRECTION
```

## 19.226.3 Description

The *smd/ulsph* style computes particle interactions according to continuum mechanics constitutive laws and an updated Lagrangian Smooth-Particle Hydrodynamics algorithm.

This pair style is invoked similar to the following command:

```
pair_style smd/ulsph *DENSITY_CONTINUITY *VELOCITY_GRADIENT *NO_GRADIENT_CORRECTION
pair_coeff i j *COMMON rho0 c0 Q1 Cp hg &
*END
```

Here, *i* and *j* denote the *LAMMPS* particle types for which this pair style is defined. Note that *i* and *j* can be different, i.e., *ulsph* cross interactions between different particle types are allowed. However, *i-i* respectively *j-j* *pair\_coeff* lines have to precede a cross interaction. In contrast to the usual *LAMMPS pair coeff* definitions, which are given solely a number of floats and integers, the *ulsph pair coeff* definition is organized using keywords. These keywords mark the beginning of different sets of parameters for particle properties, material constitutive models, and damage models. The *pair coeff* line must be terminated with the *\*END* keyword. The use the line continuation operator *&* is recommended. A typical invocation of the *ulsph* for a solid body would consist of an equation of state for computing the pressure (the diagonal components of the stress tensor), and a material model to compute shear stresses (the off-diagonal components of the stress tensor).

Note that the use of *\*GRADIENT\_CORRECTION* can lead to severe numerical instabilities. For a general fluid simulation, *\*NO\_GRADIENT\_CORRECTION* is recommended.

Please see the [SMD user guide](#) for a complete listing of the possible keywords and material models.

---

## 19.226.4 Mixing, shift, table, tail correction, restart, rRESPA info

No mixing is performed automatically. Currently, no part of MACHDYN supports restarting nor minimization. rRESPA does not apply to this pair style.

---

## 19.226.5 Restrictions

This fix is part of the MACHDYN package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

## 19.226.6 Related commands

*pair\_coeff*

## 19.226.7 Default

none

## 19.227 pair\_style smtbq command

### 19.227.1 Syntax

```
pair_style smtbq
```

### 19.227.2 Examples

```
pair_style smtbq
pair_coeff * *ffield.smtbq.Al2O3 O Al
```

### 19.227.3 Description

This pair style computes a variable charge SMTB-Q (Second-Moment tight-Binding QEq) potential as described in *SMTB-Q\_1* and *SMTB-Q\_2*. Briefly, the energy of metallic-oxygen systems is given by three contributions:

$$\begin{aligned}
 E_{tot} &= E_{ES} + E_{OO} + E_{MO} \\
 E_{ES} &= \sum_i \left[ \chi_i^0 Q_i + \frac{1}{2} J_i^0 Q_i^2 + \frac{1}{2} \sum_{j \neq i} J_{ij}(r_{ij}) f_{cut}^{R_{cut}}(r_{ij}) Q_i Q_j \right] \\
 E_{OO} &= \sum_{i,j}^{i,j=O} \left[ C \exp\left(-\frac{r_{ij}}{\rho}\right) - D f_{cut}^{r_1^{OO} r_2^{OO}}(r_{ij}) \exp(B r_{ij}) \right] \\
 E_{MO} &= \sum_i E_{cov}^i + \sum_{j \neq i} A f_{cut}^{r_{c1} r_{c2}}(r_{ij}) \exp\left[-p\left(\frac{r_{ij}}{r_0} - 1\right)\right]
 \end{aligned}$$

where  $E_{tot}$  is the total potential energy of the system,  $E_{ES}$  is the electrostatic part of the total energy,  $E_{OO}$  is the interaction between oxygen atoms and  $E_{MO}$  is a short-range interaction between metal and oxygen atoms. These interactions depend on interatomic distance  $r_{ij}$  and/or the charge  $Q_i$  of atoms  $i$ . Cut-off function enables smooth convergence to zero interaction.

The parameters appearing in the upper expressions are set in the `ffield.SMTBQ.Syst` file where `Syst` corresponds to the selected system (e.g. `ffield.SMTBQ.Al2O3`). Examples for  $\text{TiO}_2$ ,  $\text{Al}_2\text{O}_3$  are provided. A single `pair_coeff` command is used with the SMTBQ styles which provides the path to the potential file with parameters for needed elements. These are mapped to LAMMPS atom types by specifying additional arguments after the potential filename in the `pair_coeff` command. Note that atom type 1 must always correspond to oxygen atoms. As an example, to simulate a  $\text{TiO}_2$  system, atom type 1 has to be oxygen and atom type 2 Ti. The following `pair_coeff` command should then be used:

```
pair_coeff * * PathToLammps/potentials/ffield.smtbq.TiO2 O Ti
```

The electrostatic part of the energy consists of two components

self-energy of atom  $i$  in the form of a second order charge dependent polynomial and a long-range Coulombic electrostatic interaction. The latter uses the wolf summation method described in *Wolf*, spherically truncated at a longer cutoff,  $R_{coul}$ . The charge of each ion is modeled by an orbital Slater which depends on the principal quantum number ( $n$ ) of the outer orbital shared by the ion.

Interaction between oxygen,  $E_{OO}$ , consists of two parts, an attractive and a repulsive part. The attractive part is effective only at short range ( $< r_2^{OO}$ ). The attractive contribution was optimized to study surfaces reconstruction (e.g. *SMTB-Q\_2* in TiO<sub>2</sub>) and is not necessary for oxide bulk modeling. The repulsive part is the Pauli interaction between the electron clouds of oxygen. The Pauli repulsion and the coulombic electrostatic interaction have same cut off value. In the ffield.SMTBQ.Syst, the keyword 'buck' allows to consider only the repulsive O-O interactions. The keyword 'buckPlusAttr' allows to consider the repulsive and the attractive O-O interactions.

The short-range interaction between metal-oxygen,  $E_{MO}$  is based on the second moment approximation of the density of states with a N-body potential for the band energy term,  $E_{cov}^i$ , and a Born-Mayer type repulsive terms as indicated by the keyword 'second\_moment' in the ffield.SMTBQ.Syst. The energy band term is given by:

$$E_{cov}^{i(i=M,O)} = - \left\{ \eta_i (\mu \xi^0)^2 \int_{cut}^{r_{c1} r_{c2}} (r_{ij}) \left( \sum_{j(j=O,M)} \exp[-2q(\frac{r_{ij}}{r_0} - 1)] \right) \delta Q_i (2 \frac{n_0}{\eta_i} - \delta Q_i) \right\}^{1/2}$$

$$\delta Q_i = |Q_i^F| - |Q_i|$$

where  $\eta_i$  is the stoichiometry of atom  $i$ ,  $\delta Q_i$  is the charge delocalization of atom  $i$ , compared to its formal charge  $Q_i^F$ .  $n_0$ , the number of hybridized orbitals, is calculated with to the atomic orbitals shared  $d_i$  and the stoichiometry  $\eta_i$ .  $r_{c1}$  and  $r_{c2}$  are the two cutoff radius around the fourth neighbors in the cutoff function.

In the formalism used here,  $\xi^0$  is the energy parameter.  $\xi^0$  is in tight-binding approximation the hopping integral between the hybridized orbitals of the cation and the anion. In the literature we find many ways to write the hopping integral depending on whether one takes the point of view of the anion or cation. These are equivalent vision. The correspondence between the two visions is explained in appendix A of the article in the SrTiO<sub>3</sub> *SMTB-Q\_3* (parameter  $\beta$  shown in this article is in fact the  $\beta_O$ ). To summarize the relationship between the hopping integral  $\xi^O$  and the others, we have in an oxide C<sub>n</sub>O<sub>m</sub> the following relationship:

$$\xi^0 = \frac{\xi_O}{m} = \frac{\xi_C}{n}$$

$$\frac{\beta_O}{\sqrt{m}} = \frac{\beta_C}{\sqrt{n}} = \xi^0 \frac{\sqrt{m} + \sqrt{n}}{2}$$

Thus parameter  $\mu$ , indicated above, is given by  $\mu = \frac{1}{2}(\sqrt{n} + \sqrt{m})$

The potential offers the possibility to consider the polarizability of the electron clouds of oxygen by changing the slater radius of the charge density around the oxygen atoms through the parameters  $rBB$ ,  $rB$  and  $rS$  in the ffield.SMTBQ.Syst. This change in radius is performed according to the method developed by E. Maras *SMTB-Q\_2*. This method needs to determine the number of nearest neighbors around the oxygen. This calculation is based on first ( $r_{1n}$ ) and second ( $r_{2n}$ ) distances neighbors.

The SMTB-Q potential is a variable charge potential. The equilibrium charge on each atom is calculated by the electronegativity equalization (QEq) method. See *Rick* for further detail. One can adjust the frequency, the maximum number of iterative loop and the convergence of the equilibrium charge calculation. To obtain the energy conservation in NVE thermodynamic ensemble, we recommend to use a convergence parameter in the interval 10e-5 - 10e-6 eV.

The ffield.SMTBQ.Syst files are provided for few systems. They consist of nine parts and the lines beginning with '#' are comments (note that the number of comment lines matter). The first sections are on the potential parameters and others are on the simulation options and might be modified. Keywords are character type and must be enclosed in quotation marks ('').

- 1) Number of different element in the oxide:



- N\_elem= 2 or 3
- Divider line

## 2) Atomic parameters

For the anion (oxygen)

- Name of element (char) and stoichiometry in oxide
- Formal charge and mass of element
- Principal quantum number of outer orbital (n), electronegativity ( $\chi_i^0$ ) and hardness ( $J_i^0$ )
- Ionic radius parameters : max coordination number ( $coordBB = 6$  by default), bulk coordination number ( $coordB$ ), surface coordination number ( $coordS$ ) and  $rBB$ ,  $rB$  and  $rS$  the slater radius for each coordination number. **(note : If you don't want to change the slater radius, use three identical radius values)**
- Number of orbital shared by the element in the oxide ( $d_i$ )
- Divider line

For each cations (metal):

- Name of element (char) and stoichiometry in oxide
- Formal charge and mass of element
- Number of electron in outer orbital ( $ne$ ), electronegativity ( $\chi_i^0$ ), hardness ( $J_i^0$ ) and  $r_{Slater}$  the slater radius for the cation.
- Number of orbitals shared by the elements in the oxide ( $d_i$ )
- Divider line

## 3) Potential parameters:

- Keyword for element1, element2 and interaction potential ('second\_moment' or 'buck' or 'buckPlusAttr') between element 1 and 2. If the potential is 'second\_moment', specify 'oxide' or 'metal' for metal-oxygen or metal-metal interactions respectively.
- Potential parameter:
  - If type of potential is 'second\_moment' : A (eV),  $p$ ,  $\zeta^0$  (eV) and  $q$ ,  $r_{c1}$ (Å),  $r_{c2}$ (Å) and  $r_0$ (Å)
  - If type of potential is 'buck' : C (eV) and  $\rho$ (Å)
  - If type of potential is 'buckPlusAttr' : C (eV) and  $\rho$ (Å) D (eV),  $B$  (Å<sup>-1</sup>),  $r_1^{OO}$ (Å) and  $r_2^{OO}$ (Å)
- Divider line

## 4) Tables parameters:

- Cutoff radius for the Coulomb interaction ( $R_{coul}$ )
- Starting radius ( $r_{min} = 1,18845\text{Å}$ ) and increments ( $dr = 0.001\text{Å}$ ) for creating the potential table.
- Divider line

## 5) Rick model parameter:

- *Nevery* : parameter to set the frequency of the charge resolution. The charges are evaluated each *Nevery* time steps.
- Max number of iterative loop (*loopmax*) and convergence criterion (*prec*) in eV of the charge resolution
- Divider line

## 6) Coordination parameter:

- First ( $r_{1n}$ ) and second ( $r_{2n}$ ) neighbor distances in angstrom
  - Divider line
- 7) Charge initialization mode:
- Keyword (*QInitMode*) and initial oxygen charge ( $Q_{init}$ ). If keyword = 'true', all oxygen charges are initially set equal to  $Q_{init}$ . The charges on the cations are initially set in order to respect the neutrality of the box. If keyword = 'false', all atom charges are initially set equal to 0 if you use the *create\_atoms* command or the charge specified in the file structure using *read\_data* command.
  - Divider line
- 8) Mode for the electronegativity equalization (Qeq)
- Keyword (*mode*) followed by:
    - QEqAll (one QEq group) | no parameters
    - QEqAllParallel (several QEq groups) | no parameters
    - Surface | zlim (QEq only for  $z > zlim$ )
  - Parameter if necessary
  - Divider line
- 9) Verbose
- If you want the code to work in verbose mode or not : 'true' or 'false'
  - If you want to print or not in the file 'Energy\_component.txt' the three main contributions to the energy of the system according to the description presented above : 'true' or 'false' and  $N_{Energy}$ . This option writes to the file every  $N_{Energy}$  time steps. If the value is 'false' then  $N_{Energy} = 0$ . The file takes into account the possibility to have several QEq groups  $g$  then it writes: time step, number of atoms in group  $g$ , electrostatic part of energy,  $E_{ES}$ , the interaction between oxygen,  $E_{OO}$ , and short range metal-oxygen interaction,  $E_{MO}$ .
  - If you want to print to the file 'Electroneg\_component.txt' the electronegativity component ( $\frac{\partial E_{tot}}{\partial Q_i}$ ) or not: 'true' or 'false' and  $N_{Electroneg}$ . This option writes to the file every  $N_{Electroneg}$  time steps. If the value is 'false' then  $N_{Electroneg} = 0$ . The file consist of atom number  $i$ , atom type (1 for oxygen and # higher than 1 for metal), atom position:  $x$ ,  $y$  and  $z$ , atomic charge of atom  $i$ , electrostatic part of atom  $i$  electronegativity, covalent part of atom  $i$  electronegativity, the hopping integral of atom  $i$  ( $Z\beta^2$ ) $_i$  and box electronegativity.

---

**Note:** This last option slows down the calculation dramatically. Use only with a single processor simulation.

---

#### 19.227.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* mix, shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you needs to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.227.5 Restrictions

This pair style is part of the SMTBQ package and is only enabled if LAMMPS is built with that package. See the *Build package* page for more info.

This potential requires using atom type 1 for oxygen and atom type higher than 1 for metal atoms.

This pair style requires the *newton* setting to be “on” for pair interactions.

The SMTB-Q potential files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*.

### 19.227.6 Citing this work

Please cite related publication: N. Salles, O. Politano, E. Amzallag and R. Tetot, *Comput. Mater. Sci.* 111 (2016) 181-189

(SMTB-Q\_1) N. Salles, O. Politano, E. Amzallag, R. Tetot, *Comput. Mater. Sci.* 111 (2016) 181-189

(SMTB-Q\_2) E. Maras, N. Salles, R. Tetot, T. Ala-Nissila, H. Jonsson, *J. Phys. Chem. C* 2015, 119, 10391-10399

(SMTB-Q\_3) R. Tetot, N. Salles, S. Landron, E. Amzallag, *Surface Science* 616, 19-8722 28 (2013)

(Wolf) D. Wolf, P. Keblinski, S. R. Phillpot, J. Eggebrecht, *J Chem Phys*, 110, 8254 (1999).

(Rick) S. W. Rick, S. J. Stuart, B. J. Berne, *J Chem Phys* 101, 6141 (1994).

## 19.228 pair\_style snap command

Accelerator Variants: *snap/kk*

### 19.228.1 Syntax

```
pair_style snap
```

### 19.228.2 Examples

```
pair_style snap
pair_coeff * * InP.snapcoeff InP.snapparam In In P P
```

### 19.228.3 Description

Pair style *snap* defines the spectral neighbor analysis potential (SNAP), a machine-learning interatomic potential (*Thompson*). Like the GAP framework of Bartok et al. (*Bartok2010*), SNAP uses bispectrum components to characterize the local neighborhood of each atom in a very general way. The mathematical definition of the bispectrum calculation and its derivatives w.r.t. atom positions is identical to that used by *compute snap*, which is used to fit SNAP

potentials to *ab initio* energy, force, and stress data. In SNAP, the total energy is decomposed into a sum over atom energies. The energy of atom  $i$  is expressed as a weighted sum over bispectrum components.

$$E_{SNAP}^i(B_1^i, \dots, B_K^i) = \beta_0^{\mu_i} + \sum_{k=1}^K \beta_k^{\mu_i} B_k^i$$

where  $B_k^i$  is the  $k$ -th bispectrum component of atom  $i$ , and  $\beta_k^{\mu_i}$  is the corresponding linear coefficient that depends on  $\mu_i$ , the SNAP element of atom  $i$ . The number of bispectrum components used and their definitions depend on the value of *twojmax* and other parameters defined in the SNAP parameter file described below. The bispectrum calculation is described in more detail in [compute sna/atom](#).

Note that unlike for other potentials, cutoffs for SNAP potentials are not set in the `pair_style` or `pair_coeff` command; they are specified in the SNAP potential files themselves.

Only a single `pair_coeff` command is used with the *snap* style which specifies a SNAP coefficient file followed by a SNAP parameter file and then N additional arguments specifying the mapping of SNAP elements to LAMMPS atom types, where N is the number of LAMMPS atom types:

- SNAP coefficient file
- SNAP parameter file
- N element names = mapping of SNAP elements to atom types

As an example, if a LAMMPS indium phosphide simulation has 4 atoms types, with the first two being indium and the third and fourth being phosphorous, the `pair_coeff` command would look like this:

```
pair_coeff * * snap InP.snapcoeff InP.snapparam In In P P
```

The first 2 arguments must be `* *` so as to span all LAMMPS atom types. The two filenames are for the coefficient and parameter files, respectively. The two trailing ‘In’ arguments map LAMMPS atom types 1 and 2 to the SNAP ‘In’ element. The two trailing ‘P’ arguments map LAMMPS atom types 3 and 4 to the SNAP ‘P’ element.

If a SNAP mapping value is specified as NULL, the mapping is not performed. This can be used when a *snap* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

The name of the SNAP coefficient file usually ends in the “.snapcoeff” extension. It may contain coefficients for many SNAP elements. The only requirement is that each of the unique element names appearing in the LAMMPS `pair_coeff` command appear exactly once in the SNAP coefficient file. It is okay if the SNAP coefficient file contains additional elements not in the `pair_coeff` command, except when using *chemflag* (see below). The name of the SNAP parameter file usually ends in the “.snapparam” extension. It contains a small number of parameters that define the overall form of the SNAP potential. See the [pair\\_coeff](#) page for alternate ways to specify the path for these files.

SNAP potentials are quite commonly combined with one or more other LAMMPS pair styles using the *hybrid/overlay* pair style. As an example, the SNAP tantalum potential provided in the LAMMPS potentials directory combines the *snap* and *zbl* pair styles. It is invoked by the following commands:

```
variable zblcutinner equal 4
variable zblcutouter equal 4.8
variable zblz equal 73
pair_style hybrid/overlay &
zbl ${zblcutinner} ${zblcutouter} snap
pair_coeff * * zbl 0.0
pair_coeff 1 1 zbl ${zblz}
pair_coeff * * snap Ta06A.snapcoeff Ta06A.snapparam Ta
```

It is convenient to keep these commands in a separate file that can be inserted in any LAMMPS input script using the *include* command.

The top of the SNAP coefficient file can contain any number of blank and comment lines (start with #), but follows a strict format after that. The first non-blank non-comment line must contain two integers:

- `nelem` = Number of elements
- `ncoeff` = Number of coefficients

This is followed by one block for each of the *nelem* elements. The first line of each block contains three entries:

- Element name (text string)
- `R` = Element radius (distance units)
- `w` = Element weight (dimensionless)

This line is followed by *ncoeff* coefficients, one per line.

The SNAP parameter file can contain blank and comment lines (start with #) anywhere. Each non-blank non-comment line must contain one keyword/value pair. The required keywords are *rcutfac* and *twojmax*. Optional keywords are *rfac0*, *rmin0*, *switchflag*, *bzeroflag*, *quadraticflag*, *chemflag*, *bnormflag*, *wselfallflag*, *switchinnerflag*, *sinner*, *dinner*, *chunksize*, and *parallethresh*.

The default values for these keywords are

- *rfac0* = 0.99363
- *rmin0* = 0.0
- *switchflag* = 1
- *bzeroflag* = 1
- *quadraticflag* = 0
- *chemflag* = 0
- *bnormflag* = 0
- *wselfallflag* = 0
- *switchinnerflag* = 0
- *chunksize* = 32768
- *parallethresh* = 8192

For detailed definitions of all of these keywords, see the [compute sna/atom](#) doc page.

If *quadraticflag* is set to 1, then the SNAP energy expression includes additional quadratic terms that have been shown to increase the overall accuracy of the potential without much increase in computational cost ([Wood](#)).

$$E_{SNAP}^i(\mathbf{B}^i) = \beta_0^{\mu_i} + \beta^{\mu_i} \cdot \mathbf{B}_i + \frac{1}{2} \mathbf{B}_i^t \cdot \boldsymbol{\alpha}^{\mu_i} \cdot \mathbf{B}_i$$

where  $\mathbf{B}_i$  is the  $K$ -vector of bispectrum components,  $\beta^{\mu_i}$  is the  $K$ -vector of linear coefficients for element  $\mu_i$ , and  $\boldsymbol{\alpha}^{\mu_i}$  is the symmetric  $K$  by  $K$  matrix of quadratic coefficients. The SNAP coefficient file should contain  $K(K+1)/2$  additional coefficients in each element block, the upper-triangular elements of  $\boldsymbol{\alpha}^{\mu_i}$ .

If *chemflag* is set to 1, then the energy expression is written in terms of explicit multi-element bispectrum components indexed on ordered triplets of elements, which has been shown to increase the ability of the SNAP potential to capture energy differences in chemically complex systems, at the expense of a significant increase in computational cost ([Cusentino](#)).

$$E_{SNAP}^i(\mathbf{B}^i) = \beta_0^{\mu_i} + \sum_{\kappa, \lambda, \mu} \beta_{\mu_i}^{\kappa \lambda \mu} \cdot \mathbf{B}_i^{\kappa \lambda \mu}$$

where  $\mathbf{B}_i^{\kappa\lambda\mu}$  is the  $K$ -vector of bispectrum components for neighbors of elements  $\kappa$ ,  $\lambda$ , and  $\mu$  and  $\beta_{\mu_i}^{\kappa\lambda\mu}$  is the corresponding  $K$ -vector of linear coefficients for element  $\mu_i$ . The SNAP coefficient file should contain a total of  $KN_{elem}^3$  coefficients in each element block, where  $N_{elem}$  is the number of elements in the SNAP coefficient file, which must equal the number of unique elements appearing in the LAMMPS `pair_coeff` command, to avoid ambiguity in the number of coefficients.

The keyword `switchinnerflag` activates an additional switching function that smoothly turns off contributions to the SNAP potential from neighbor atoms at short separations. If `switchinnerflag` is set to 1 then the additional keywords `sinner` and `dinner` must also be provided. Each of these is followed by `nelements` values, where `nelements` is the number of unique elements appearing in the LAMMPS `pair_coeff` command. The element order should correspond to the order in which elements first appear in the `pair_coeff` command reading from left to right.

The keywords `chunksize` and `parallelthresh` are only applicable when using the pair style `snap` with the KOKKOS package on GPUs and are ignored otherwise. The `chunksize` keyword controls the number of atoms in each pass used to compute the bispectrum components and is used to avoid running out of memory. For example if there are 8192 atoms in the simulation and the `chunksize` is set to 4096, the bispectrum calculation will be broken up into two passes (running on a single GPU). The `parallelthresh` keyword controls a crossover threshold for performing extra parallelism. For small systems, exposing additional parallelism can be beneficial when there is not enough work to fully saturate the GPU threads otherwise. However, the extra parallelism also leads to more divergence and can hurt performance when the system is already large enough to saturate the GPU threads. Extra parallelism will be performed if the `chunksize` (or total number of atoms per GPU) is smaller than `parallelthresh`.

---

**Note:** The previously used `diagonalstyle` keyword was removed in 2019, since all known SNAP potentials use the default value of 3.

---

#### 19.228.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, mixing is performed by LAMMPS with user-specifiable parameters as described above. You never need to specify a `pair_coeff` command with I != J arguments for this style.

This pair style does not support the `pair_modify` shift, table, and tail options.

This pair style does not write its information to `binary restart files`, since it is stored in potential files. Thus, you need to re-specify the `pair_style` and `pair_coeff` commands in an input script that reads a restart file.

This pair style can only be used via the `pair` keyword of the `run_style respa` command. It does not support the `inner`, `middle`, `outer` keywords.

---

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix command-line switch` when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 19.228.5 Restrictions

This style is part of the ML-SNAP package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.228.6 Related commands

*compute sna/atom, compute snad/atom, compute snav/atom, compute snap*

### 19.228.7 Default

none

(Thompson) Thompson, Swiler, Trott, Foiles, Tucker, J Comp Phys, 285, 316 (2015).

(Bartok2010) Bartok, Payne, Kondor, Csanyi, Phys Rev Lett, 104, 136403 (2010).

(Wood) Wood and Thompson, J Chem Phys, 148, 241721, (2018)

(Cusentino) Cusentino, Wood, and Thompson, J Phys Chem A, xxx, xxxxx, (2020)

## 19.229 pair\_style soft command

Accelerator Variants: *soft/gpu, soft/omp*

### 19.229.1 Syntax

```
pair_style soft cutoff
```

- cutoff = global cutoff for soft interactions (distance units)

### 19.229.2 Examples

```
pair_style soft 1.0
pair_coeff * * 10.0
pair_coeff 1 1 10.0 3.0

pair_style soft 1.0
pair_coeff * * 0.0
variable prefactor equal ramp(0,30)
fix 1 all adapt 1 pair soft a * * v_prefactor
```

### 19.229.3 Description

Style *soft* computes pairwise interactions with the formula

$$E = A \left[ 1 + \cos \left( \frac{\pi r}{r_c} \right) \right] \quad r < r_c$$

It is useful for pushing apart overlapping atoms, since it does not blow up as  $r$  goes to 0.  $A$  is a prefactor that can be made to vary in time from the start to the end of the run (see discussion below), e.g. to start with a very soft potential and slowly harden the interactions over time.  $R_c$  is the cutoff. See the *fix nve/limit* command for another way to push apart overlapping atoms.

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $A$  (energy units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global soft cutoff is used.

---

**Note:** The syntax for *pair\_coeff* with a single  $A$  coeff is different in the current version of LAMMPS than in older versions which took two values,  $A_{start}$  and  $A_{stop}$ , to ramp between them. This functionality is now available in a more general form through the *fix adapt* command, as explained below. Note that if you use an old input script and specify  $A_{start}$  and  $A_{stop}$  without a cutoff, then LAMMPS will interpret that as  $A$  and a cutoff, which is probably not what you want.

---

The *fix adapt* command can be used to vary  $A$  for one or more pair types over the course of a simulation, in which case *pair\_coeff* settings for  $A$  must still be specified, but will be overridden. For example these commands will vary the prefactor  $A$  for all pairwise interactions from 0.0 at the beginning to 30.0 at the end of a run:

```
variable prefactor equal ramp(0,30)
fix 1 all adapt 1 pair soft a * * v_prefactor
```

Note that a formula defined by an *equal-style variable* can use the current timestep, elapsed time in the current run, elapsed time since the beginning of a series of runs, as well as access other variables.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---



### 19.229.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , the A coefficient and cutoff distance for this pair style can be mixed. A is always mixed via a *geometric* rule. The cutoff is mixed according to the `pair_modify` mix value. The default mix value is *geometric*. See the “`pair_modify`” command for details.

This pair style does not support the *pair\_modify* shift option, since the pair interaction goes to 0.0 at the cutoff.

The *pair\_modify* table and tail options are not relevant for this pair style.

This pair style writes its information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.229.5 Restrictions

none

### 19.229.6 Related commands

*pair\_coeff*, *fix nve/limit*, *fix adapt*

### 19.229.7 Default

none

## 19.230 pair\_style sph/heatconduction command

### 19.230.1 Syntax

```
pair_style sph/heatconduction
```

### 19.230.2 Examples

```
pair_style sph/heatconduction
pair_coeff * * 1.0 2.4
```

### 19.230.3 Description

The sph/heatconduction style computes heat transport between SPH particles. The transport model is the diffusion equation for the internal energy.

See [this PDF guide](#) to using SPH in LAMMPS.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above.

- D diffusion coefficient (length<sup>2</sup>/time units)
  - h kernel function cutoff (distance units)
- 

### 19.230.4 Mixing, shift, table, tail correction, restart, rRESPA info

This style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This style does not support the *pair\_modify* shift, table, and tail options.

This style does not write information to *binary restart files*. Thus, you need to re-specify the pair\_style and pair\_coeff commands in an input script that reads a restart file.

This style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.230.5 Restrictions

This pair style is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 19.230.6 Related commands

*pair\_coeff*, pair\_sph/rhosum

### 19.230.7 Default

none

## 19.231 pair\_style sph/idealgas command

### 19.231.1 Syntax

```
pair_style sph/idealgas
```

## 19.231.2 Examples

```
pair_style sph/idealgas
pair_coeff * * 1.0 2.4
```

## 19.231.3 Description

The sph/idealgas style computes pressure forces between particles according to the ideal gas equation of state:

$$p = (\gamma - 1)\rho e$$

where  $\gamma = 1.4$  is the heat capacity ratio,  $\rho$  is the local density, and  $e$  is the internal energy per unit mass. This pair style also computes Monaghan's artificial viscosity to prevent particles from interpenetrating (*Monaghan*).

See [this PDF guide](#) to using SPH in LAMMPS.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above.

- $\nu$  artificial viscosity (no units)
- h kernel function cutoff (distance units)

## 19.231.4 Mixing, shift, table, tail correction, restart, rRESPA info

This style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This style does not support the *pair\_modify* shift, table, and tail options.

This style does not write information to *binary restart files*. Thus, you need to re-specify the pair\_style and pair\_coeff commands in an input script that reads a restart file.

This style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

## 19.231.5 Restrictions

This pair style is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

## 19.231.6 Related commands

*pair\_coeff*, *pair\_sph/rhsum*

## 19.231.7 Default

none

---

(Monaghan) Monaghan and Gingold, Journal of Computational Physics, 52, 374-389 (1983).

## 19.232 pair\_style sph/lj command

### 19.232.1 Syntax

```
pair_style sph/lj
```

---

### 19.232.2 Examples

```
pair_style sph/lj
pair_coeff * * 1.0 2.4
```

---

### 19.232.3 Description

The sph/lj style computes pressure forces between particles according to the Lennard-Jones equation of state, which is computed according to Ree's 1980 polynomial fit (*Ree*). The Lennard-Jones parameters epsilon and sigma are set to unity. This pair style also computes Monaghan's artificial viscosity to prevent particles from interpenetrating (*Monaghan*).

See [this PDF guide](#) to using SPH in LAMMPS.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above.

- $\nu$  artificial viscosity (no units)
  - h kernel function cutoff (distance units)
- 

### 19.232.4 Mixing, shift, table, tail correction, restart, rRESPA info

This style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This style does not support the *pair\_modify* shift, table, and tail options.

This style does not write information to *binary restart files*. Thus, you need to re-specify the pair\_style and pair\_coeff commands in an input script that reads a restart file.

This style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.232.5 Restrictions

As noted above, the Lennard-Jones parameters epsilon and sigma are set to unity.

This pair style is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

### 19.232.6 Related commands

`pair_coeff`, `pair_sph/rhosum`

### 19.232.7 Default

none

---

**(Ree)** Ree, Journal of Chemical Physics, 73, 5401 (1980).

**(Monaghan)** Monaghan and Gingold, Journal of Computational Physics, 52, 374-389 (1983).

## 19.233 `pair_style sph/rhosum` command

### 19.233.1 Syntax

```
pair_style sph/rhosum Nstep
```

- Nstep = timestep interval

### 19.233.2 Examples

```
pair_style sph/rhosum 10  
pair_coeff * * 2.4
```

### 19.233.3 Description

The `sph/rhosum` style computes the local particle mass density rho for SPH particles by kernel function interpolation, every Nstep timesteps.

See [this PDF guide](#) to using SPH in LAMMPS.

The following coefficients must be defined for each pair of atoms types via the `pair_coeff` command as in the examples above.

- h (distance units)
-

### 19.233.4 Mixing, shift, table, tail correction, restart, rRESPA info

This style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This style does not support the *pair\_modify* shift, table, and tail options.

This style does not write information to *binary restart files*. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.233.5 Restrictions

This pair style is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.233.6 Related commands

*pair\_coeff*, *pair\_sph/taitwater*

### 19.233.7 Default

none

## 19.234 *pair\_style sph/taitwater* command

### 19.234.1 Syntax

```
pair_style sph/taitwater
```

### 19.234.2 Examples

```
pair_style sph/taitwater  
pair_coeff * * 1000.0 1430.0 1.0 2.4
```

### 19.234.3 Description

The *sph/taitwater* style computes pressure forces between SPH particles according to Tait's equation of state:

$$p = B \left[ \left( \frac{\rho}{\rho_0} \right)^\gamma - 1 \right]$$

where  $\gamma = 7$  and  $B = c_0^2 \rho_0 / \gamma$ , with  $\rho_0$  being the reference density and  $c_0$  the reference speed of sound.

This pair style also computes Monaghan's artificial viscosity to prevent particles from interpenetrating (*Monaghan*).

See [this PDF guide](#) to using SPH in LAMMPS.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above.

- $\rho_0$  reference density (mass/volume units)
  - $c_0$  reference soundspeed (distance/time units)
  - $\nu$  artificial viscosity (no units)
  - h kernel function cutoff (distance units)
- 

#### 19.234.4 Mixing, shift, table, tail correction, restart, rRESPA info

This style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This style does not support the *pair\_modify* shift, table, and tail options.

This style does not write information to *binary restart files*. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

#### 19.234.5 Restrictions

This pair style is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

#### 19.234.6 Related commands

*pair\_coeff*, *pair\_sph/rhosum*

#### 19.234.7 Default

none

---

(Monaghan) Monaghan and Gingold, Journal of Computational Physics, 52, 374-389 (1983).

### 19.235 pair\_style sph/taitwater/morris command

#### 19.235.1 Syntax

```
pair_style sph/taitwater/morris
```

## 19.235.2 Examples

```
pair_style sph/taitwater/morris
pair_coeff * * 1000.0 1430.0 1.0 2.4
```

## 19.235.3 Description

The sph/taitwater/morris style computes pressure forces between SPH particles according to Tait's equation of state:

$$p = B \left[ \left( \frac{\rho}{\rho_0} \right)^\gamma - 1 \right]$$

where  $\gamma = 7$  and  $B = c_0^2 \rho_0 / \gamma$ , with  $\rho_0$  being the reference density and  $c_0$  the reference speed of sound.

This pair style also computes laminar viscosity (*Morris*).

See [this PDF guide](#) to using SPH in LAMMPS.

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above.

- $\rho_0$  reference density (mass/volume units)
- $c_0$  reference soundspeed (distance/time units)
- $\nu$  dynamic viscosity (mass\*distance/time units)
- h kernel function cutoff (distance units)

---

## 19.235.4 Mixing, shift, table, tail correction, restart, rRESPA info

This style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

This style does not support the *pair\_modify* shift, table, and tail options.

This style does not write information to *binary restart files*. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

## 19.235.5 Restrictions

This pair style is part of the SPH package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.



## 19.235.6 Related commands

*pair\_coeff*, *pair\_sph/rhosome*

## 19.235.7 Default

none

---

(Morris) Morris, Fox, Zhu, J Comp Physics, 136, 214-226 (1997).

## 19.236 pair\_style lj/spica command

Accelerator Variants: *lj/spica/gpu*, *lj/spica/kk*, *lj/spica/omp*

## 19.237 pair\_style lj/spica/coul/long command

Accelerator Variants: *lj/spica/coul/long/gpu*, *lj/spica/coul/long/omp*

## 19.238 pair\_style lj/spica/coul/msm command

Accelerator Variants: *lj/spica/coul/msm/omp*

### 19.238.1 Syntax

```
pair_style style args
```

- style = *lj/spica* or *lj/spica/coul/long*
- args = list of arguments for a particular style

*lj/spica* args = cutoff

cutoff = global cutoff for Lennard Jones interactions (distance units)

*lj/spica/coul/long* args = cutoff (cutoff2)

cutoff = global cutoff for LJ (and Coulombic if only 1 arg) (distance units)

cutoff2 = global cutoff for Coulombic (optional) (distance units)

### 19.238.2 Examples

```
pair_style lj/spica 2.5
pair_coeff 1 1 lj12_6 1 1.1 2.8

pair_style lj/spica/coul/long 10.0
pair_style lj/spica/coul/long 10.0 12.0
pair_coeff 1 1 lj9_6 100.0 3.5 12.0
```

(continues on next page)

(continued from previous page)

```

pair_style lj/spica/coul/msm 10.0
pair_style lj/spica/coul/msm 10.0 12.0
pair_coeff 1 1 lj9_6 100.0 3.5 12.0

```

### 19.238.3 Description

The *lj/spica* styles compute a 9/6, 12/4, 12/5, or 12/6 Lennard-Jones potential, given by

$$E = \frac{27}{4}\epsilon \left[ \left(\frac{\sigma}{r}\right)^9 - \left(\frac{\sigma}{r}\right)^6 \right] \quad r < r_c$$

$$E = \frac{3\sqrt{3}}{2}\epsilon \left[ \left(\frac{\sigma}{r}\right)^{12} - \left(\frac{\sigma}{r}\right)^4 \right] \quad r < r_c$$

$$E = \frac{12}{7} \left(\frac{12}{5}\right)^{\left(\frac{5}{7}\right)} \epsilon \left[ \left(\frac{\sigma}{r}\right)^{12} - \left(\frac{\sigma}{r}\right)^5 \right] \quad r < r_c$$

$$E = 4\epsilon \left[ \left(\frac{\sigma}{r}\right)^{12} - \left(\frac{\sigma}{r}\right)^6 \right] \quad r < r_c$$

as required for the SPICA (formerly called SDK) and the pSPICA Coarse-grained MD parameterization discussed in (*Shinoda*), (*DeVane*), (*Seo*), and (*Miyazaki*).  $r_c$  is the cutoff. Summary information on these force fields can be found at <https://www.spica-ff.org>

Style *lj/spica/coul/long* computes the adds Coulombic interactions with an additional damping factor applied so it can be used in conjunction with the *kpace\_style* command and its *ewald* or *pppm* or *pppm/cg* option. The Coulombic cutoff specified for this style means that pairwise interactions within this distance are computed directly; interactions outside that distance are computed in reciprocal space.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- *cg\_type* (lj9\_6, lj12\_4, lj12\_5, or lj12\_6)
- *epsilon* (energy units)
- *sigma* (distance units)
- *cutoff1* (distance units)

Note that *sigma* is defined in the LJ formula as the zero-crossing distance for the potential, not as the energy minimum. The prefactors are chosen so that the potential minimum is at  $-\epsilon$ .

The latter 2 coefficients are optional. If not specified, the global LJ and Coulombic cutoffs specified in the *pair\_style* command are used. If only one cutoff is specified, it is used as the cutoff for both LJ and Coulombic interactions for this type pair. If both coefficients are specified, they are used as the LJ and Coulombic cutoffs for this type pair.

For *lj/spica/coul/long* and *lj/spica/coul/msm* only the LJ cutoff can be specified since a Coulombic cutoff cannot be specified for an individual I,J type pair. All type pairs use the same global Coulombic cutoff specified in the *pair\_style* command.

The original implementation of the above styles are style *lj/sdk*, *lj/sdk/coul/long*, and *lj/sdk/coul/msm*, and available for backward compatibility.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages*

page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.238.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of the lj/spica pair styles *cannot* be mixed, since different pairs may have different exponents. So all parameters for all pairs have to be specified explicitly through the “pair\_coeff” command. Defining them in a data file is also not supported, due to limitations of that file format.

All of the lj/spica pair styles support the *pair\_modify* shift option for the energy of the Lennard-Jones portion of the pair interaction.

The *lj/spica/coul/long* pair styles support the *pair\_modify* table option since they can tabulate the short-range portion of the long-range Coulombic interaction.

All of the lj/spica pair styles write their information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

The lj/spica and lj/cut/coul/long pair styles do not support the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command.

---

#### 19.238.5 Restrictions

All of the lj/spica pair styles are part of the CG-SPICA package. The *lj/spica/coul/long* style also requires the KSPACE package to be built (which is enabled by default). They are only enabled if LAMMPS was built with that package. See the *Build package* doc page for more info.

#### 19.238.6 Related commands

*pair\_coeff*, *angle\_style spica*

#### 19.238.7 Default

none

---

(Shinoda) Shinoda, DeVane, Klein, Mol Sim, 33, 27-36 (2007).

(DeVane) Shinoda, DeVane, Klein, Soft Matter, 4, 2453-2462 (2008).

(Seo) Seo, Shinoda, J Chem Theory Comput, 15, 762-774 (2019).

(Miyazaki) Miyazaki, Okazaki, Shinoda, J Chem Theory Comput, 16, 782-793 (2020).

## 19.239 pair\_style spin/dipole/cut command

## 19.240 pair\_style spin/dipole/long command

### 19.240.1 Syntax

```
pair_style spin/dipole/cut cutoff
pair_style spin/dipole/long cutoff
```

- cutoff = global cutoff for magnetic dipole energy and forces (optional) (distance units)

### 19.240.2 Examples

```
pair_style spin/dipole/cut 10.0
pair_coeff * * 10.0
pair_coeff 2 3 8.0

pair_style spin/dipole/long 9.0
pair_coeff * * 10.0
pair_coeff 2 3 6.0
```

### 19.240.3 Description

Style *spin/dipole/cut* computes a short-range dipole-dipole interaction between pairs of magnetic particles that each have a magnetic spin. The magnetic dipole-dipole interactions are computed by the following formulas for the magnetic energy, magnetic precession vector omega and mechanical force between particles I and J.

$$\mathcal{H}_{\text{long}} = -\frac{\mu_0(\mu_B)^2}{4\pi} \sum_{i,j,i \neq j}^N \frac{g_i g_j}{r_{ij}^3} \left( 3(\vec{e}_{ij} \cdot \vec{s}_i)(\vec{e}_{ij} \cdot \vec{s}_j) - \vec{s}_i \cdot \vec{s}_j \right)$$

$$\omega_i = \frac{\mu_0(\mu_B)^2}{4\pi\hbar} \sum_j \frac{g_i g_j}{r_{ij}^3} \left( 3(\vec{e}_{ij} \cdot \vec{s}_j)\vec{e}_{ij} - \vec{s}_j \right)$$

$$\mathbf{F}_i = \frac{3\mu_0(\mu_B)^2}{4\pi} \sum_j \frac{g_i g_j}{r_{ij}^4} \left[ ((\vec{s}_i \cdot \vec{s}_j) - 5(\vec{e}_{ij} \cdot \vec{s}_i)(\vec{e}_{ij} \cdot \vec{s}_j))\vec{e}_{ij} + ((\vec{e}_{ij} \cdot \vec{s}_i)\vec{s}_j + (\vec{e}_{ij} \cdot \vec{s}_j)\vec{s}_i) \right]$$

where  $\vec{s}_i$  and  $\vec{s}_j$  are the spin on two magnetic particles,  $r$  is their separation distance, and the vector  $\vec{e}_{ij} = \frac{\vec{r}_i - \vec{r}_j}{|\vec{r}_i - \vec{r}_j|}$  is the direction vector between the two particles.

Style *spin/dipole/long* computes long-range magnetic dipole-dipole interaction. A *kpace\_style* must be defined to use this pair style. Currently, *kpace\_style ewald/dipole/spin* and *kpace\_style ppm/dipole/spin* support long-range magnetic dipole-dipole interactions.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

## 19.240.4 Restrictions

The *spin/dipole/cut* and *spin/dipole/long* styles are part of the SPIN package. They are only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

Using dipole/spin pair styles with *electron units* is not currently supported.

## 19.240.5 Related commands

*pair\_coeff*, *kpace\_style fix nve/spin*

## 19.240.6 Default

none

## 19.241 pair\_style spin/dmi command

### 19.241.1 Syntax

```
pair_style spin/dmi cutoff
```

- cutoff = global cutoff pair (distance in metal units)

### 19.241.2 Examples

```
pair_style spin/dmi 4.0
pair_coeff * * dmi 2.6 0.001 1.0 0.0 0.0
pair_coeff 1 2 dmi 4.0 0.00109 0.0 0.0 1.0
```

### 19.241.3 Description

Style *spin/dmi* computes the Dzyaloshinskii-Moriya (DM) interaction between pairs of magnetic spins. According to the expression reported in ([Rohart](#)), one has the following DM energy:

$$\mathbf{H}_{dm} = \sum_{i,j=1,i \neq j}^N \left( \vec{e}_{ij} \times \vec{D} \right) \cdot (\vec{s}_i \times \vec{s}_j),$$

where  $\vec{s}_i$  and  $\vec{s}_j$  are two neighboring magnetic spins of two particles,  $\vec{e}_{ij} = \frac{r_i - r_j}{|r_i - r_j|}$  is the unit vector between sites  $i$  and  $j$ , and  $\vec{D}$  is the DM vector defining the intensity (in eV) and the direction of the interaction.

In ([Rohart](#)),  $\vec{D}$  is defined as the direction normal to the film oriented from the high spin-orbit layer to the magnetic ultra-thin film.

The application of a spin-lattice Poisson bracket to this energy (as described in ([Tranchida](#))) allows to derive a magnetic torque  $\vec{\omega}$ , and a mechanical force  $\vec{F}$  (for spin-lattice calculations only) for each magnetic particle  $i$ :

$$\vec{\omega}_i = -\frac{1}{\hbar} \sum_j^{Neighb} \vec{s}_j \times \left( \vec{e}_{ij} \times \vec{D} \right) \quad \text{and} \quad \vec{F}_i = - \sum_j^{Neighb} \frac{1}{r_{ij}} \vec{D} \times (\vec{s}_i \times \vec{s}_j)$$

More details about the derivation of these torques/forces are reported in (*Tranchida*).

For the *spin/dmi* pair style, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, and set in the following order:

- *rc* (distance units)
- $|D|$  (energy units)
- *Dx*, *Dy*, *Dz* (direction of *D*)

Note that *rc* is the radius cutoff of the considered DM interaction,  $|D|$  is the norm of the DM vector (in eV), and *Dx*, *Dy* and *Dz* define its direction.

None of those coefficients is optional. If not specified, the *spin/dmi* pair style cannot be used.

---

#### 19.241.4 Restrictions

All the *pair/spin* styles are part of the SPIN package. These styles are only enabled if LAMMPS was built with this package, and if the atom\_style “spin” was declared. See the *Build package* page for more info.

#### 19.241.5 Related commands

*atom\_style spin*, *pair\_coeff*, *pair\_eam*,

#### 19.241.6 Default

none

---

**(Rohart)** Rohart and Thiaville, Physical Review B, 88(18), 184422. (2013).

**(Tranchida)** Tranchida, Plimpton, Thibaudeau and Thompson, Journal of Computational Physics, 372, 406-425, (2018).

### 19.242 pair\_style spin/exchange command

### 19.243 pair\_style spin/exchange/biquadratic command

#### 19.243.1 Syntax

```
pair_style spin/exchange cutoff
pair_style spin/exchange/biquadratic cutoff
```

- cutoff = global cutoff pair (distance in metal units)

## 19.243.2 Examples

```

pair_style spin/exchange 4.0
pair_coeff * * exchange 4.0 0.0446928 0.003496 1.4885
pair_coeff 1 2 exchange 6.0 -0.01575 0.0 1.965 offset yes

pair_style spin/exchange/biquadratic 4.0
pair_coeff * * biquadratic 4.0 0.05 0.03 1.48 0.05 0.03 1.48 offset no
pair_coeff 1 2 biquadratic 6.0 -0.01 0.0 1.9 0.0 0.1 19

```

## 19.243.3 Description

Style *spin/exchange* computes the exchange interaction between pairs of magnetic spins:

$$H_{ex} = - \sum_{i,j}^N J_{ij}(r_{ij}) \vec{s}_i \cdot \vec{s}_j$$

where  $\vec{s}_i$  and  $\vec{s}_j$  are two unit vectors representing the magnetic spins of two particles (usually atoms), and  $r_{ij} = |\vec{r}_i - \vec{r}_j|$  is the inter-atomic distance between those two particles. The summation is over pairs of nearest neighbors.  $J(r_{ij})$  is a function defining the intensity and the sign of the exchange interaction for different neighboring shells.

Style *spin/exchange/biquadratic* computes a biquadratic exchange interaction between pairs of magnetic spins:

$$H_{bi} = - \sum_{i,j}^N J_{ij}(r_{ij}) \vec{s}_i \cdot \vec{s}_j - \sum_{i,j}^N K_{ij}(r_{ij}) (\vec{s}_i \cdot \vec{s}_j)^2$$

where  $\vec{s}_i$ ,  $\vec{s}_j$ ,  $r_{ij}$  and  $J(r_{ij})$  have the same definitions as above, and  $K(r_{ij})$  is a second function, defining the intensity and the sign of the biquadratic term.

The interatomic dependence of  $J(r_{ij})$  and  $K(r_{ij})$  in both interactions above is defined by the following function:

$$f(r_{ij}) = 4a \left(\frac{r_{ij}}{d}\right)^2 \left(1 - b \left(\frac{r_{ij}}{d}\right)^2\right) e^{-\left(\frac{r_{ij}}{d}\right)^2} \Theta(R_c - r_{ij})$$

where  $a$ ,  $b$  and  $d$  are the three constant coefficients defined in the associated “pair\_coeff” command, and  $R_c$  is the radius cutoff associated to the pair interaction (see below for more explanations).

The coefficients  $a$ ,  $b$ , and  $d$  need to be fitted so that the function above matches with the value of the exchange interaction for the  $N$  neighbor shells taken into account. Examples and more explanations about this function and its parameterization are reported in ([Tranchida](#)).

When a *spin/exchange/biquadratic* pair style is defined, six coefficients (three for  $J(r_{ij})$ , and three for  $K(r_{ij})$ ) have to be fitted.

From this exchange interaction, each spin  $i$  will be submitted to a magnetic torque  $\vec{\omega}_i$ , and its associated atom can be submitted to a force  $\vec{F}_i$  for spin-lattice calculations (see *fix nve/spin*), such as:

$$\vec{\omega}_i = \frac{1}{\hbar} \sum_j^{Neighb} J(r_{ij}) \vec{s}_j \quad \text{and} \quad \vec{F}_i = \sum_j^{Neighb} \frac{\partial J(r_{ij})}{\partial r_{ij}} (\vec{s}_i \cdot \vec{s}_j) \vec{e}_{ij}$$

with  $\hbar$  the Planck constant (in metal units), and  $\vec{e}_{ij} = \frac{\vec{r}_i - \vec{r}_j}{|\vec{r}_i - \vec{r}_j|}$  the unit vector between sites  $i$  and  $j$ . Equivalent forces and magnetic torques are generated for the biquadratic term when a *spin/exchange/biquadratic* pair style is defined.

More details about the derivation of these torques/forces are reported in ([Tranchida](#)).

For the *spin/exchange* and *spin/exchange/biquadratic* pair styles, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, and set in the following order:

- $R_c$  (distance units)
- $a$  (energy units)
- $b$  (adim parameter)
- $d$  (distance units)

for the *spin/exchange* pair style, and:

- $R_c$  (distance units)
- $a_j$  (energy units)
- $b_j$  (adim parameter)
- $d_j$  (distance units)
- $a_k$  (energy units)
- $b_k$  (adim parameter)
- $d_k$  (distance units)

for the *spin/exchange/biquadratic* pair style.

Note that  $R_c$  is the radius cutoff of the considered exchange interaction, and  $a$ ,  $b$  and  $d$  are the three coefficients performing the parameterization of the function  $J(r_{ij})$  defined above (in the *biquadratic* style,  $a_j$ ,  $b_j$ ,  $d_j$  and  $a_k$ ,  $b_k$ ,  $d_k$  are the coefficients of  $J(r_{ij})$  and  $K(r_{ij})$  respectively).

None of those coefficients is optional. If not specified, the *spin/exchange* pair style cannot be used.

---

### Offsetting magnetic forces and energies:

For spin-lattice simulation, it can be useful to offset the mechanical forces and energies generated by the exchange interaction. The *offset* keyword allows to apply this offset. By setting *offset* to *yes*, the energy definitions above are replaced by:

$$H_{ex} = - \sum_{i,j}^N J_{ij}(r_{ij}) [\vec{s}_i \cdot \vec{s}_j - 1]$$

for the *spin/exchange* pair style, and:

$$H_{bi} = - \sum_{i,j}^N J_{ij}(r_{ij}) [\vec{s}_i \cdot \vec{s}_j - 1] - \sum_{i,j}^N K_{ij}(r_{ij}) [(\vec{s}_i \cdot \vec{s}_j)^2 - 1]$$

for the *spin/exchange/biquadratic* pair style.

Note that this offset only affects the calculation of the energy and mechanical forces. It does not modify the calculation of the precession vectors (and thus does not impact the purely magnetic properties). This ensures that when all spins are aligned, the magnetic energy and the associated mechanical forces (and thus the pressure generated by the magnetic potential) are null.

---

**Note:** This offset term can be very important when calculations such as equations of state (energy vs volume, or energy vs pressure) are being performed. Indeed, setting the *offset* term ensures that at the ground state of the crystal

---



and at the equilibrium magnetic configuration (typically ferromagnetic), the pressure is null, as expected. Otherwise, magnetic forces could generate a residual pressure.

When the *offset* option is set to *no*, no offset is applied (also corresponding to the default option).

### 19.243.4 Restrictions

All the *pair/spin* styles are part of the SPIN package. These styles are only enabled if LAMMPS was built with this package, and if the atom\_style “spin” was declared. See the *Build package* page for more info.

### 19.243.5 Related commands

*atom\_style spin*, *pair\_coeff*, *pair\_eam*,

### 19.243.6 Default

The default *offset* keyword value is *no*.

(Tranchida) Tranchida, Plimpton, Thibaudau and Thompson, Journal of Computational Physics, 372, 406-425, (2018).

## 19.244 pair\_style spin/magelec command

### 19.244.1 Syntax

```
pair_style spin/magelec cutoff
```

- cutoff = global cutoff pair (distance in metal units)

### 19.244.2 Examples

```
pair_style spin/magelec 4.5
pair_coeff * * magelec 4.5 0.00109 1.0 1.0 1.0
```

### 19.244.3 Description

Style *spin/me* computes a magneto-electric interaction between pairs of magnetic spins. According to the derivation reported in (*Katsura*), this interaction is defined as:

$$\vec{\omega}_i = -\frac{1}{\hbar} \sum_j^{Neighb} \vec{s}_j \times \vec{D}(r_{ij})$$

$$\vec{F}_i = - \sum_j^{Neighb} \frac{\partial D(r_{ij})}{\partial r_{ij}} (\vec{s}_i \times \vec{s}_j) \cdot \vec{r}_{ij}$$

where  $\vec{s}_i$  and  $\vec{s}_j$  are neighboring magnetic spins of two particles.

From this magneto-electric interaction, each spin  $i$  will be submitted to a magnetic torque  $\vec{\omega}$ , and its associated atom can be submitted to a force  $\vec{F}$  for spin-lattice calculations (see *fix nve/spin*), such as:

$$\vec{F}^i = - \sum_j^{Neighbor} (\vec{s}_i \times \vec{s}_j) \times \vec{E}$$
$$\vec{\omega}^i = -\frac{1}{\hbar} \sum_j^{Neighbor} \vec{s}_j \times (\vec{E} \times r_{ij})$$

with  $\hbar$  the Planck constant (in metal units) and  $\vec{E}$  an electric polarization vector. The norm and direction of  $\vec{E}$  are giving the intensity and the direction of a screened dielectric atomic polarization (in eV).

More details about the derivation of these torques/forces are reported in (*Tranchida*).

---

## 19.244.4 Restrictions

All the *pair/spin* styles are part of the SPIN package. These styles are only enabled if LAMMPS was built with this package, and if the atom\_style “spin” was declared. See the *Build package* page for more info.

## 19.244.5 Related commands

*atom\_style spin*, *pair\_coeff*, *pair\_style spin/exchange*, *pair\_eam*,

## 19.244.6 Default

none

---

**(Katsura)** H. Katsura, N. Nagaosa, A.V. Balatsky. Phys. Rev. Lett., 95(5), 057205. (2005)

**(Tranchida)** Tranchida, Plimpton, Thibaudau, and Thompson, Journal of Computational Physics, 372, 406-425, (2018).

## 19.245 pair\_style spin/neel command

### 19.245.1 Syntax

```
pair_style spin/neel cutoff
```

- cutoff = global cutoff pair (distance in metal units)

## 19.245.2 Examples

```
pair_style spin/neel 4.0
pair_coeff * * neel 4.0 0.0048 0.234 1.168 2.6905 0.705 0.652
pair_coeff 1 2 neel 4.0 0.0048 0.234 1.168 0.0 0.0 1.0
```

## 19.245.3 Description

Style *spin/neel* computes the Neel pair anisotropy model between pairs of magnetic spins:

$$\mathcal{H}_{Neel} = - \sum_{i,j=1,i \neq j}^N g_1(r_{ij}) \left( (\mathbf{e}_{ij} \cdot \mathbf{s}_i)(\mathbf{e}_{ij} \cdot \mathbf{s}_j) - \frac{\mathbf{s}_i \cdot \mathbf{s}_j}{3} \right) + q_1(r_{ij}) \left( (\mathbf{e}_{ij} \cdot \mathbf{s}_i)^2 - \frac{\mathbf{s}_i \cdot \mathbf{s}_j}{3} \right) \left( (\mathbf{e}_{ij} \cdot \mathbf{s}_i)^2 - \frac{\mathbf{s}_i \cdot \mathbf{s}_j}{3} \right) + q_2(r_{ij}) \left( (\mathbf{e}_{ij} \cdot \mathbf{s}_i) \right)$$

where  $\mathbf{s}_i$  and  $\mathbf{s}_j$  are two neighboring magnetic spins of two particles,  $r_{ij} = |\mathbf{r}_i - \mathbf{r}_j|$  is the inter-atomic distance between the two particles,  $\mathbf{e}_{ij} = \frac{\mathbf{r}_i - \mathbf{r}_j}{|\mathbf{r}_i - \mathbf{r}_j|}$  is their normalized separation vector and  $g_1$ ,  $q_1$  and  $q_2$  are three functions defining the intensity of the dipolar and quadrupolar contributions, with:

$$\begin{aligned} g_1(r_{ij}) &= g(r_{ij}) + \frac{12}{35}q(r_{ij}) \\ q_1(r_{ij}) &= \frac{9}{5}q(r_{ij}) \\ q_2(r_{ij}) &= -\frac{2}{5}q(r_{ij}) \end{aligned}$$

With the functions  $g(r_{ij})$  and  $q(r_{ij})$  defined and fitted according to the same Bethe-Slater function used to fit the exchange interaction:

$$J(r_{ij}) = 4a \left( \frac{r_{ij}}{d} \right)^2 \left( 1 - b \left( \frac{r_{ij}}{d} \right)^2 \right) e^{-\left( \frac{r_{ij}}{d} \right)^2} \Theta(R_c - r_{ij})$$

where  $a$ ,  $b$  and  $d$  are the three constant coefficients defined in the associated “pair\_coeff” command.

The coefficients  $a$ ,  $b$ , and  $d$  need to be fitted so that the function above matches with the values of the magneto-elastic constant of the materials at stake.

Examples and more explanations about this function and its parameterization are reported in (*Tranchida*). More examples of parameterization will be provided in future work.

From this DM interaction, each spin  $i$  will be submitted to a magnetic torque  $\omega$  and its associated atom to a force  $\mathbf{F}$  (for spin-lattice calculations only).

More details about the derivation of these torques/forces are reported in (*Tranchida*).

## 19.245.4 Restrictions

All the *pair/spin* styles are part of the SPIN package. These styles are only enabled if LAMMPS was built with this package, and if the atom\_style “spin” was declared. See the *Build package* page for more info.

## 19.245.5 Related commands

*atom\_style spin, pair\_coeff, pair\_eam,*

## 19.245.6 Default

none

---

(Tranchida) Tranchida, Plimpton, Thibaudau and Thompson, Journal of Computational Physics, 372, 406-425, (2018).

## 19.246 pair\_style srp command

## 19.247 pair\_style srp/react command

### 19.247.1 Syntax

```
pair_style srp cutoff btype dist keyword value ...
pair_style srp/react cutoff btype dist react-id keyword value ...
```

- cutoff = global cutoff for SRP interactions (distance units)
- btype = bond type to apply SRP interactions to (can be wildcard, see below)
- distance = *min* or *mid*
- react-id = id of either fix bond/break or fix bond/create
- zero or more keyword/value pairs may be appended
- keyword = *exclude*

*bptype* value = atom type for bond particles  
*exclude* value = *yes* or *no*

### 19.247.2 Examples

```
pair_style hybrid dpd 1.0 1.0 12345 srp 0.8 1 mid exclude yes
pair_coeff 1 1 dpd 60.0 4.5 1.0
pair_coeff 1 2 none
pair_coeff 2 2 srp 100.0 0.8

pair_style hybrid dpd 1.0 1.0 12345 srp 0.8 * min exclude yes
pair_coeff 1 1 dpd 60.0 50 1.0
pair_coeff 1 2 none
pair_coeff 2 2 srp 40.0

fix          create all bond/create 100 1 2 1.0 1 prob 0.2 19852
pair_style hybrid dpd 1.0 1.0 12345 srp/react 0.8 * min create exclude yes
pair_coeff 1 1 dpd 60.0 50 1.0
```

(continues on next page)

(continued from previous page)

```

pair_coeff 1 2 none
pair_coeff 2 2 srp/react 40.0

pair_style hybrid srp 0.8 2 mid
pair_coeff 1 1 none
pair_coeff 1 2 none
pair_coeff 2 2 srp 100.0 0.8

```

### Description

Style *srp* computes a soft segmental repulsive potential (SRP) that acts between pairs of bonds. This potential is useful for preventing bonds from passing through one another when a soft non-bonded potential acts between beads in, for example, DPD polymer chains. An example input script that uses this command is provided in `examples/PACKAGES/srp`.

Bonds of specified type *btype* interact with one another through a bond-pairwise potential, such that the force on bond *i* due to bond *j* is as follows

$$F_{ij}^{\text{SRP}} = C(1 - r/r_c)\hat{r}_{ij} \quad r < r_c$$

where  $r$  and  $\hat{r}_{ij}$  are the distance and unit vector between the two bonds. Note that *btype* can be specified as an asterisk “\*”, which case the interaction is applied to all bond types. The *mid* option computes  $r$  and  $\hat{r}_{ij}$  from the midpoint distance between bonds. The *min* option computes  $r$  and  $\hat{r}_{ij}$  from the minimum distance between bonds. The force acting on a bond is mapped onto the two bond atoms according to the lever rule,

$$F_{i1}^{\text{SRP}} = F_{ij}^{\text{SRP}}(L)$$

$$F_{i2}^{\text{SRP}} = F_{ij}^{\text{SRP}}(1 - L)$$

where  $L$  is the normalized distance from the atom to the point of closest approach of bond *i* and *j*. The *mid* option takes  $L$  as 0.5 for each interaction as described in (Sirk).

The following coefficients must be defined via the *pair\_coeff* command as in the examples above, or in the data file or restart file read by the *read\_data* or *read\_restart* commands:

- $C$  (force units)
- $r_c$  (distance units)

The last coefficient is optional. If not specified, the global cutoff is used.

---

**Note:** Pair style *srp* considers each bond of type *btype* to be a fictitious “particle” of type *bptype*, where *bptype* is either the largest atom type in the system, or the type set by the *bptype* flag. Any actual existing particles with this atom type will be deleted at the beginning of a run. This means you must specify the number of types in your system accordingly; usually to be one larger than what would normally be the case, e.g. via the *create\_box* or by changing the header in your *data file*. The fictitious “bond particles” are inserted at the beginning of the run, and serve as placeholders that define the position of the bonds. This allows neighbor lists to be constructed and pairwise interactions to be computed in almost the same way as is done for actual particles. Because bonds interact only with other bonds, *pair\_style hybrid* should be used to turn off interactions between atom type *bptype* and all other types of atoms. An error will be flagged if *pair\_style hybrid* is not used.

---

The optional *exclude* keyword determines if forces are computed between first neighbor (directly connected) bonds. For a setting of *no*, first neighbor forces are computed; for *yes* they are not computed. A setting of *no* cannot be used with the *min* option for distance calculation because the minimum distance between directly connected bonds is zero.

Pair style *srp* turns off normalization of thermodynamic properties by particle number, as if the command *thermo\_modify norm no* had been issued.

The pairwise energy associated with style *srp* is shifted to be zero at the cutoff distance  $r_c$ .

---

Pair style *srp/react* interfaces the pair style *srp* with the bond breaking and formation mechanisms provided by fix *bond/break* and fix *bond/create*, respectively. When using this pair style, whenever a bond breaking (or formation) reaction occurs, the corresponding fictitious particle is deleted (or inserted) during the same simulation time step as the reaction. This is useful in the simulation of reactive systems involving large polymeric molecules (*Palkar*) where the segmental repulsive potential is necessary to minimize topological violations, and also needs to be turned on and off according to the progress of the reaction.

---

### 19.247.3 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support mixing.

This pair style does not support the *pair\_modify* shift option for the energy of the pair interaction. Note that as discussed above, the energy term is already shifted to be 0.0 at the cutoff distance  $r_c$ .

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes global and per-atom information to *binary restart files*. Pair *srp* should be used with *pair\_style hybrid*, thus the *pair\_coeff* commands need to be specified in the input script when reading a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.247.4 Restrictions

This pair style is part of the MISC package. It is only enabled if LAMMPS was built with that package. See the Making LAMMPS section for more info.

This pair style must be used with *pair\_style hybrid*.

This pair style requires the *newton* command to be *on* for non-bonded interactions.

This pair style is not compatible with *rigid body integrators*

### 19.247.5 Related commands

*pair\_style hybrid*, *pair\_coeff*, *pair dpd*

### 19.247.6 Default

The default keyword value is *exclude = yes*.

---

(**Sirk**) Sirk TW, Sliozberg YR, Brennan JK, Lisal M, Andzelm JW, J Chem Phys, 136 (13) 134903, 2012.

(**Palkar**) Palkar V, Kuksenok O, J. Phys. Chem. B, 126 (1), 336-346, 2022

## 19.248 pair\_style sw command

Accelerator Variants: *sw/gpu, sw/intel, sw/kk, sw/omp*

## 19.249 pair\_style sw/mod command

Accelerator Variants: *sw/mod/omp*

### 19.249.1 Syntax

```
pair_style style keyword values
```

- style = *sw* or *sw/mod*
- keyword = *maxdelcs* or *threebody*

*maxdelcs* value = delta1 delta2 (optional, *sw/mod* only)

delta1 = The minimum threshold for the variation of cosine of three-body angle

delta2 = The maximum threshold for the variation of cosine of three-body angle

*threebody* value = *on* or *off* (optional, *sw* only)

*on* (default) = Compute both the three-body and two-body terms of the potential

*off* = Compute only the two-body term of the potential

### 19.249.2 Examples

```
pair_style sw
pair_coeff * * si.sw Si
pair_coeff * * GaN.sw Ga N Ga

pair_style sw/mod maxdelcs 0.25 0.35
pair_coeff * * tmd.sw.mod Mo S S

pair_style hybrid sw threebody on sw threebody off
pair_coeff * * sw 1 mW_xL.sw mW NULL
pair_coeff 1 2 sw 2 mW_xL.sw mW xL
pair_coeff 2 2 sw 2 mW_xL.sw mW xL
```

### 19.249.3 Description

The *sw* style computes a 3-body *Stillinger-Weber* potential for the energy  $E$  of a system of atoms as

$$E = \sum_i \sum_{j>i} \phi_2(r_{ij}) + \sum_i \sum_{j \neq i} \sum_{k>j} \phi_3(r_{ij}, r_{ik}, \theta_{ijk})$$

$$\phi_2(r_{ij}) = A_{ij} \epsilon_{ij} \left[ B_{ij} \left( \frac{\sigma_{ij}}{r_{ij}} \right)^{p_{ij}} - \left( \frac{\sigma_{ij}}{r_{ij}} \right)^{q_{ij}} \right] \exp \left( \frac{\sigma_{ij}}{r_{ij} - a_{ij} \sigma_{ij}} \right)$$

$$\phi_3(r_{ij}, r_{ik}, \theta_{ijk}) = \lambda_{ijk} \epsilon_{ijk} [\cos \theta_{ijk} - \cos \theta_{0ijk}]^2 \exp \left( \frac{\gamma_{ij} \sigma_{ij}}{r_{ij} - a_{ij} \sigma_{ij}} \right) \exp \left( \frac{\gamma_{ik} \sigma_{ik}}{r_{ik} - a_{ik} \sigma_{ik}} \right)$$

where  $\phi_2$  is a two-body term and  $\phi_3$  is a three-body term. The summations in the formula are over all neighbors J and K of atom I within a cutoff distance  $a'\sigma$ .

New in version 14Dec2021.

The *sw/mod* style is designed for simulations of materials when distinguishing three-body angles are necessary, such as borophene and transition metal dichalcogenides, which cannot be described by the original code for the Stillinger-Weber potential. For instance, there are several types of angles around each Mo atom in *MoS\_2*, and some unnecessary angle types should be excluded in the three-body interaction. Such exclusion may be realized by selecting proper angle types directly. The exclusion of unnecessary angles is achieved here by the cut-off function ( $f_C(\delta)$ ), which induces only minimum modifications for LAMMPS.

Validation, benchmark tests, and applications of the *sw/mod* style can be found in (Jiang2) and (Jiang3).

The *sw/mod* style computes the energy E of a system of atoms, whose potential function is mostly the same as the Stillinger-Weber potential. The only modification is in the three-body term, where the value of  $\delta = \cos \theta_{ijk} - \cos \theta_{0ijk}$  used in the original energy and force expression is scaled by a switching factor  $f_C(\delta)$ :

$$f_C(\delta) = \begin{cases} 1 & : |\delta| < \delta_1 \\ \frac{1}{2} + \frac{1}{2} \cos\left(\pi \frac{|\delta| - \delta_1}{\delta_2 - \delta_1}\right) & : \delta_1 < |\delta| < \delta_2 \\ 0 & : |\delta| > \delta_2 \end{cases}$$

This cut-off function decreases smoothly from 1 to 0 over the range  $[\delta_1, \delta_2]$ . This smoothly turns off the energy and force contributions for  $|\delta| > \delta_2$ . It is suggested that  $\delta_1$  and  $\delta_2$  to be the value around  $0.5 |\cos \theta_1 - \cos \theta_2|$ , with  $\theta_1$  and  $\theta_2$  as the different types of angles around an atom. For borophene and transition metal dichalcogenides,  $\delta_1 = 0.25$  and  $\delta_2 = 0.35$ . This value enables the cut-off function to exclude unnecessary angles in the three-body SW terms.

---

**Note:** The cut-off function is just to be used as a technique to exclude some unnecessary angles, and it has no physical meaning. It should be noted that the force and potential are inconsistent with each other in the decaying range of the cut-off function, as the angle dependence for the cut-off function is not implemented in the force (first derivation of potential). However, the angle variation is much smaller than the given threshold value for actual simulations, so the inconsistency between potential and force can be neglected in actual simulations.

---

The *threebody* keyword is optional and determines whether or not the three-body term of the potential is calculated. The default value is “on” and it is only available for the plain *sw* pair style variants, but not available for the *sw/mod* and *sw/angle/table* pair style variants. To turn off the threebody contributions all  $\lambda_{ijk}$  parameters from the potential file are forcibly set to 0. In addition the pair style implementation may employ code optimizations for the *threebody off* setting that can result in significant speedups versus the default. These code optimizations are currently only available for the MANYBODY and OPENMP packages.

Only a single `pair_coeff` command is used with the *sw* and *sw/mod* styles which specifies a Stillinger-Weber potential file with parameters for all needed elements, except for when the *threebody off* setting is used (see note below). These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of SW elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the potential file.

As an example, imagine a file `SiC.sw` has Stillinger-Weber values for Si and C. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Si, and the fourth to be C, you would use the following `pair_coeff` command:

```
pair_style sw
pair_coeff * * SiC.sw Si Si Si C
```



The first 2 arguments must be \* \* so as to span all LAMMPS atom types. The first three Si arguments map LAMMPS atom types 1, 2, and 3 to the Si element in the SW file. The final C argument maps LAMMPS atom type 4 to the C element in the SW file. If an argument value is specified as NULL, the mapping is not performed. This can be used when an *sw* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

---

**Note:** When the *threebody off* keyword is used, multiple *pair\_coeff* commands may be used to specific the pairs of atoms which don't require three-body term. In these cases, the first 2 arguments are not required to be \* \*.

---

Stillinger-Weber files in the *potentials* directory of the LAMMPS distribution have a “.sw” suffix. Lines that are not blank or comments (starting with #) define parameters for a triplet of elements. The parameters in a single entry correspond to the two-body and three-body coefficients in the formula above:

- element 1 (the center atom in a 3-body interaction)
- element 2
- element 3
- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- $a$
- $\lambda$
- $\gamma$
- $\cos \theta_0$
- A
- B
- p
- q
- tol

The A, B, p, and q parameters are used only for two-body interactions. The  $\lambda$  and  $\cos \theta_0$  parameters are used only for three-body interactions. The  $\epsilon$ ,  $\sigma$  and  $a$  parameters are used for both two-body and three-body interactions.  $\gamma$  is used only in the three-body interactions, but is defined for pairs of atoms. The non-annotated parameters are unitless.

LAMMPS introduces an additional performance-optimization parameter *tol* that is used for both two-body and three-body interactions. In the Stillinger-Weber potential, the interaction energies become negligibly small at atomic separations substantially less than the theoretical cutoff distances. LAMMPS therefore defines a virtual cutoff distance based on a user defined tolerance *tol*. The use of the virtual cutoff distance in constructing atom neighbor lists can significantly reduce the neighbor list sizes and therefore the computational cost. LAMMPS provides a *tol* value for each of the three-body entries so that they can be separately controlled. If *tol* = 0.0, then the standard Stillinger-Weber cutoff is used.

The Stillinger-Weber potential file must contain entries for all the elements listed in the *pair\_coeff* command. It can also contain entries for additional elements not being used in a particular simulation; LAMMPS ignores those entries.

For a single-element simulation, only a single entry is required (e.g. SiSiSi). For a two-element simulation, the file must contain 8 entries (for SiSiSi, SiSiC, SiCSi, SiCC, CSiSi, CSiC, CCSi, CCC), that specify SW parameters for all permutations of the two elements interacting in three-body configurations. Thus for 3 elements, 27 entries would be required, etc.

As annotated above, the first element in the entry is the center atom in a three-body interaction. Thus an entry for SiCC means a Si atom with 2 C atoms as neighbors. The parameter values used for the two-body interaction come from the entry where the second and third elements are the same. Thus the two-body parameters for Si interacting with C, comes from the SiCC entry. The three-body parameters can in principle be specific to the three elements of the configuration. In the literature, however, the three-body parameters are usually defined by simple formulas involving two sets of pairwise parameters, corresponding to the ij and ik pairs, where i is the center atom. The user must ensure that the correct combining rule is used to calculate the values of the three-body parameters for alloys. Note also that the function  $\phi_3$  contains two exponential screening factors with parameter values from the ij pair and ik pairs. So  $\phi_3$  for a C atom bonded to a Si atom and a second C atom will depend on the three-body parameters for the CSiC entry, and also on the two-body parameters for the CCC and CSiSi entries. Since the order of the two neighbors is arbitrary, the three-body parameters for entries CSiC and CCSi should be the same. Similarly, the two-body parameters for entries SiCC and CSiSi should also be the same. The parameters used only for two-body interactions (A, B, p, and q) in entries whose second and third element are different (e.g. SiCSi) are not used for anything and can be set to 0.0 if desired. This is also true for the parameters in  $\phi_3$  that are taken from the ij and ik pairs ( $\sigma$ ,  $a$ ,  $\gamma$ )

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

**Note:** When using the INTEL package with this style, there is an additional 5 to 10 percent performance improvement when the Stillinger-Weber parameters p and q are set to 4 and 0 respectively. These parameters are common for modeling silicon and water.

---

#### 19.249.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, mixing is performed by LAMMPS as described above from values in the potential file.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

The *single()* function of the *sw* pair style is only enabled and supported for the case of the *threebody off* setting.

---

## 19.249.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

The Stillinger-Weber potential files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use the *sw* or *sw/mod* pair styles with any LAMMPS units, but you would need to create your own SW potential file with coefficients listed in the appropriate units if your simulation does not use “metal” units. If the potential file contains a ‘UNITS:’ metadata tag in the first line of the potential file, then LAMMPS can convert it transparently between “metal” and “real” units.

## 19.249.6 Related commands

*pair\_coeff*

## 19.249.7 Default

The default value for the *threebody* setting of the “sw” pair style is “on”, the default values for the “*maxdelcs*” setting of the *sw/mod* pair style are *delta1* = 0.25 and *delta2* = 0.35.

---

**(Stillinger)** Stillinger and Weber, Phys Rev B, 31, 5262 (1985).

**(Jiang2)** J.-W. Jiang, Nanotechnology 26, 315706 (2015).

**(Jiang3)** J.-W. Jiang, Acta Mech. Solida. Sin 32, 17 (2019).

## 19.250 pair\_style sw/angle/table command

### 19.250.1 Syntax

```
pair_style style
```

- style = *sw/angle/table*

### 19.250.2 Examples

```
pair_style sw/angle/table
pair_coeff * * spce.sw type
```

Used in example input script:

```
examples/PACKAGES/manybody_table/in.spce_sw
```

### 19.250.3 Description

New in version 2Jun2022.

The *sw/angle/table* style is a modification of the original *pair\_style sw*. It has been developed for coarse-grained simulations (of water) (*Scherer1*), but can be employed for all kinds of systems. It computes a modified 3-body *Stillinger-Weber* potential for the energy  $E$  of a system of atoms as

$$E = \sum_i \sum_{j>i} \phi_2(r_{ij}) + \sum_i \sum_{j \neq i} \sum_{k>j} \phi_3(r_{ij}, r_{ik}, \theta_{ijk})$$

$$\phi_2(r_{ij}) = A_{ij} \epsilon_{ij} \left[ B_{ij} \left( \frac{\sigma_{ij}}{r_{ij}} \right)^{p_{ij}} - \left( \frac{\sigma_{ij}}{r_{ij}} \right)^{q_{ij}} \right] \exp \left( \frac{\sigma_{ij}}{r_{ij} - a_{ij} \sigma_{ij}} \right)$$

$$\phi_3(r_{ij}, r_{ik}, \theta_{ijk}) = f^{3b}(\theta_{ijk}) \exp \left( \frac{\gamma_{ij} \sigma_{ij}}{r_{ij} - a_{ij} \sigma_{ij}} \right) \exp \left( \frac{\gamma_{ik} \sigma_{ik}}{r_{ik} - a_{ik} \sigma_{ik}} \right)$$

where  $\phi_2$  is a two-body term and  $\phi_3$  is a three-body term. The summations in the formula are over all neighbors  $J$  and  $K$  of atom  $I$  within a cutoff distance  $a\sigma$ . In contrast to the original *sw* style, *sw/angle/table* allows for a flexible three-body term  $f^{3b}(\theta_{ijk})$  which is read in as a tabulated interaction. It can be parameterized with the *csg\_fmacth* app of VOTCA as available at: <https://gitlab.mpcdf.mpg.de/votca/votca>.

Only a single *pair\_coeff* command is used with the *sw/angle/table* style which specifies a modified Stillinger-Weber potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying  $N_{el}$  additional arguments after the “.sw” filename in the *pair\_coeff* command, where  $N_{el}$  is the number of LAMMPS atom types:

- “.sw” filename
- $N_{el}$  element names = mapping of SW elements to atom types

See the *pair\_coeff* page for alternate ways to specify the path for the potential file.

As an example, imagine a file *SiC.sw* has Stillinger-Weber values for Si and C. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Si, and the fourth to be C, you would use the following *pair\_coeff* command:

```
pair_coeff * * SiC.sw Si Si Si C
```

The first 2 arguments must be *\*\** so as to span all LAMMPS atom types. The first three Si arguments map LAMMPS atom types 1,2,3 to the Si element in the SW file. The final C argument maps LAMMPS atom type 4 to the C element in the SW file. If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *sw/angle/table* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

The (modified) Stillinger-Weber files have a “.sw” suffix. Lines that are not blank or comments (starting with #) define parameters for a triplet of elements. The parameters in a single entry correspond to the two-body and three-body coefficients in the formula above. Here, also the suffix “.sw” is used though the original Stillinger-Weber file format is supplemented with four additional lines per parameter block to specify the tabulated three-body interaction. A single entry then contains:

- element 1 (the center atom in a 3-body interaction)
- element 2
- element 3
- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- $a$
- $\lambda$

- $\gamma$
- $\cos \theta_0$
- A
- B
- p
- q
- tol
- filename
- keyword
- style
- N

The A, B, p, and q parameters are used only for two-body interactions. The  $\lambda$  and  $\cos \theta_0$  parameters, only used for three-body interactions in the original Stillinger-Weber style, are read in but ignored in this modified pair style. The  $\epsilon$  parameter is only used for two-body interactions in this modified pair style and not for the three-body terms. The  $\sigma$  and  $a$  parameters are used for both two-body and three-body interactions.  $\gamma$  is used only in the three-body interactions, but is defined for pairs of atoms. The non-annotated parameters are unitless.

LAMMPS introduces an additional performance-optimization parameter *tol* that is used for both two-body and three-body interactions. In the Stillinger-Weber potential, the interaction energies become negligibly small at atomic separations substantially less than the theoretical cutoff distances. LAMMPS therefore defines a virtual cutoff distance based on a user defined tolerance *tol*. The use of the virtual cutoff distance in constructing atom neighbor lists can significantly reduce the neighbor list sizes and therefore the computational cost. LAMMPS provides a *tol* value for each of the three-body entries so that they can be separately controlled. If *tol* = 0.0, then the standard Stillinger-Weber cutoff is used.

The additional parameters *filename*, *keyword*, *style*, and *N* refer to the tabulated angular potential  $f^{3b}(\theta_{ijk})$ . The tabulated angular potential has to be of the format as used in the *angle\_style table* command:

An interpolation tables of length *N* is created. The interpolation is done in one of 2 *styles*: *linear* or *spline*. For the *linear* style, the angle is used to find 2 surrounding table values from which an energy or its derivative is computed by linear interpolation. For the *spline* style, a cubic spline coefficients are computed and stored at each of the *N* values in the table. The angle is used to find the appropriate set of coefficients which are used to evaluate a cubic polynomial which computes the energy or derivative.

The *filename* specifies the file containing the tabulated energy and derivative values of  $f^{3b}(\theta_{ijk})$ . The *keyword* then specifies a section of the file. The format of this file is as follows (without the parenthesized comments):

```
# Angle potential for harmonic (one or more comment or blank lines)

HAM                               (keyword is the first text on line)
N 181 FP 0 0 EQ 90.0             (N, FP, EQ parameters)
                                   (blank line)
1 0.0 200.5 2.5                  (index, angle, energy, derivative)
2 1.0 198.0 2.5
...
181 180.0 0.0 0.0
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The next line lists (in any order) one or more parameters for the table. Each parameter is a keyword followed by one or more numeric values.

The parameter “N” is required and its value is the number of table entries that follow. Note that this may be different than the  $N$  specified in the Stillinger-Weber potential file. Let  $N_{sw} = N$  in the “.sw” file, and  $N_{file} = “N”$  in the tabulated angular file. What LAMMPS does is a preliminary interpolation by creating splines using the  $N_{file}$  tabulated values as nodal points. It uses these to interpolate as needed to generate energy and derivative values at  $N_{table}$  different points. The resulting tables of length  $N_{sw}$  are then used as described above, when computing energy and force for individual angles and their atoms. This means that if you want the interpolation tables of length  $N_{sw}$  to match exactly what is in the tabulated file (with effectively no preliminary interpolation), you should set  $N_{sw} = N_{file}$ .

The “FP” parameter is optional. If used, it is followed by two values  $f_{plo}$  and  $f_{phi}$ , which are the second derivatives at the innermost and outermost angle settings. These values are needed by the spline construction routines. If not specified by the “FP” parameter, they are estimated (less accurately) by the first two and last two derivative values in the table.

The “EQ” parameter is also optional. If used, it is followed by a the equilibrium angle value, which is used, for example, by the *fix shake* command. If not used, the equilibrium angle is set to 180.0.

Following a blank line, the next  $N$  lines of the angular table file list the tabulated values. On each line, the first value is the index from 1 to  $N$ , the second value is the angle value (in degrees), the third value is the energy (in energy units), and the fourth is  $-dE/d(\theta)$  (also in energy units). The third term is the energy of the 3-atom configuration for the specified angle. The last term is the derivative of the energy with respect to the angle (in degrees, not radians). Thus the units of the last term are still energy, not force. The angle values must increase from one line to the next. The angle values must also begin with 0.0 and end with 180.0, i.e. span the full range of possible angles.

Note that one angular potential file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified *keyword* of appropriate section of the “.sw” file.

The Stillinger-Weber potential file must contain entries for all the elements listed in the `pair_coeff` command. It can also contain entries for additional elements not being used in a particular simulation; LAMMPS ignores those entries.

For a single-element simulation, only a single entry is required (e.g. SiSiSi). For a two-element simulation, the file must contain 8 entries (for SiSiSi, SiSiC, SiCSi, SiCC, CSiSi, CSiC, CCSi, CCC), that specify SW parameters for all permutations of the two elements interacting in three-body configurations. Thus for 3 elements, 27 entries would be required, etc.

As annotated above, the first element in the entry is the center atom in a three-body interaction. Thus an entry for SiCC means a Si atom with 2 C atoms as neighbors. The parameter values used for the two-body interaction come from the entry where the second and third elements are the same. Thus the two-body parameters for Si interacting with C, comes from the SiCC entry. The three-body angular potential  $f^{3b}(\theta_{ijk})$  can in principle be specific to the three elements of the configuration. However, the user must ensure that it makes physically sense. Note also that the function  $\phi_3$  contains two exponential screening factors with parameter values from the  $ij$  pair and  $ik$  pairs. So  $\phi_3$  for a C atom bonded to a Si atom and a second C atom will depend on the three-body parameters for the CSiC entry, and also on the two-body parameters for the CCC and CSiSi entries. Since the order of the two neighbors is arbitrary, the three-body parameters and the tabulated angular potential for entries CSiC and CCSi should be the same. Similarly, the two-body parameters for entries SiCC and CSiSi should also be the same. The parameters used only for two-body interactions ( $A$ ,  $B$ ,  $p$ , and  $q$ ) in entries whose second and third element are different (e.g. SiCSi) are not used for anything and can be set to 0.0 if desired. This is also true for the parameters in  $\phi_3$  that are taken from the  $ij$  and  $ik$  pairs ( $\sigma$ ,  $a$ ,  $\gamma$ )

Additional input files and reference data can be found at: [https://gitlab.mpcdf.mpg.de/votca/votca/-/tree/master/csg-tutorials/spce/3body\\_sw](https://gitlab.mpcdf.mpg.de/votca/votca/-/tree/master/csg-tutorials/spce/3body_sw)

---

### 19.250.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, mixing is performed by LAMMPS as described above from values in the potential file, but not for the tabulated angular potential file.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.250.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

### 19.250.6 Related commands

*pair\_coeff*, *pair\_style sw*, *pair\_style threebody/table*

---

(Stillinger) Stillinger and Weber, Phys Rev B, 31, 5262 (1985).

(Scherer1) C. Scherer and D. Andrienko, Phys. Chem. Chem. Phys. 20, 22387-22394 (2018).

## 19.251 pair\_style table command

Accelerator Variants: *table/gpu*, *table/kk*, *table/omp*

### 19.251.1 Syntax

```
pair_style table style N keyword ...
```

- style = *lookup* or *linear* or *spline* or *bitmap* = method of interpolation
- N = use N values in *lookup*, *linear*, *spline* tables
- N = use 2^N values in *bitmap* tables
- zero or more keywords may be appended
- keyword = *ewald* or *pppm* or *msm* or *dispersion* or *tip4p*

## 19.251.2 Examples

```
pair_style table linear 1000
pair_style table linear 1000 ppm
pair_style table bitmap 12
pair_coeff * 3 morse.table ENTRY1
pair_coeff * 3 morse.table ENTRY1 7.0
```

## 19.251.3 Description

Style *table* creates interpolation tables from potential energy and force values listed in a file(s) as a function of distance. When performing dynamics or minimization, the interpolation tables are used to evaluate energy and forces for pairwise interactions between particles, similar to how analytic formulas are used for other pair styles.

The interpolation tables are created as a pre-computation by fitting cubic splines to the file values and interpolating energy and force values at each of  $N$  distances. During a simulation, the tables are used to interpolate energy and force values as needed for each pair of particles separated by a distance  $R$ . The interpolation is done in one of 4 styles: *lookup*, *linear*, *spline*, or *bitmap*.

For the *lookup* style, the distance  $R$  is used to find the nearest table entry, which is the energy or force.

For the *linear* style, the distance  $R$  is used to find the 2 surrounding table values from which an energy or force is computed by linear interpolation.

For the *spline* style, cubic spline coefficients are computed and stored for each of the  $N$  values in the table, one set of splines for energy, another for force. Note that these splines are different than the ones used to pre-compute the  $N$  values. Those splines were fit to the  $N_{file}$  values in the tabulated file, where often  $N_{file} < N$ . The distance  $R$  is used to find the appropriate set of spline coefficients which are used to evaluate a cubic polynomial which computes the energy or force.

For the *bitmap* style, the specified  $N$  is used to create interpolation tables that are  $2^N$  in length. The distance  $R$  is used to index into the table via a fast bit-mapping technique due to (Wolff), and a linear interpolation is performed between adjacent table values.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above.

- filename
- keyword
- cutoff (distance units)

The filename specifies a file containing tabulated energy and force values. The keyword specifies a section of the file. The cutoff is an optional coefficient. If not specified, the outer cutoff in the table itself (see below) will be used to build an interpolation table that extend to the largest tabulated distance. If specified, only file values up to the cutoff are used to create the interpolation table. The format of this file is described below.

If your tabulated potential(s) are designed to be used as the short-range part of one of the long-range solvers specified by the *kpace\_style* command, then you must use one or more of the optional keywords listed above for the *pair\_style* command. These are *ewald* or *pppm* or *msm* or *dispersion* or *tip4p*. This is so LAMMPS can insure the short-range potential and long-range solver are compatible with each other, as it does for other short-range pair styles, such as *pair\_style lj/cut/coul/long*. Note that it is up to you to insure the tabulated values for each pair of atom types has the correct functional form to be compatible with the matching long-range solver.

---

Here are some guidelines for using the *pair\_style* table command to best effect:



- Vary the number of table points; you may need to use more than you think to get good resolution.
- Always use the *pair\_write* command to produce a plot of what the final interpolated potential looks like. This can show up interpolation “features” you may not like.
- Start with the linear style; it’s the style least likely to have problems.
- Use  $N$  in the *pair\_style* command equal to the “ $N$ ” in the tabulation file, and use the “RSQ” or “BITMAP” parameter, so additional interpolation is not needed. See discussion below.
- Make sure that your tabulated forces and tabulated energies are consistent ( $dE/dr = -F$ ) over the entire range of  $r$  values. LAMMPS will warn if this is not the case.
- Use as large an inner cutoff as possible. This avoids fitting splines to very steep parts of the potential.

The format of a tabulated file has an (optional) header followed by a series of one or more sections, defined as follows (without the parenthesized comments). The header must start with a # character and the DATE: and UNITS: tags will be parsed and used:

```
# DATE: 2020-06-10 UNITS: real CONTRIBUTOR: ... (header line)
# Morse potential for Fe (one or more comment or blank lines)

MORSE_FE (keyword is first text on line)
N 500 R 1.0 10.0 (N, R, RSQ, BITMAP, FPRIME parameters)
                (blank)
1 1.0 25.5 102.34 (index, r, energy, force)
2 1.02 23.4 98.5
...
500 10.0 0.001 0.003
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The line can contain additional text, but the initial text must match the argument specified in the *pair\_coeff* command. The next line lists (in any order) one or more parameters for the table. Each parameter is a keyword followed by one or more numeric values.

The parameter “ $N$ ” is required and its value is the number of table entries that follow. Note that this may be different than the  $N$  specified in the *pair\_style table* command. Let  $N_{\text{table}} = N$  in the *pair\_style* command, and  $N_{\text{file}} = “N”$  in the tabulated file. What LAMMPS does is a preliminary interpolation by creating splines using the  $N_{\text{file}}$  tabulated values as nodal points. It uses these to interpolate energy and force values at  $N_{\text{table}}$  different points. The resulting tables of length  $N_{\text{table}}$  are then used as described above, when computing energy and force for individual pair distances. This means that if you want the interpolation tables of length  $N_{\text{table}}$  to match exactly what is in the tabulated file (with effectively no preliminary interpolation), you should set  $N_{\text{table}} = N_{\text{file}}$ , and use the “RSQ” or “BITMAP” parameter. This is because the internal table abscissa is always RSQ (separation distance squared), for efficient lookup.

All other parameters are optional. If “R” or “RSQ” or “BITMAP” does not appear, then the distances in each line of the table are used as-is to perform spline interpolation. In this case, the table values can be spaced in  $r$  uniformly or however you wish to position table values in regions of large gradients.

If used, the parameters “R” or “RSQ” are followed by 2 values *rlo* and *rhi*. If specified, the distance associated with each energy and force value is computed from these 2 values (at high accuracy), rather than using the (low-accuracy) value listed in each line of the table. The distance values in the table file are ignored in this case. For “R”, distances uniformly spaced between *rlo* and *rhi* are computed; for “RSQ”, squared distances uniformly spaced between *rlo*\**rlo* and *rhi*\**rhi* are computed.

**Note:** If you use “R” or “RSQ”, the tabulated distance values in the file are effectively ignored, and replaced by new values as described in the previous paragraph. If the distance value in the table is not very close to the new value (i.e.

round-off difference), then you will be assigning energy/force values to a different distance, which is probably not what you want. LAMMPS will warn if this is occurring.

---

If used, the parameter “BITMAP” is also followed by 2 values *rlo* and *rhi*. These values, along with the “N” value determine the ordering of the N lines that follow and what distance is associated with each. This ordering is complex, so it is not documented here, since this file is typically produced by the *pair\_write* command with its *bitmap* option. When the table is in BITMAP format, the “N” parameter in the file must be equal to  $2^M$  where M is the value specified in the *pair\_style* command. Also, a cutoff parameter cannot be used as an optional third argument in the *pair\_coeff* command; the entire table extent as specified in the file must be used.

If used, the parameter “FPRIME” is followed by 2 values *fplo* and *fphi* which are the derivative of the force at the innermost and outermost distances listed in the table. These values are needed by the spline construction routines. If not specified by the “FPRIME” parameter, they are estimated (less accurately) by the first 2 and last 2 force values in the table. This parameter is not used by BITMAP tables.

Following a blank line, the next N lines list the tabulated values. On each line, the first value is the index from 1 to N, the second value is r (in distance units), the third value is the energy (in energy units), and the fourth is the force (in force units). The r values must increase from one line to the next (unless the BITMAP parameter is specified).

Note that one file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified keyword.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.251.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

The *pair\_modify* shift, table, and tail options are not relevant for this pair style.

This pair style writes the settings for the “pair\_style table” command to *binary restart files*, so a pair\_style command does not need to be specified in an input script that reads a restart file. However, the coefficient information is not stored in the restart file, since it is tabulated in the potential files. Thus, pair\_coeff commands do need to be specified in the restart input script.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.251.5 Restrictions

none

### 19.251.6 Related commands

*pair\_coeff*, *pair\_write*

### 19.251.7 Default

none

---

(Wolff) Wolff and Rudd, Comp Phys Comm, 120, 200-32 (1999).

## 19.252 *pair\_style table/rx* command

Accelerator Variants: *table/rx/kk*

### 19.252.1 Syntax

```
pair_style table style N ...
```

- style = *lookup* or *linear* or *spline* or *bitmap* = method of interpolation
- N = use N values in *lookup*, *linear*, *spline* tables
- weighting = fractional or molecular (optional)

### 19.252.2 Examples

```
pair_style table/rx linear 1000
pair_style table/rx linear 1000 fractional
pair_style table/rx linear 1000 molecular
pair_coeff * * rxn.table ENTRY1 h2o h2o 10.0
pair_coeff * * rxn.table ENTRY1 1fluid 1fluid 10.0
pair_coeff * 3 rxn.table ENTRY1 h2o no2 10.0
```

### 19.252.3 Description

Style *table/rx* is used in reaction DPD simulations, where the coarse-grained (CG) particles are composed of  $m$  species whose reaction rate kinetics are determined from a set of  $n$  reaction rate equations through the *fix rx* command. The species of one CG particle can interact with a species in a neighboring CG particle through a site-site interaction potential model. Style *table/rx* creates interpolation tables of length  $N$  from pair potential and force values listed in a file(s) as a function of distance. The files are read by the *pair\_coeff* command.

The interpolation tables are created by fitting cubic splines to the file values and interpolating energy and force values at each of  $N$  distances. During a simulation, these tables are used to interpolate energy and force values as needed. The interpolation is done in one of 4 styles: *lookup*, *linear*, *spline*, or *bitmap*.

For the *lookup* style, the distance between 2 atoms is used to find the nearest table entry, which is the energy or force.

For the *linear* style, the pair distance is used to find 2 surrounding table values from which an energy or force is computed by linear interpolation.

For the *spline* style, a cubic spline coefficients are computed and stored at each of the  $N$  values in the table. The pair distance is used to find the appropriate set of coefficients which are used to evaluate a cubic polynomial which computes the energy or force.

For the *bitmap* style, the  $N$  means to create interpolation tables that are  $2^N$  in length. The pair distance is used to index into the table via a fast bit-mapping technique (*Wolff*) and a linear interpolation is performed between adjacent table values.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above.

- filename
- keyword
- species1
- species2
- cutoff (distance units)

The filename specifies a file containing tabulated energy and force values. The keyword specifies a section of the file. The cutoff is an optional coefficient. If not specified, the outer cutoff in the table itself (see below) will be used to build an interpolation table that extend to the largest tabulated distance. If specified, only file values up to the cutoff are used to create the interpolation table. The format of this file is described below.

The species tags define the site-site interaction potential between two species contained within two different particles. The species tags must either correspond to the species defined in the reaction kinetics files specified with the *fix rx* command or they must correspond to the tag “1fluid”, signifying interaction with a product species mixture determined through a one-fluid approximation. The interaction potential is weighted by the geometric average of either the mole fraction concentrations or the number of molecules associated with the interacting coarse-grained particles (see the *fractional* or *molecular* weighting pair style options). The coarse-grained potential is stored before and after the reaction kinetics solver is applied, where the difference is defined to be the internal chemical energy (uChem).

---

Here are some guidelines for using the *pair\_style table/rx* command to best effect:

- Vary the number of table points; you may need to use more than you think to get good resolution.
  - Always use the *pair\_write* command to produce a plot of what the final interpolated potential looks like. This can show up interpolation “features” you may not like.
  - Start with the linear style; it’s the style least likely to have problems.
  - Use  $N$  in the *pair\_style* command equal to the “ $N$ ” in the tabulation file, and use the “RSQ” or “BITMAP” parameter, so additional interpolation is not needed. See discussion below.
  - Make sure that your tabulated forces and tabulated energies are consistent ( $dE/dr = -F$ ) along the entire range of  $r$  values.
  - Use as large an inner cutoff as possible. This avoids fitting splines to very steep parts of the potential.
- 

The format of a tabulated file is a series of one or more sections, defined as follows (without the parenthesized comments):

```

# Morse potential for Fe   (one or more comment or blank lines)

MORSE_FE                 (keyword is first text on line)
N 500 R 1.0 10.0         (N, R, RSQ, BITMAP, FPRIME parameters)
                          (blank)
1 1.0 25.5 102.34       (index, r, energy, force)
2 1.02 23.4 98.5
...
500 10.0 0.001 0.003

```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The line can contain additional text, but the initial text must match the argument specified in the `pair_coeff` command. The next line lists (in any order) one or more parameters for the table. Each parameter is a keyword followed by one or more numeric values.

The parameter “N” is required and its value is the number of table entries that follow. Note that this may be different than the  $N$  specified in the `pair_style table/rx` command. Let  $N_{\text{table}} = N$  in the `pair_style` command, and  $N_{\text{file}} = “N”$  in the tabulated file. What LAMMPS does is a preliminary interpolation by creating splines using the  $N_{\text{file}}$  tabulated values as nodal points. It uses these to interpolate as needed to generate energy and force values at  $N_{\text{table}}$  different points. The resulting tables of length  $N_{\text{table}}$  are then used as described above, when computing energy and force for individual pair distances. This means that if you want the interpolation tables of length  $N_{\text{table}}$  to match exactly what is in the tabulated file (with effectively no preliminary interpolation), you should set  $N_{\text{table}} = N_{\text{file}}$ , and use the “RSQ” or “BITMAP” parameter. The internal table abscissa is RSQ (separation distance squared).

All other parameters are optional. If “R” or “RSQ” or “BITMAP” does not appear, then the distances in each line of the table are used as-is to perform spline interpolation. In this case, the table values can be spaced in  $r$  uniformly or however you wish to position table values in regions of large gradients.

If used, the parameters “R” or “RSQ” are followed by 2 values  $r_{lo}$  and  $r_{hi}$ . If specified, the distance associated with each energy and force value is computed from these 2 values (at high accuracy), rather than using the (low-accuracy) value listed in each line of the table. The distance values in the table file are ignored in this case. For “R”, distances uniformly spaced between  $r_{lo}$  and  $r_{hi}$  are computed; for “RSQ”, squared distances uniformly spaced between  $r_{lo} * r_{lo}$  and  $r_{hi} * r_{hi}$  are computed.

If used, the parameter “BITMAP” is also followed by 2 values  $r_{lo}$  and  $r_{hi}$ . These values, along with the “N” value determine the ordering of the  $N$  lines that follow and what distance is associated with each. This ordering is complex, so it is not documented here, since this file is typically produced by the `pair_write` command with its `bitmap` option. When the table is in BITMAP format, the “N” parameter in the file must be equal to  $2^M$  where  $M$  is the value specified in the `pair_style` command. Also, a cutoff parameter cannot be used as an optional third argument in the `pair_coeff` command; the entire table extent as specified in the file must be used.

If used, the parameter “FPRIME” is followed by 2 values  $f_{plo}$  and  $f_{phi}$  which are the derivative of the force at the innermost and outermost distances listed in the table. These values are needed by the spline construction routines. If not specified by the “FPRIME” parameter, they are estimated (less accurately) by the first 2 and last 2 force values in the table. This parameter is not used by BITMAP tables.

Following a blank line, the next  $N$  lines list the tabulated values. On each line, the first value is the index from 1 to  $N$ , the second value is  $r$  (in distance units), the third value is the energy (in energy units), and the fourth is the force (in force units). The  $r$  values must increase from one line to the next (unless the BITMAP parameter is specified).

Note that one file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified keyword.

### 19.252.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

The *pair\_modify* shift, table, and tail options are not relevant for this pair style.

This pair style writes the settings for the “pair\_style table/rx” command to *binary restart files*, so a pair\_style command does not need to be specified in an input script that reads a restart file. However, the coefficient information is not stored in the restart file, since it is tabulated in the potential files. Thus, pair\_coeff commands do need to be specified in the restart input script.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 19.252.5 Restrictions

This command is part of the DPD-REACT package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.252.6 Related commands

*pair\_coeff*

### 19.252.7 Default

fractional weighting

---

(Wolff) Wolff and Rudd, Comp Phys Comm, 120, 200-32 (1999).

## 19.253 pair\_style tersoff command

Accelerator Variants: *tersoff/gpu*, *tersoff/intel*, *tersoff/kk*, *tersoff/omp*

## 19.254 pair\_style tersoff/table command

Accelerator Variants: *tersoff/table/omp*

### 19.254.1 Syntax

```
pair_style style keywords values
```

- style = *tersoff* or *tersoff/table*
- keyword = *shift*
  - shift* value = delta
  - delta = negative shift in equilibrium bond length

### 19.254.2 Examples

```
pair_style tersoff
pair_coeff * * Si.tersoff Si
pair_coeff * * SiC.tersoff Si C Si

pair_style tersoff/table
pair_coeff * * SiGe.tersoff Si(D)

pair_style tersoff shift 0.05
pair_coeff * * Si.tersoff Si
```

### 19.254.3 Description

The *tersoff* style computes a 3-body Tersoff potential (*Tersoff\_1*) for the energy  $E$  of a system of atoms as

$$\begin{aligned}
 E &= \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij} \\
 V_{ij} &= f_C(r_{ij} + \delta) [f_R(r_{ij} + \delta) + b_{ij} f_A(r_{ij} + \delta)] \\
 f_C(r) &= \begin{cases} 1 & : r < R - D \\ \frac{1}{2} - \frac{1}{2} \sin\left(\frac{\pi}{2} \frac{r-R}{D}\right) & : R - D < r < R + D \\ 0 & : r > R + D \end{cases} \\
 f_R(r) &= A \exp(-\lambda_1 r) \\
 f_A(r) &= -B \exp(-\lambda_2 r) \\
 b_{ij} &= (1 + \beta^n \zeta_{ij}^n)^{-\frac{1}{2n}} \\
 \zeta_{ij} &= \sum_{k \neq i, j} f_C(r_{ik} + \delta) g[\theta_{ijk}(r_{ij}, r_{ik})] \exp[\lambda_3^m (r_{ij} - r_{ik})^m] \\
 g(\theta) &= \gamma_{ijk} \left( 1 + \frac{c^2}{d^2} - \frac{c^2}{[d^2 + (\cos \theta - \cos \theta_0)^2]} \right)
 \end{aligned}$$

where  $f_R$  is a two-body term and  $f_A$  includes three-body interactions. The summations in the formula are over all neighbors  $J$  and  $K$  of atom  $I$  within a cutoff distance =  $R + D$ .  $\delta$  is an optional negative shift of the equilibrium bond length, as described below.

The *tersoff/table* style uses tabulated forms for the two-body, environment and angular functions. Linear interpolation is performed between adjacent table entries. The table length is chosen to be accurate within  $10^{-6}$  with respect to the *tersoff* style energy. The *tersoff/table* should give better performance in terms of speed.

Only a single `pair_coeff` command is used with the *tersoff* style which specifies a Tersoff potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying  $N$  additional arguments after the filename in the `pair_coeff` command, where  $N$  is the number of LAMMPS atom types:

- filename
- $N$  element names = mapping of Tersoff elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the potential file.

As an example, imagine the `SiC.tersoff` file has Tersoff values for Si and C. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Si, and the fourth to be C, you would use the following `pair_coeff` command:

```
pair_coeff * * SiC.tersoff Si Si Si C
```

The first 2 arguments must be `**` so as to span all LAMMPS atom types. The first three Si arguments map LAMMPS atom types 1,2,3 to the Si element in the Tersoff file. The final C argument maps LAMMPS atom type 4 to the C element in the Tersoff file. If a mapping value is specified as `NULL`, the mapping is not performed. This can be used when a *tersoff* potential is used as part of the *hybrid* pair style. The `NULL` values are placeholders for atom types that will be used with other potentials.

Tersoff files in the *potentials* directory of the LAMMPS distribution have a “.tersoff” suffix. Lines that are not blank or comments (starting with #) define parameters for a triplet of elements. The parameters in a single entry correspond to coefficients in the formula above:

- element 1 (the center atom in a 3-body interaction)
- element 2 (the atom bonded to the center atom)
- element 3 (the atom influencing the 1-2 bond in a bond-order sense)



- m
- $\gamma$
- $\lambda_3$  (1/distance units)
- c
- d
- $\cos \theta_0$  (can be a value < -1 or > 1)
- n
- $\beta$
- $\lambda_2$  (1/distance units)
- B (energy units)
- R (distance units)
- D (distance units)
- $\lambda_1$  (1/distance units)
- A (energy units)

The n,  $\beta$ ,  $\lambda_2$ , B,  $\lambda_1$ , and A parameters are only used for two-body interactions. The m,  $\gamma$ ,  $\lambda_3$ , c, d, and  $\cos \theta_0$  parameters are only used for three-body interactions. The R and D parameters are used for both two-body and three-body interactions. The non-annotated parameters are unitless. The value of m must be 3 or 1.

The Tersoff potential file must contain entries for all the elements listed in the pair\_coeff command. It can also contain entries for additional elements not being used in a particular simulation; LAMMPS ignores those entries.

For a single-element simulation, only a single entry is required (e.g. SiSiSi). For a two-element simulation, the file must contain 8 entries (for SiSiSi, SiSiC, SiCSi, SiCC, CSiSi, CSiC, CCSi, CCC), that specify Tersoff parameters for all permutations of the two elements interacting in three-body configurations. Thus for 3 elements, 27 entries would be required, etc.

As annotated above, the first element in the entry is the center atom in a three-body interaction and it is bonded to the second atom and the bond is influenced by the third atom. Thus an entry for SiCC means Si bonded to a C with another C atom influencing the bond. Thus three-body parameters for SiCSi and SiSiC entries will not, in general, be the same. The parameters used for the two-body interaction come from the entry where the second element is repeated. Thus the two-body parameters for Si interacting with C, comes from the SiCC entry.

The parameters used for a particular three-body interaction come from the entry with the corresponding three elements. The parameters used only for two-body interactions (n,  $\beta$ ,  $\lambda_2$ , B,  $\lambda_1$ , and A) in entries whose second and third element are different (e.g. SiCSi) are not used for anything and can be set to 0.0 if desired.

Note that the twobody parameters in entries such as SiCC and CSiSi are often the same, due to the common use of symmetric mixing rules, but this is not always the case. For example, the beta and n parameters in Tersoff\_2 (*Tersoff\_2*) are not symmetric. Similarly, the threebody parameters in entries such as SiCSi and SiSiC are often the same, but this is not always the case, particularly the value of R, which is sometimes typed on the first and second elements, sometimes on the first and third elements. Hence the need to specify R and D explicitly for all element triples. For example, while Tersoff's notation in Tersoff\_2 (*Tersoff\_2*) is ambiguous on this point, and properties of the zincblende lattice are the same for either choice, Tersoff's results for rocksalt are consistent with typing on the first and third elements. *Albe et al.* adopts the same convention. Conversely, the potential for B/N/C from the Cagin group uses the opposite convention, typing on the first and second elements.

We chose the above form so as to enable users to define all commonly used variants of the Tersoff potential. In particular, our form reduces to the original Tersoff form when  $m = 3$  and  $\gamma = 1$ , while it reduces to the form of *Albe et al.* when  $\beta = 1$  and  $m = 1$ . Note that in the current Tersoff implementation in LAMMPS, m must be specified as either

3 or 1. Tersoff used a slightly different but equivalent form for alloys, which we will refer to as Tersoff\_2 potential (*Tersoff\_2*). The *tersoff/table* style implements Tersoff\_2 parameterization only.

LAMMPS parameter values for Tersoff\_2 can be obtained as follows:  $\gamma_{ijk} = \omega_{ik}$ ,  $\lambda_3 = 0$  and the value of m has no effect. The parameters for species i and j can be calculated using the Tersoff\_2 mixing rules:

$$\begin{aligned}\lambda_1^{i,j} &= \frac{1}{2}(\lambda_1^i + \lambda_1^j) \\ \lambda_2^{i,j} &= \frac{1}{2}(\lambda_2^i + \lambda_2^j) \\ A_{i,j} &= (A_i A_j)^{1/2} \\ B_{i,j} &= \chi_{ij} (B_i B_j)^{1/2} \\ R_{i,j} &= (R_i R_j)^{1/2} \\ S_{i,j} &= (S_i S_j)^{1/2}\end{aligned}$$

Tersoff\_2 parameters R and S must be converted to the LAMMPS parameters R and D (R is different in both forms), using the following relations:  $R=(R'+S')/2$  and  $D=(S'-R')/2$ , where the primes indicate the Tersoff\_2 parameters.

In the potentials directory, the file `SiCGe.tersoff` provides the LAMMPS parameters for Tersoff's various versions of Si, as well as his alloy parameters for Si, C, and Ge. This file can be used for pure Si, (three different versions), pure C, pure Ge, binary SiC, and binary SiGe. LAMMPS will generate an error if this file is used with any combination involving C and Ge, since there are no entries for the GeC interactions (Tersoff did not publish parameters for this cross-interaction.) Tersoff files are also provided for the SiC alloy (`SiC.tersoff`) and the GaN (`GaN.tersoff`) alloys.

Many thanks to Rutuparna Narulkar, David Farrell, and Xiaowang Zhou for helping clarify how Tersoff parameters for alloys have been defined in various papers.

The *shift* keyword computes the energy E of a system of atoms, whose formula is the same as the Tersoff potential. The only modification is that the original equilibrium bond length ( $r_0$ ) of the system is shifted to  $r_0 - \delta$ . The minus sign arises because each radial distance  $r$  is replaced by  $r + \delta$ .

The *shift* keyword is designed for simulations of closely matched van der Waals heterostructures. For instance, consider the case of a system with few-layers graphene atop a thick hexagonal boron nitride (h-BN) substrate simulated using periodic boundary conditions. The experimental lattice mismatch of ~1.8% between graphene and h-BN is not well captured by the equilibrium lattice constants of available potentials, thus a small in-plane strain will be introduced in the system when building a periodic supercell. To minimize the effect of strain on simulation results, the *shift* keyword allows adjusting the equilibrium bond length of one of the two materials (e.g., h-BN). Validation, benchmark tests, and applications of the *shift* keyword can be found in ([Mandelli\\_1](#)) and ([Ouyang\\_1](#)).

For the specific case discussed above, the force field can be defined as

```
pair_style hybrid/overlay rebo tersoff shift -0.00407 ilp/graphene/hbn 16.0 coul/shield_
↪16.0
pair_coeff * * rebo CH.rebo NULL NULL C
pair_coeff * * tersoff BNC.tersoff B N NULL
pair_coeff * * ilp/graphene/hbn BNCH.ILP B N C
pair_coeff 1 1 coul/shield 0.70
pair_coeff 1 2 coul/shield 0.695
pair_coeff 2 2 coul/shield 0.69
```

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.254.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, mixing is performed by LAMMPS as described above from values in the potential file.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.254.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the [Build package](#) page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

The *shift* keyword is not supported by the *tersoff/gpu*, *tersoff/intel*, *tersoff/kk*, *tersoff/table* or *tersoff/table/omp* variants.

The *tersoff/intel* pair style is only available when compiling LAMMPS with the Intel compilers.

The Tersoff potential files provided with LAMMPS (see the potentials directory) are parameterized for “*metal*” units. In addition the pair style supports converting potential parameters on-the-fly between “metal” and “real” units. You can use the *tersoff* pair style variants with any LAMMPS units setting, but you would need to create your own Tersoff potential file with coefficients listed in the appropriate units if your simulation does not use “metal” or “real” units.

#### 19.254.6 Related commands

*pair\_coeff*

#### 19.254.7 Default

shift delta = 0.0

---

(**Tersoff\_1**) J. Tersoff, Phys Rev B, 37, 6991 (1988).

(**Albe**) J. Nord, K. Albe, P. Erhart, and K. Nordlund, J. Phys.: Condens. Matter, 15, 5649(2003).

(**Tersoff\_2**) J. Tersoff, Phys Rev B, 39, 5566 (1989); errata (PRB 41, 3248)

(**Mandelli\_1**) D. Mandelli, W. Ouyang, M. Urbakh, and O. Hod, ACS Nano 13(7), 7603-7609 (2019).

(**Ouyang\_1**) W. Ouyang et al., J. Chem. Theory Comput. 16(1), 666-676 (2020).

## 19.255 pair\_style tersoff/mod command

Accelerator Variants: *tersoff/mod/gpu*, *tersoff/mod/kk*, *tersoff/mod/omp*

## 19.256 pair\_style tersoff/mod/c command

Accelerator Variants: *tersoff/mod/c/omp*

### 19.256.1 Syntax

```
pair_style style keywords values
```

- style = *tersoff/mod* or *tersoff/mod/c*
- keyword = *shift*

*shift* value = delta

delta = negative shift in equilibrium bond length

### 19.256.2 Examples

```
pair_style tersoff/mod
pair_coeff * * Si.tersoff.mod Si Si

pair_style tersoff/mod/c
pair_coeff * * Si.tersoff.modc Si Si
```

### 19.256.3 Description

The *tersoff/mod* and *tersoff/mod/c* styles computes a bond-order type interatomic potential (*Kumagai*) based on a 3-body Tersoff potential (*Tersoff\_1*), (*Tersoff\_2*) with modified cutoff function and angular-dependent term, giving the energy E of a system of atoms as

$$\begin{aligned}
 E &= \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij} \\
 V_{ij} &= f_C(r_{ij} + \delta) [f_R(r_{ij} + \delta) + b_{ij} f_A(r_{ij} + \delta)] \\
 f_C(r) &= \begin{cases} 1 & : r < R - D \\ \frac{1}{2} - \frac{9}{16} \sin\left(\frac{\pi}{2} \frac{r-R}{D}\right) - \frac{1}{16} \sin\left(\frac{3\pi}{2} \frac{r-R}{D}\right) & : R - D < r < R + D \\ 0 & : r > R + D \end{cases} \\
 f_R(r) &= A \exp(-\lambda_1 r) \\
 f_A(r) &= -B \exp(-\lambda_2 r) \\
 b_{ij} &= (1 + \zeta_{ij}^\eta)^{-\frac{1}{2\eta}} \\
 \zeta_{ij} &= \sum_{k \neq i, j} f_C(r_{ik} + \delta) g(\theta_{ijk}) \exp[\alpha(r_{ij} - r_{ik})^\beta] \\
 g(\theta) &= c_1 + g_o(\theta) g_a(\theta) \\
 g_o(\theta) &= \frac{c_2 (h - \cos \theta)^2}{c_3 + (h - \cos \theta)^2} \\
 g_a(\theta) &= 1 + c_4 \exp[-c_5 (h - \cos \theta)^2]
 \end{aligned}$$

where  $f_R$  is a two-body term and  $f_A$  includes three-body interactions.  $\delta$  is an optional negative shift of the equilibrium bond length, as described below.

The summations in the formula are over all neighbors J and K of atom I within a cutoff distance = R + D. The *tersoff/mod/c* style differs from *tersoff/mod* only in the formulation of the  $V_{ij}$  term, where it contains an additional  $c_0$  term.

$$V_{ij} = f_C(r_{ij} + \delta) [f_R(r_{ij} + \delta) + b_{ij} f_A(r_{ij} + \delta) + c_0]$$

The modified cutoff function  $f_C$  proposed by (*Murty*) and having a continuous second-order differential is employed. The angular-dependent term  $g(\theta)$  was modified to increase the flexibility of the potential.

The *tersoff/mod* potential is fitted to both the elastic constants and melting point by employing the modified Tersoff potential function form in which the angular-dependent term is improved. The model performs extremely well in describing the crystalline, liquid, and amorphous phases (*Schelling*).

Only a single `pair_coeff` command is used with the *tersoff/mod* style which specifies a Tersoff/MOD potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of Tersoff/MOD elements to atom types

As an example, imagine the `Si.tersoff_mod` file has Tersoff values for Si. If your LAMMPS simulation has 3 Si atoms types, you would use the following `pair_coeff` command:

```
pair_coeff * * Si.tersoff_mod Si Si Si
```

The first 2 arguments must be `**` so as to span all LAMMPS atom types. The three Si arguments map LAMMPS atom types 1,2,3 to the Si element in the Tersoff/MOD file. If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *tersoff/mod* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

Tersoff/MOD file in the *potentials* directory of the LAMMPS distribution have a “.tersoff.mod” suffix. Potential files for the *tersoff/mod/c* style have the suffix “.tersoff.modc”. Lines that are not blank or comments (starting with #) define parameters for a triplet of elements. The parameters in a single entry correspond to coefficients in the formulae above:

- element 1 (the center atom in a 3-body interaction)
- element 2 (the atom bonded to the center atom)
- element 3 (the atom influencing the 1-2 bond in a bond-order sense)
- $\beta$
- $\alpha$
- h
- $\eta$
- $\beta_{ters} = 1$  (dummy parameter)
- $\lambda_2$  (1/distance units)
- B (energy units)
- R (distance units)
- D (distance units)
- $\lambda_1$  (1/distance units)
- A (energy units)
- n
- c1
- c2
- c3
- c4
- c5
- c0 (energy units, tersoff/mod/c only):ul

The n,  $\eta$ ,  $\lambda_2$ , B,  $\lambda_1$ , and A parameters are only used for two-body interactions. The  $\beta$ ,  $\alpha$ , c1, c2, c3, c4, c5, h parameters are only used for three-body interactions. The R and D parameters are used for both two-body and three-body interactions. The c0 term applies to *tersoff/mod/c* only. The non-annotated parameters are unitless.

The Tersoff/MOD potential file must contain entries for all the elements listed in the pair\_coeff command. It can also contain entries for additional elements not being used in a particular simulation; LAMMPS ignores those entries.

For a single-element simulation, only a single entry is required (e.g. SiSiSi). As annotated above, the first element in the entry is the center atom in a three-body interaction and it is bonded to the second atom and the bond is influenced by the third atom. Thus an entry for SiSiSi means Si bonded to a Si with another Si atom influencing the bond.

The *shift* keyword computes the energy E of a system of atoms, whose formula is the same as the Tersoff potential. The only modification is that the original equilibrium bond length ( $r_0$ ) of the system is shifted to  $r_0 - \delta$ . The minus sign arises because each radial distance  $r$  is replaced by  $r + \delta$ . More information on this option is given on the main [pair\\_tersoff](#) page.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#)

page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.256.4 Mixing, shift, table, tail correction, restart, rRESPA info

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.256.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

The *shift* keyword is not supported by the *tersoff/gpu*, *tersoff/intel*, *tersoff/kk*, *tersoff/table* or *tersoff/table/omp* variants.

The *tersoff/mod* potential files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use the *tersoff/mod* pair style with any LAMMPS units, but you would need to create your own Tersoff/MOD potential file with coefficients listed in the appropriate units if your simulation does not use “metal” units.

#### 19.256.6 Related commands

*pair\_coeff*

#### 19.256.7 Default

none

---

**(Kumagai)** T. Kumagai, S. Izumi, S. Hara, S. Sakai, *Comp. Mat. Science*, 39, 457 (2007).

**(Tersoff\_1)** J. Tersoff, *Phys Rev B*, 37, 6991 (1988).

**(Tersoff\_2)** J. Tersoff, *Phys Rev B*, 38, 9902 (1988).

**(Murty)** M.V.R. Murty, H.A. Atwater, *Phys Rev B*, 51, 4889 (1995).

**(Schelling)** Patrick K. Schelling, *Comp. Mat. Science*, 44, 274 (2008).

## 19.257 pair\_style tersoff/zbl command

Accelerator Variants: *tersoff/zbl/gpu*, *tersoff/zbl/kk*, *tersoff/zbl/omp*

### 19.257.1 Syntax

```
pair_style tersoff/zbl keywords values
```

- keyword = *shift*  
*shift* value = delta  
delta = negative shift in equilibrium bond length

### 19.257.2 Examples

```
pair_style tersoff/zbl  
pair_coeff * * SiC.tersoff.zbl Si C Si
```

### 19.257.3 Description

The *tersoff/zbl* style computes a 3-body Tersoff potential (*Tersoff\_1*) with a close-separation pairwise modification based on a Coulomb potential and the Ziegler-Biersack-Littmark universal screening function (*ZBL*), giving the energy



E of a system of atoms as

$$E = \frac{1}{2} \sum_i \sum_{j \neq i} V_{ij}$$

$$V_{ij} = (1 - f_F(r_{ij} + \delta)) V^{ZBL}(r_{ij} + \delta) + f_F(r_{ij} + \delta) V^{Tersoff}(r_{ij} + \delta)$$

$$f_F(r) = \frac{1}{1 + e^{-A_F(r-r_C)}}$$

$$V^{ZBL}(r) = \frac{1}{4\pi\epsilon_0} \frac{Z_1 Z_2 e^2}{r} \phi(r/a)$$

$$a = \frac{0.8854 a_0}{Z_1^{0.23} + Z_2^{0.23}}$$

$$\phi(x) = 0.1818e^{-3.2x} + 0.5099e^{-0.9423x} + 0.2802e^{-0.4029x} + 0.02817e^{-0.2016x}$$

$$V^{Tersoff}(r) = f_C(r) [f_R(r) + b_{ij} f_A(r)]$$

$$f_C(r) = \begin{cases} 1 & : r < R - D \\ \frac{1}{2} - \frac{1}{2} \sin\left(\frac{\pi}{2} \frac{r-R}{D}\right) & : R - D < r < R + D \\ 0 & : r > R + D \end{cases}$$

$$f_R(r) = A \exp(-\lambda_1 r)$$

$$f_A(r) = -B \exp(-\lambda_2 r)$$

$$b_{ij} = (1 + \beta^n \zeta_{ij}^n)^{-\frac{1}{2n}}$$

$$\zeta_{ij} = \sum_{k \neq i, j} f_C(r_{ik} + \delta) g(\theta_{ijk}) \exp[\lambda_3^m (r_{ij} - r_{ik})^m]$$

$$g(\theta) = \gamma_{ijk} \left( 1 + \frac{c^2}{d^2} - \frac{c^2}{[d^2 + (\cos \theta - \cos \theta_0)^2]} \right)$$

The  $f_F$  term is a fermi-like function used to smoothly connect the ZBL repulsive potential with the Tersoff potential. There are 2 parameters used to adjust it:  $A_F$  and  $r_C$ .  $A_F$  controls how “sharp” the transition is between the two, and  $r_C$  is essentially the cutoff for the ZBL potential.

For the ZBL portion, there are two terms. The first is the Coulomb repulsive term, with  $Z_1$ ,  $Z_2$  as the number of protons in each nucleus,  $e$  as the electron charge (1 for metal and real units) and  $\epsilon_0$  as the permittivity of vacuum. The second part is the ZBL universal screening function, with  $a_0$  being the Bohr radius (typically 0.529 Angstroms), and the remainder of the coefficients provided by the original paper. This screening function should be applicable to most systems. However, it is only accurate for small separations (i.e. less than 1 Angstrom).

For the Tersoff portion,  $f_R$  is a two-body term and  $f_A$  includes three-body interactions. The summations in the formula are over all neighbors  $J$  and  $K$  of atom  $I$  within a cutoff distance =  $R + D$ .

$\delta$  is an optional negative shift of the equilibrium bond length, as described below.

Only a single `pair_coeff` command is used with the `tersoff/zbl` style which specifies a Tersoff/ZBL potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying  $N$  additional arguments after the filename in the `pair_coeff` command, where  $N$  is the number of LAMMPS atom types:

- filename
- $N$  element names = mapping of Tersoff/ZBL elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the potential file.

As an example, imagine the SiC.tersoff.zbl file has Tersoff/ZBL values for Si and C. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Si, and the fourth to be C, you would use the following pair\_coeff command:

```
pair_coeff * * SiC.tersoff Si Si Si C
```

The first 2 arguments must be \* \* so as to span all LAMMPS atom types. The first three Si arguments map LAMMPS atom types 1,2,3 to the Si element in the Tersoff/ZBL file. The final C argument maps LAMMPS atom type 4 to the C element in the Tersoff/ZBL file. If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *tersoff/zbl* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

Tersoff/ZBL files in the *potentials* directory of the LAMMPS distribution have a “.tersoff.zbl” suffix. Lines that are not blank or comments (starting with #) define parameters for a triplet of elements. The parameters in a single entry correspond to coefficients in the formula above:

- element 1 (the center atom in a 3-body interaction)
- element 2 (the atom bonded to the center atom)
- element 3 (the atom influencing the 1-2 bond in a bond-order sense)
- m
- $\gamma$
- $\lambda_3$  (1/distance units)
- c
- d
- $\cos \theta_0$  (can be a value  $< -1$  or  $> 1$ )
- n
- $\beta$
- $\lambda_2$  (1/distance units)
- B (energy units)
- R (distance units)
- D (distance units)
- $\lambda_1$  (1/distance units)
- A (energy units)
- $Z_i$
- $Z_j$
- ZBLcut (distance units)
- ZBLexpscale (1/distance units)

The n,  $\beta$ ,  $\lambda_2$ , B,  $\lambda_1$ , and A parameters are only used for two-body interactions. The m,  $\gamma$ ,  $\lambda_3$ , c, d, and  $\cos \theta_0$  parameters are only used for three-body interactions. The R and D parameters are used for both two-body and three-body interactions. The  $Z_i$ ,  $Z_j$ , ZBLcut, ZBLexpscale parameters are used in the ZBL repulsive portion of the potential and in the Fermi-like function. The non-annotated parameters are unitless. The value of m must be 3 or 1.

The Tersoff/ZBL potential file must contain entries for all the elements listed in the pair\_coeff command. It can also contain entries for additional elements not being used in a particular simulation; LAMMPS ignores those entries.

For a single-element simulation, only a single entry is required (e.g. SiSiSi). For a two-element simulation, the file must contain 8 entries (for SiSiSi, SiSiC, SiCSi, SiCC, CSiSi, CSiC, CCSi, CCC), that specify Tersoff parameters for

all permutations of the two elements interacting in three-body configurations. Thus for 3 elements, 27 entries would be required, etc.

As annotated above, the first element in the entry is the center atom in a three-body interaction and it is bonded to the second atom and the bond is influenced by the third atom. Thus an entry for SiCC means Si bonded to a C with another C atom influencing the bond. Thus three-body parameters for SiCSi and SiSiC entries will not, in general, be the same. The parameters used for the two-body interaction come from the entry where the second element is repeated. Thus the two-body parameters for Si interacting with C, comes from the SiCC entry.

The parameters used for a particular three-body interaction come from the entry with the corresponding three elements. The parameters used only for two-body interactions ( $n$ ,  $\beta$ ,  $\lambda_2$ ,  $B$ ,  $\lambda_1$ , and  $A$ ) in entries whose second and third element are different (e.g. SiCSi) are not used for anything and can be set to 0.0 if desired.

Note that the twobody parameters in entries such as SiCC and CSiSi are often the same, due to the common use of symmetric mixing rules, but this is not always the case. For example, the beta and n parameters in Tersoff\_2 (*Tersoff\_2*) are not symmetric.

We chose the above form so as to enable users to define all commonly used variants of the Tersoff portion of the potential. In particular, our form reduces to the original Tersoff form when  $m = 3$  and  $\gamma = 1$ , while it reduces to the form of *Albe et al.* when  $\beta = 1$  and  $m = 1$ . Note that in the current Tersoff implementation in LAMMPS,  $m$  must be specified as either 3 or 1. Tersoff used a slightly different but equivalent form for alloys, which we will refer to as Tersoff\_2 potential (*Tersoff\_2*).

LAMMPS parameter values for Tersoff\_2 can be obtained as follows:  $\gamma = \omega_{ijk}$ ,  $\lambda_3 = 0$  and the value of  $m$  has no effect. The parameters for species  $i$  and  $j$  can be calculated using the Tersoff\_2 mixing rules:

$$\begin{aligned}\lambda_1^{i,j} &= \frac{1}{2}(\lambda_1^i + \lambda_1^j) \\ \lambda_2^{i,j} &= \frac{1}{2}(\lambda_2^i + \lambda_2^j) \\ A_{i,j} &= (A_i A_j)^{1/2} \\ B_{i,j} &= \chi_{ij} (B_i B_j)^{1/2} \\ R_{i,j} &= (R_i R_j)^{1/2} \\ S_{i,j} &= (S_i S_j)^{1/2}\end{aligned}$$

Tersoff\_2 parameters  $R$  and  $S$  must be converted to the LAMMPS parameters  $R$  and  $D$  ( $R$  is different in both forms), using the following relations:  $R=(R'+S')/2$  and  $D=(S'-R')/2$ , where the primes indicate the Tersoff\_2 parameters.

In the potentials directory, the file SiCGe.tersoff provides the LAMMPS parameters for Tersoff's various versions of Si, as well as his alloy parameters for Si, C, and Ge. This file can be used for pure Si, (three different versions), pure C, pure Ge, binary SiC, and binary SiGe. LAMMPS will generate an error if this file is used with any combination involving C and Ge, since there are no entries for the GeC interactions (Tersoff did not publish parameters for this cross-interaction.) Tersoff files are also provided for the SiC alloy (SiC.tersoff) and the GaN (GaN.tersoff) alloys.

Many thanks to Rutuparna Narulkar, David Farrell, and Xiaowang Zhou for helping clarify how Tersoff parameters for alloys have been defined in various papers. Also thanks to Ram Devanathan for providing the base ZBL implementation.

The *shift* keyword computes the energy  $E$  of a system of atoms, whose formula is the same as the Tersoff potential. The only modification is that the original equilibrium bond length ( $r_0$ ) of the system is shifted to  $r_0 - \delta$ . The minus sign arises because each radial distance  $r$  is replaced by  $r + \delta$ . More information on this option is given on the main [pair\\_tersoff](#) page.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.257.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, where types I and J correspond to two different element types, mixing is performed by LAMMPS as described above from values in the potential file.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.257.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

The *shift* keyword is currently not supported for the *tersoff/gpu* and *tersoff/kk* variants of this pair style.

The *tersoff/zbl* potential files provided with LAMMPS (see the potentials directory) are parameterized for “*metal*” *units*. Also the pair style supports converting potential file parameters on-the-fly between “metal” and “real” units. You can use the *tersoff/zbl* pair style with any LAMMPS units, but you would need to create your own *tersoff/zbl* potential file with coefficients listed in the appropriate units if your simulation does not use “metal” or “real” units.

#### 19.257.6 Related commands

*pair\_coeff*

#### 19.257.7 Default

none

---

(**Tersoff\_1**) J. Tersoff, Phys Rev B, 37, 6991 (1988).

(**ZBL**) J.F. Ziegler, J.P. Biersack, U. Littmark, ‘Stopping and Ranges of Ions in Matter’ Vol 1, 1985, Pergamon Press.

(**Albe**) J. Nord, K. Albe, P. Erhart and K. Nordlund, J. Phys.: Condens. Matter, 15, 5649(2003).

(**Tersoff\_2**) J. Tersoff, Phys Rev B, 39, 5566 (1989); errata (PRB 41, 3248)

## 19.258 `pair_style thole` command

## 19.259 `pair_style lj/cut/thole/long` command

Accelerator Variants: *lj/cut/thole/long/omp*

### 19.259.1 Syntax

```
pair_style style args
```

- `style` = *thole* or *lj/cut/thole/long*
- `args` = list of arguments for a particular style

*thole* args = damp cutoff

damp = global damping parameter

cutoff = global cutoff (distance units)

*lj/cut/thole/long* args = damp cutoff (cutoff2)

damp = global damping parameter

cutoff = global cutoff for LJ (and Thole if only 1 arg) (distance units)

cutoff2 = global cutoff for Thole (optional) (distance units)

### 19.259.2 Examples

```
pair_style hybrid/overlay ... thole 2.6 12.0
pair_coeff 1 1 thole 1.0
pair_coeff 1 2 thole 1.0 2.6 10.0
pair_coeff * 2 thole 1.0 2.6

pair_style lj/cut/thole/long 2.6 12.0
```

Example input scripts available: `examples/PACKAGES/drude`

### 19.259.3 Description

The *thole* pair styles are meant to be used with force fields that include explicit polarization through Drude dipoles. This link describes how to use the *thermalized Drude oscillator model* in LAMMPS and polarizable models in LAMMPS are discussed on the *Howto polarizable* doc page.

The *thole* pair style should be used as a sub-style within in the *pair\_style hybrid/overlay* command, in conjunction with a main pair style including Coulomb interactions, i.e. any pair style containing *coul/cut* or *coul/long* in its style name.

The *lj/cut/thole/long* pair style is equivalent to, but more convenient than the frequent combination *hybrid/overlay lj/cut/coul/long cutoff thole damp cutoff2*. It is not only a shorthand for this `pair_style` combination, but it also allows for mixing pair coefficients instead of listing them all. The *lj/cut/thole/long* pair style is also a bit faster because it avoids an overlay and can benefit from OMP acceleration. Moreover, it uses a more precise approximation of the direct Coulomb interaction at short range similar to *coul/long/cs*, which stabilizes the temperature of Drude particles.

The *thole* pair styles compute the Coulomb interaction damped at short distances by a function

$$T_{ij}(r_{ij}) = 1 - \left(1 + \frac{s_{ij}r_{ij}}{2}\right) \exp(-s_{ij}r_{ij})$$

This function results from an adaptation to point charges (*Noskov*) of the dipole screening scheme originally proposed by *Thole*. The scaling coefficient  $s_{ij}$  is determined by the polarizability of the atoms,  $\alpha_i$ , and by a Thole damping parameter  $a$ . This Thole damping parameter usually takes a value of 2.6, but in certain force fields the value can depend upon the atom types. The mixing rule for Thole damping parameters is the arithmetic average, and for polarizabilities the geometric average between the atom-specific values.

$$s_{ij} = \frac{a_{ij}}{(\alpha_{ij})^{1/3}} = \frac{(a_i + a_j)/2}{[(\alpha_i \alpha_j)^{1/2}]^{1/3}}$$

The damping function is only applied to the interactions between the point charges representing the induced dipoles on polarizable sites, that is, charges on Drude particles,  $q_{D,i}$ , and opposite charges,  $-q_{D,i}$ , located on the respective core particles (to which each Drude particle is bonded). Therefore, Thole screening is not applied to the full charge of the core particle  $q_i$ , but only to the  $-q_{D,i}$  part of it.

The interactions between core charges are subject to the weighting factors set by the *special\_bonds* command. The interactions between Drude particles and core charges or non-polarizable atoms are also subject to these weighting factors. The Drude particles inherit the 1-2, 1-3 and 1-4 neighbor relations from their respective cores.

For pair\_style *thole*, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the example above.

- $\alpha$  (distance units<sup>3</sup>)
- damp
- cutoff (distance units)

The last two coefficients are optional. If not specified the global Thole damping parameter or global cutoff specified in the pair\_style command are used. In order to specify a cutoff (third argument) a damp parameter (second argument) must also be specified.

For pair style *lj/cut/thole/long*, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command.

- $\epsilon$  (energy units)
- $\sigma$  (length units)
- $\alpha$  (distance units<sup>3</sup>)
- damp
- LJ cutoff (distance units)

The last two coefficients are optional and default to the global values from the *pair\_style* command line.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

### 19.259.4 Mixing, shift, table, tail correction, restart, rRESPA info

The *thole* pair style does not support mixing. Thus, coefficients for all I,J pairs must be specified explicitly.

The *lj/cut/thole/long* pair style does support mixing. Mixed coefficients are defined using

$$\alpha_{ij} = \sqrt{\alpha_i \alpha_j}$$

$$a_{ij} = \frac{1}{2}(a_i + a_j)$$

### 19.259.5 Restrictions

These pair styles are part of the DRUDE package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This `pair_style` should currently not be used with the *charmm dihedral style* if the latter has non-zero 1-4 weighting factors. This is because the *thole* pair style does not know which pairs are 1-4 partners of which dihedrals.

The *lj/cut/thole/long* pair style should be used with a *Kspace solver* like PPPM or Ewald, which is only enabled if LAMMPS was built with the *kpace* package.

### 19.259.6 Related commands

*fix drude*, *fix langevin/drude*, *fix drude/transform*, *compute temp/drude pair\_style lj/cut/coul/long*

### 19.259.7 Default

none

---

(**Noskov**) Noskov, Lamoureux and Roux, J Phys Chem B, 109, 6705 (2005).

(**Thole**) Chem Phys, 59, 341 (1981).

## 19.260 pair\_style threebody/table command

### 19.260.1 Syntax

```
pair_style style
```

- `style = threebody/table`

## 19.260.2 Examples

```
pair_style threebody/table
pair_coeff * * spce2.3b type1 type2

pair_style hybrid/overlay table linear 1200 threebody/table
pair_coeff      1 1 table table_CG_CG.txt VOTCA
pair_coeff      * * threebody/table spce.3b type
```

Used in example input scripts:

```
examples/PACKAGES/manybody_table/in.spce
examples/PACKAGES/manybody_table/in.spce2
```

## 19.260.3 Description

New in version 2Jun2022.

The *threebody/table* style is a pair style for generic tabulated three-body interactions. It has been developed for (coarse-grained) simulations (of water) with Kernel-based machine learning (ML) potentials (*Scherer2*). As for many other MANYBODY package pair styles the energy of a system is computed as a sum over three-body terms:

$$E = \sum_i \sum_{j \neq i} \sum_{k > j} \phi_3(r_{ij}, r_{ik}, \theta_{ijk})$$

The summations in the formula are over all neighbors J and K of atom I within a cutoff distance *cut*. In contrast to the Stillinger-Weber potential, all forces are not calculated analytically, but read in from a three-body force/energy table which can be generated with the *csg\_ml* app of VOTCA as available at: <https://gitlab.mpcdf.mpg.de/votca/votca>.

Only a single *pair\_coeff* command is used with the *threebody/table* style which specifies a threebody potential (“*.3b*”) file with parameters for all needed elements. These are then mapped to LAMMPS atom types by specifying *N\_el* additional arguments after the “*.3b*” filename in the *pair\_coeff* command, where *N\_el* is the number of LAMMPS atom types:

- “*.3b*” filename
- *N\_el* element names = mapping of threebody elements to atom types

See the *pair\_coeff* page for alternate ways to specify the path for the potential file.

As an example, imagine a file *SiC.3b* has three-body values for Si and C. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Si, and the fourth to be C, you would use the following *pair\_coeff* command:

```
pair_coeff * * SiC.3b Si Si Si C
```

The first 2 arguments must be *\*\** so as to span all LAMMPS atom types. The first three Si arguments map LAMMPS atom types 1,2,3 to the Si element in the “*.3b*” file. The final C argument maps LAMMPS atom type 4 to the C element in the threebody file. If a mapping value is specified as NULL, the mapping is not performed. This can be used when a *threebody/table* potential is used as part of the *hybrid* pair style. The NULL values are placeholders for atom types that will be used with other potentials.

The three-body files have a “*.3b*” suffix. Lines that are not blank or comments (starting with #) define parameters for a triplet of elements. The parameters in a single entry specify to the (three-body) cutoff distance and the tabulated three-body interaction. A single entry then contains:

- element 1 (the center atom in a 3-body interaction)
- element 2



- element 3
- cut (distance units)
- filename
- keyword
- style
- N

The parameter *cut* is the (three-body) cutoff distance. When set to 0, no interaction is calculated for this element triplet. The parameters *filename*, *keyword*, *style*, and *N* refer to the tabulated three-body potential.

The tabulation is done on a three-dimensional grid of the two distances  $r_{ij}$  and  $r_{ik}$  as well as the angle  $\theta_{ijk}$  which is constructed in the following way. There are two different cases. If element 2 and element 3 are of the same type (e.g. SiCC), the distance  $r_{ij}$  is varied in “N” steps from *rmin* to *rmax* and the distance  $r_{ik}$  is varied from  $r_{ij}$  to *rmax*. This can be done, due to the symmetry of the triplet. If element 2 and element 3 are not of the same type (e.g. SiCSi), there is no additional symmetry and the distance  $r_{ik}$  is also varied from *rmin* to *rmax* in “N” steps. The angle  $\theta_{ijk}$  is always varied in “2N” steps from  $(0.0 + 180.0/(4N))$  to  $(180.0 - 180.0/(4N))$ . Therefore, the total number of table entries is “M = N \* N \* (N+1)” for the symmetric (element 2 and element 3 are of the same type) and “M = 2 \* N \* N \* N” for the general case (element 2 and element 3 are not of the same type).

The forces on all three particles I, J, and K of a triplet of this type of three-body interaction potential ( $\phi_3(r_{ij}, r_{ik}, \theta_{ijk})$ ) lie within the plane defined by the three inter-particle distance vectors  $\mathbf{r}_{ij}$ ,  $\mathbf{r}_{ik}$ , and  $\mathbf{r}_{jk}$ . This property is used to project the forces onto the inter-particle distance vectors as follows

$$\begin{pmatrix} \mathbf{f}_i \\ \mathbf{f}_j \\ \mathbf{f}_k \end{pmatrix} = \begin{pmatrix} f_{i1} & f_{i2} & 0 \\ f_{j1} & 0 & f_{j2} \\ 0 & f_{k1} & f_{k2} \end{pmatrix} \begin{pmatrix} \mathbf{r}_{ij} \\ \mathbf{r}_{ik} \\ \mathbf{r}_{jk} \end{pmatrix}$$

and then tabulate the 6 force constants  $f_{i1}$ ,  $f_{i2}$ ,  $f_{j1}$ ,  $f_{j2}$ ,  $f_{k1}$ , and  $f_{k2}$ , as well as the energy of a triplet *e*. Due to symmetry reasons, the following relations hold:  $f_{i1} = -f_{j1}$ ,  $f_{i2} = -f_{k1}$ , and  $f_{j2} = -f_{k2}$ . As in this pair style the forces are read in directly, a correct MD simulation is also performed in the case that the triplet energies are set to *e*=0.

The *filename* specifies the file containing the tabulated energy and derivative values of  $\phi_3(r_{ij}, r_{ik}, \theta_{ijk})$ . The *keyword* then specifies a section of the file. The format of this file is as follows (without the parenthesized comments):

```
# Tabulated three-body potential for spce water (one or more comment or blank lines)

ENTRY1                                     (keyword is_
↪the first text on line)
N 12 rmin 2.55 rmax 3.65                   (N, rmin,_
↪rmax parameters)
                                           (blank line)
1 2.55 2.55 3.75 -867.212 -611.273 867.212 21386.8 611.273 -21386.8 0.0 (index, r_ij,
↪ r_ik, theta, f_i1, f_i2, f_j1, f_j2, f_k1, f_k2, e)
2 2.55 2.55 11.25 -621.539 -411.189 621.539 5035.95 411.189 -5035.95 0.0
...
1872 3.65 3.65 176.25 -0.00215132 -0.00412886 0.00215137 0.00111754 0.00412895 -0.
↪00111757 0.0
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The next line lists (in any order) one or more parameters for the table. Each parameter is a keyword followed by one or more numeric values.

The parameter “N” is required. It should be the same than the parameter “N” of the “.3b” file, otherwise its value is overwritten. “N” determines the number of table entries “M” that follow: “M = N \* N \* (N+1)” (symmetric triplet,

e.g. SiCC) or “ $M = 2 * N * N * N$ ” (asymmetric triplet, e.g. SiCSi). Therefore “ $M = 12 * 12 * 13 = 1872$ ” in the above symmetric example. The parameters “rmin” and “rmax” are also required and determine the minimum and maximum of the inter-particle distances  $r_{ij}$  and  $r_{ik}$ .

Following a blank line, the next  $M$  lines of the angular table file list the tabulated values. On each line, the first value is the index from 1 to  $M$ , the second value is the distance  $r_{ij}$ , the third value is the distance  $r_{ik}$ , the fourth value is the angle  $\theta_{ijk}$ , the next six values are the force constants  $f_{i1}$ ,  $f_{i2}$ ,  $f_{j1}$ ,  $f_{j2}$ ,  $f_{k1}$ , and  $f_{k2}$ , and the last value is the energy  $e$ .

Note that one three-body potential file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified *keyword* of appropriate section of the “.3b” file.

At the moment, only the *style linear* is allowed and implemented. After reading in the force table, it is internally stored in LAMMPS as a lookup table. For each triplet configuration occurring in the simulation within the cutoff distance, the next nearest tabulated triplet configuration is looked up. No interpolation is done. This allows for a very efficient force calculation with the stored force constants and energies. Due to the know table structure, the lookup can be done efficiently. It has been tested (*Scherer2*) that with a reasonably small bin size, the accuracy and speed is comparable to that of a Stillinger-Weber potential with tabulated three-body interactions (*pair\_style sw/angle/table*) while the table format of this pair style allows for more flexible three-body interactions.

As for the Stillinger-Weber potential, the three-body potential file must contain entries for all the elements listed in the *pair\_coeff* command. It can also contain entries for additional elements not being used in a particular simulation; LAMMPS ignores those entries.

For a single-element simulation, only a single entry is required (e.g. SiSiSi). For a two-element simulation, the file must contain 8 entries (for SiSiSi, SiSiC, SiCSi, SiCC, CSiSi, CSiC, CCSi, CCC), that specify threebody parameters for all permutations of the two elements interacting in three-body configurations. Thus for 3 elements, 27 entries would be required, etc.

As annotated above, the first element in the entry is the center atom in a three-body interaction. Thus an entry for SiCC means a Si atom with 2 C atoms as neighbors. The tabulated three-body forces can in principle be specific to the three elements of the configuration. However, the user must ensure that it makes physically sense. E.g., the tabulated three-body forces for the entries CSiC and CCSi should be the same exchanging  $r_{ij}$  with  $r_{ik}$ ,  $f_{j1}$  with  $f_{k1}$ , and  $f_{j2}$  with  $f_{k2}$ .

Additional input files and reference data can be found at: <https://gitlab.mpcdf.mpg.de/votca/votca/-/tree/master/csg-tutorials/ml>

---

#### 19.260.4 Mixing, shift, table, tail correction, restart, rRESPA info

As all interactions are tabulated, no mixing is performed.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

## 19.260.5 Restrictions

This pair style is part of the MANYBODY package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires the *newton* setting to be “on” for pair interactions.

## 19.260.6 Related commands

*pair\_coeff*, *pair sw/angle/table*

(Scherer<sup>2</sup>) C. Scherer, R. Scheid, D. Andrienko, and T. Bereau, J. Chem. Theor. Comp. 16, 3194-3204 (2020).

## 19.261 pair\_style tracker command

### 19.261.1 Syntax

```
pair_style tracker fix_ID N keyword values attribute1 attribute2 ...
```

- fix\_ID = ID of associated internal fix to store data
- N = prepare data for output every this many timesteps
- zero or more keywords may be appended
- keyword = *finite* or *time/min* or *type/include*

*finite* value = none

pair style uses atomic diameters to identify contacts

*time/min* value = T

T = minimum number of timesteps of interaction

*type/include* value = list1 list2

list1,list2 = separate lists of types (see below)

- one or more attributes may be appended

```
possible attributes = id1 id2 time/created time/broken time/total
                    r/min r/ave x y z
```

id1, id2 = IDs of the 2 atoms in each pair interaction

time/created = the timestep that the 2 atoms began interacting

time/broken = the timestep that the 2 atoms stopped interacting

time/total = the total number of timesteps the 2 atoms interacted

r/min = the minimum radial distance between the 2 atoms during the interaction.  
↪(distance units)

r/ave = the average radial distance between the 2 atoms during the interaction.  
↪(distance units)

x, y, z = the center of mass position of the 2 atoms when they stopped interacting.  
↪(distance units)

## 19.261.2 Examples

```
pair_style hybrid/overlay tracker myfix 1000 id1 id2 type/include 1 * type/include 2 3,4_
→ lj/cut 2.5
pair_coeff 1 1 tracker 2.0

pair_style hybrid/overlay tracker myfix 1000 finite x y z time/min 100 granular
pair_coeff * * tracker

dump 1 all local 1000 dump.local f_myfix[1] f_myfix[2] f_myfix[3]
dump_modify 1 write_header no
```

## 19.261.3 Description

Style *tracker* monitors information about pairwise interactions. It does not calculate any forces on atoms. *Pair hybrid/overlay* can be used to combine this pair style with any other pair style, as shown in the examples above.

At each timestep, if two neighboring atoms move beyond the interaction cutoff, pairwise data is processed and transferred to an internal fix labeled *fix\_ID*. This allows the local data to be accessed by other LAMMPS commands. Additional filters can be applied using the *time/min* or *type/include* keywords described below. Note that this is the interaction cutoff defined by this pair style, not the short-range cutoff defined by the pair style that is calculating forces on atoms.

Following any optional keyword/value arguments, a list of one or more attributes is specified. These include the IDs of the two atoms in the pair. The other attributes for the pair of atoms are the duration of time they were “interacting” or at the point in time they started or stopped interacting. In this context, “interacting” means the time window during which the two atoms were closer than the interaction cutoff distance. The attributes for *time/\** refer to timesteps.

Data is continuously accumulated by the internal fix over intervals of  $N$  timesteps. At the end of each interval, all of the saved accumulated data is deleted to make room for new data. Individual datum may therefore persist anywhere between  $1$  to  $N$  timesteps depending on when they are saved. This data can be accessed using the *fix\_ID* and a *dump local* command. To ensure all data is output, the dump frequency should correspond to the same interval of  $N$  timesteps. A dump frequency of an integer multiple of  $N$  can be used to regularly output a sample of the accumulated data.

---

The following optional keywords may be used.

If the *finite* keyword is not used, the following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- cutoff (distance units)

If the *finite* keyword is used, there are no additional coefficients to set for each pair of atom types via the *pair\_coeff* command. Interaction cutoffs are instead calculated based on the diameter of finite particles. However you must still use the *pair\_coeff* for all atom types. For example the command

```
pair_coeff * *
```

should be used.

The *time/min* keyword sets a minimum amount of time that an interaction must persist to be included. This setting can be used to censor short-lived interactions.

The *type/include* keyword filters interactions based on the types of the two atoms. Data is only saved for interactions between atoms whose two atom types appear in *list1* and *list2*. Atom type 1 must be in *list1* and atom type 2 in *list2*. Or vice versa.

Each type list consists of a series of type ranges separated by commas. Each range can be specified as a single numeric value, or a wildcard asterisk can be used to specify a range of values. This takes the form “\*” or “\*n” or “n\*” or “m\*n”. For example, if  $M$  = the number of atom types, then an asterisk with no numeric values means all types from 1 to  $M$ . A leading asterisk means all types from 1 to  $n$  (inclusive). A trailing asterisk means all types from  $n$  to  $M$  (inclusive). A middle asterisk means all types from  $m$  to  $n$  (inclusive). Note that the *type/include* keyword can be specified multiple times.

#### 19.261.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs  $I, J$  and  $I \neq J$ , the cutoff coefficient and cutoff distance for this pair style can be mixed. The cutoff is always mixed via a *geometric* rule. The cutoff is mixed according to the `pair_modify` mix value. The default mix value is *geometric*. See the “`pair_modify`” command for details.

This pair style writes its information to *binary restart files*, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

The *pair\_modify* shift, table, and tail options are not relevant for this pair style.

The accumulated data is not written to restart files and should be output before a restart file is written to avoid missing data.

The internal fix calculates a local vector or local array depending on the number of input values. The length of the vector or number of rows in the array is the number of recorded, lost interactions. If a single input is specified, a local vector is produced. If two or more inputs are specified, a local array is produced where the number of columns = the number of inputs. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector or array will be floating point values that correspond to the specified attribute.

#### 19.261.5 Restrictions

This fix is part of the MISC package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style is currently incompatible with granular pair styles that extend beyond the contact (e.g. JKR and DMT).

#### 19.261.6 Related commands

#### 19.261.7 Default

none

### 19.262 pair\_style tri/lj command

#### 19.262.1 Syntax

```
pair_style tri/lj cutoff
```

cutoff = global cutoff for interactions (distance units)

## 19.262.2 Examples

```
pair_style tri/lj 3.0
pair_coeff * * 1.0 1.0
pair_coeff 1 1 1.0 1.5 2.5
```

## 19.262.3 Description

Style *tri/lj* treats particles which are triangles as a set of small spherical particles that tile the triangle surface as explained below. Interactions between two triangles, each with N1 and N2 spherical particles, are calculated as the pairwise sum of N1\*N2 Lennard-Jones interactions. Interactions between a triangle with N spherical particles and a point particle are treated as the pairwise sum of N Lennard-Jones interactions. See the *pair\_style lj/cut* doc page for the definition of Lennard-Jones interactions.

The cutoff distance for an interaction between 2 triangles, or between a triangle and a point particle, is calculated from the position of the triangle (its centroid), not between pairs of individual spheres comprising the triangle. Thus an interaction is either calculated in its entirety or not at all.

The set of non-overlapping spherical particles that represent a triangle, for purposes of this pair style, are generated in the following manner. Assume the triangle is of type I, and `sigma_II` has been specified. We want a set of spheres with centers in the plane of the triangle, none of them larger in diameter than `sigma_II`, which completely cover the triangle's area, but with minimal overlap and a minimal total number of spheres. This is done in a recursive manner. Place a sphere at the centroid of the original triangle. Calculate what diameter it must have to just cover all 3 corner points of the triangle. If that diameter is equal to or smaller than `sigma_II`, then include a sphere of the calculated diameter in the set of covering spheres. If the diameter is larger than `sigma_II`, then split the triangle into 2 triangles by bisecting its longest side. Repeat the process on each sub-triangle, recursing as far as needed to generate a set of covering spheres. When finished, the original criteria are met, and the set of covering spheres should be near minimal in number and overlap, at least for input triangles with a reasonable aspect-ratio.

The LJ interaction between 2 spheres on different triangles of types I,J is computed with an arithmetic mixing of the sigma values of the 2 spheres and using the specified epsilon value for I,J atom types. Note that because the sigma values for triangles spheres is computed using only `sigma_II` values, specific to the triangles's type, this means that any specified `sigma_IJ` values (for I != J) are effectively ignored.

For style *tri/lj*, the following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- epsilon (energy units)
- sigma (distance units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global cutoff is used.

### 19.262.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the epsilon and sigma coefficients and cutoff distance for all of this pair style can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

### 19.262.5 Restrictions

This style is part of the ASPHERE package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

Defining particles to be triangles so they participate in tri/tri or tri/particle interactions requires the use the *atom\_style tri* command.

### 19.262.6 Related commands

*pair\_coeff*, *pair\_style line/lj*

### 19.262.7 Default

none

## 19.263 pair\_style ufm command

Accelerator Variants: *ufm/gpu*, *ufm/omp*, *ufm/opt*

### 19.263.1 Syntax

```
pair_style ufm cutoff
```

- cutoff = global cutoff for *ufm* interactions (distance units)

### 19.263.2 Examples

```
pair_style ufm 4.0
pair_coeff 1 1 100.0 1.0 2.5
pair_coeff * * 100.0 1.0

pair_style ufm 4.0
pair_coeff * * 10.0 1.0
variable prefactor equal ramp(10,100)
fix 1 all adapt 1 pair ufm epsilon * * v_prefactor
```

### 19.263.3 Description

Style *ufm* computes pairwise interactions using the Uhlenbeck-Ford model (UFM) potential ([Paula Leite2016](#)) which is given by

$$E = -\epsilon \ln [1 - \exp(-r^2/\sigma^2)] \quad r < r_c$$
$$\epsilon = p k_B T$$

where  $r_c$  is the cutoff,  $\sigma$  is a distance-scale and  $\epsilon$  is an energy-scale, i.e., a product of Boltzmann constant  $k_B$ , temperature  $T$  and the Uhlenbeck-Ford  $p$ -parameter which is responsible to control the softness of the interactions ([Paula Leite2017](#)). This model is useful as a reference system for fluid-phase free-energy calculations ([Paula Leite2016](#)).

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global *ufm* cutoff is used.

The *fix adapt* command can be used to vary epsilon and sigma for this pair style over the course of a simulation, in which case *pair\_coeff* settings for epsilon and sigma must still be specified, but will be overridden. For example these commands will vary the prefactor epsilon for all pairwise interactions from 10.0 at the beginning to 100.0 at the end of a run:

```
variable prefactor equal ramp(10,100)
fix 1 all adapt 1 pair ufm epsilon * * v_prefactor
```

---

**Note:** The thermodynamic integration procedure can be performed with this potential using *fix adapt*. This command will rescale the force on each atom by varying a scale variable, which always starts with value 1.0. The syntax is the same described above, however, changing epsilon to scale. A detailed explanation of how to use this command and perform nonequilibrium thermodynamic integration in LAMMPS is given in the paper by ([Freitas](#)).

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---



### 19.263.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , the epsilon and sigma coefficients and cutoff distance for this pair style can be mixed. The default mix value is *geometric*. See the “pair\_modify” command for details.

This pair style support the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table and tail are not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

### 19.263.5 Restrictions

This pair style is part of the EXTRA-PAIR package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

### 19.263.6 Related commands

*pair\_coeff*, *fix adapt*

### 19.263.7 Default

none

**(Paula Leite2017)** Paula Leite, Santos-Florez, and de Koning, Phys Rev E, 96, 32115 (2017).

**(Paula Leite2016)** Paula Leite, Freitas, Azevedo, and de Koning, J Chem Phys, 126, 044509 (2016).

**(Freitas)** Freitas, Asta, and de Koning, Computational Materials Science, 112, 333 (2016).

## 19.264 pair\_style vashishta command

Accelerator Variants: *vashishta/gpu*, *vashishta/omp*, *vashishta/kk*

## 19.265 pair\_style vashishta/table command

Accelerator Variants: *vashishta/table/omp*

## 19.265.1 Syntax

```
pair_style style args
```

- style = *vashishta* or *vashishta/table*
- args = list of arguments for a particular style

*vashishta* args = none

*vashishta/table* args = Ntable cutinner

Ntable = # of tabulation points

cutinner = tabulate from cutinner to cutoff

## 19.265.2 Examples

```
pair_style vashishta
pair_coeff * * SiC.vashishta Si C

pair_style vashishta/table 100000 0.2
pair_coeff * * SiC.vashishta Si C
```

## 19.265.3 Description

The *vashishta* and *vashishta/table* styles compute the combined 2-body and 3-body family of potentials developed in the group of Priya Vashishta and collaborators. By combining repulsive, screened Coulombic, screened charge-dipole, and dispersion interactions with a bond-angle energy based on the Stillinger-Weber potential, this potential has been used to describe a variety of inorganic compounds, including SiO2 *Vashishta1990*, SiC *Vashishta2007*, and InP *Branicio2009*.

The potential for the energy  $U$  of a system of atoms is

$$U = \sum_i \sum_{j>i}^N U_{ij}^{(2)}(r_{ij}) + \sum_i \sum_{j \neq i} \sum_{k>j, k \neq i}^N U_{ijk}^{(3)}(r_{ij}, r_{ik}, \theta_{ijk})$$

$$U_{ij}^{(2)}(r) = \frac{H_{ij}}{r^{\eta_{ij}}} + \frac{Z_i Z_j}{r} \exp(-r/\lambda_{1,ij}) - \frac{D_{ij}}{r^4} \exp(-r/\lambda_{4,ij}) - \frac{W_{ij}}{r^6}, r < r_{c,ij}$$

$$U_{ijk}^{(3)}(r_{ij}, r_{ik}, \theta_{ijk}) = B_{ijk} \frac{[\cos \theta_{ijk} - \cos \theta_{0ijk}]^2}{1 + C_{ijk} [\cos \theta_{ijk} - \cos \theta_{0ijk}]^2} \times$$

$$\exp\left(\frac{\gamma_{ij}}{r_{ij} - r_{0,ij}}\right) \exp\left(\frac{\gamma_{ik}}{r_{ik} - r_{0,ik}}\right), r_{ij} < r_{0,ij}, r_{ik} < r_{0,ik}$$

where we follow the notation used in *Branicio2009*.  $U^2$  is a two-body term and  $U^3$  is a three-body term. The summation over two-body terms is over all neighbors  $J$  within a cutoff distance =  $r_c$ . The twobody terms are shifted and tilted by a linear function so that the energy and force are both zero at  $r_c$ . The summation over three-body terms is over all neighbors  $i$  and  $k$  within a cut-off distance =  $r_0$ , where the exponential screening function becomes zero.

The *vashishta* style computes these formulas analytically. The *vashishta/table* style tabulates the analytic values for  $Ntable$  points from cutinner to the cutoff of the potential. The points are equally spaced in  $R^2$  space from cutinner<sup>2</sup> to cutoff<sup>2</sup>. For the two-body term in the above equation, a linear interpolation for each pairwise distance between adjacent points in the table. In practice the tabulated version can run 3-5x faster than the analytic version with moderate to little loss of accuracy for  $Ntable$  values between 10000 and 1000000. It is not recommended to use less than 5000 tabulation points.

Only a single `pair_coeff` command is used with either style which specifies a Vashishta potential file with parameters for all needed elements. These are mapped to LAMMPS atom types by specifying N additional arguments after the filename in the `pair_coeff` command, where N is the number of LAMMPS atom types:

- filename
- N element names = mapping of Vashishta elements to atom types

See the [pair\\_coeff](#) page for alternate ways to specify the path for the potential file.

As an example, imagine a file `SiC.vashishta` has parameters for Si and C. If your LAMMPS simulation has 4 atoms types and you want the first 3 to be Si, and the fourth to be C, you would use the following `pair_coeff` command:

```
pair_coeff * * SiC.vashishta Si Si Si C
```

The first 2 arguments must be `**` so as to span all LAMMPS atom types. The first three Si arguments map LAMMPS atom types 1,2,3 to the Si element in the file. The final C argument maps LAMMPS atom type 4 to the C element in the file. If a mapping value is specified as `NULL`, the mapping is not performed. This can be used when a *vashishta* potential is used as part of the *hybrid* pair style. The `NULL` values are placeholders for atom types that will be used with other potentials.

Vashishta files in the *potentials* directory of the LAMMPS distribution have a “.vashishta” suffix. Lines that are not blank or comments (starting with #) define parameters for a triplet of elements. The parameters in a single entry correspond to the two-body and three-body coefficients in the formulae above:

- element 1 (the center atom in a 3-body interaction)
- element 2
- element 3
- $H$  (energy units)
- $\eta$
- $Z_i$  (electron charge units)
- $Z_j$  (electron charge units)
- $\lambda_1$  (distance units)
- $D$  (energy units)
- $\lambda_4$  (distance units)
- $W$  (energy units)
- $r_c$  (distance units)
- $B$  (energy units)
- $\gamma$
- $r_0$  (distance units)
- $C$
- $\cos \theta_0$

The non-annotated parameters are unitless. The Vashishta potential file must contain entries for all the elements listed in the `pair_coeff` command. It can also contain entries for additional elements not being used in a particular simulation; LAMMPS ignores those entries. For a single-element simulation, only a single entry is required (e.g. SiSiSi). For a two-element simulation, the file must contain 8 entries (for SiSiSi, SiSiC, SiCSi, SiCC, CSiSi, CSiC, CCSi, CCC), that specify parameters for all permutations of the two elements interacting in three-body configurations. Thus for 3 elements, 27 entries would be required, etc.

Depending on the particular version of the Vashishta potential, the values of these parameters may be keyed to the identities of zero, one, two, or three elements. In order to make the input file format unambiguous, general, and simple to code, LAMMPS uses a slightly confusing method for specifying parameters. All parameters are divided into two classes: two-body and three-body. Two-body and three-body parameters are handled differently, as described below. The two-body parameters are  $H$ ,  $\eta$ ,  $\lambda_1$ ,  $D$ ,  $\lambda_4$ ,  $W$ ,  $r_c$ ,  $\gamma$ , and  $r_0$ . They appear in the above formulae with two subscripts. The parameters  $Z_i$  and  $Z_j$  are also classified as two-body parameters, even though they only have 1 subscript. The three-body parameters are  $B$ ,  $C$ ,  $\cos \theta_0$ . They appear in the above formulae with three subscripts. Two-body and three-body parameters are handled differently, as described below.

The first element in each entry is the center atom in a three-body interaction, while the second and third elements are two neighbor atoms. Three-body parameters for a central atom I and two neighbors J and K are taken from the IJK entry. Note that even though three-body parameters do not depend on the order of J and K, LAMMPS stores three-body parameters for both IJK and IKJ. The user must ensure that these values are equal. Two-body parameters for an atom I interacting with atom J are taken from the IJJ entry, where the second and third elements are the same. Thus the two-body parameters for Si interacting with C come from the SiCC entry. Note that even though two-body parameters (except possibly gamma and r0 in U3) do not depend on the order of the two elements, LAMMPS will get the Si-C value from the SiCC entry and the C-Si value from the CSiSi entry. The user must ensure that these values are equal. Two-body parameters appearing in entries where the second and third elements are different are stored but never used. It is good practice to enter zero for these values. Note that the three-body function U3 above contains the two-body parameters  $\gamma$  and  $r_0$ . So U3 for a central C atom bonded to an Si atom and a second C atom will take three-body parameters from the CSiC entry, but two-body parameters from the CCC and CSiSi entries.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 19.265.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and  $I \neq J$ , where types I and J correspond to two different element types, mixing is performed by LAMMPS as described above from values in the potential file.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style does not write its information to *binary restart files*, since it is stored in potential files. Thus, you need to re-specify the *pair\_style* and *pair\_coeff* commands in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

## 19.265.5 Restrictions

These pair styles are part of the MANYBODY package. They are only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

These pair styles requires the *newton* setting to be “on” for pair interactions.

The Vashishta potential files provided with LAMMPS (see the potentials directory) are parameterized for metal *units*. You can use the Vashishta potential with any LAMMPS units, but you would need to create your own potential file with coefficients listed in the appropriate units if your simulation does not use “metal” units.

## 19.265.6 Related commands

*pair\_coeff*

## 19.265.7 Default

none

(Vashishta1990) P. Vashishta, R. K. Kalia, J. P. Rino, Phys. Rev. B 41, 12197 (1990).

(Vashishta2007) P. Vashishta, R. K. Kalia, A. Nakano, J. P. Rino. J. Appl. Phys. 101, 103515 (2007).

(Branicio2009) Branicio, Rino, Gan and Tsuzuki, J. Phys Condensed Matter 21 (2009) 095002

## 19.266 pair\_style wf/cut command

### 19.266.1 Syntax

```
pair_style wf/cut cutoff
```

- cutoff = cutoff for wf interactions (distance units)

### 19.266.2 Examples

```
pair_style      wf/cut 2.0
pair_coeff      1 1 1.0 1.0 1 1 2.0
```

### 19.266.3 Description

The *wf/cut* (Wang-Frenkel) style computes LJ-like potentials as described in *Wang2020*. This potential is by construction finite ranged and it vanishes quadratically at the cutoff distance, avoiding truncation, shifting, interpolation and other typical procedures with the LJ potential. The *wf/cut* can be used when a typical short-ranged potential with attraction is required. The potential is given by which is given by:

$$\phi(r) = \epsilon\alpha \left( \left[ \frac{\sigma}{r} \right]^{2\mu} - 1 \right) \left( \left[ \frac{r_c}{r} \right]^{2\mu} - 1 \right)^{2\nu}$$

with

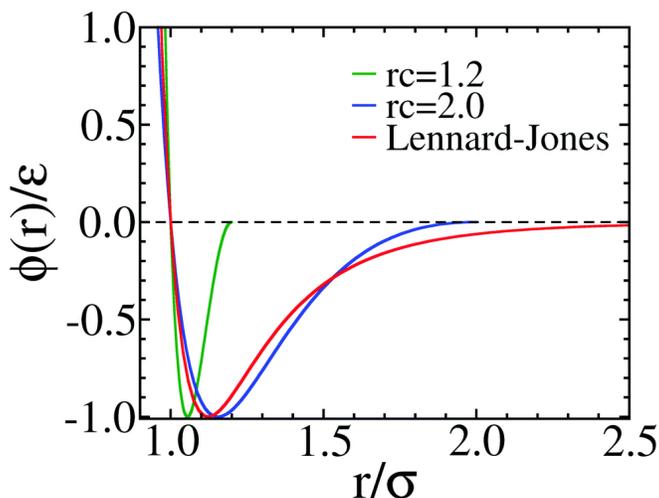
$$\alpha = 2\nu \left(\frac{r_c}{\sigma}\right)^{2\mu} \left[ \frac{1 + 2\nu}{2\nu [(r_c/\sigma)^{2\mu} - 1]} \right]^{2\nu+1}$$

and

$$r_{min} = r_c \left[ \frac{1 + 2\nu}{1 + 2\nu(r_c/\sigma)^{2\nu}} \right]^{1/2\nu}$$

$r_c$  is the cutoff.

Comparison of the non-truncated Lennard-Jones 12-6 potential (red curve), and the WF potentials with  $\mu = 1$  and  $\nu = 1$  are shown in the figure below. The blue curve has  $r_c = 2.0$  and the green curve has  $r_c = 1.2$  and can be used to describe colloidal interactions.



The following coefficients must be defined for each pair of atoms types via the `pair_coeff` command as in the example above, or in the data file or restart files read by the `read_data` or `read_restart` commands:

- $\epsilon$  (energy units)
- $\sigma$  (distance units)
- $\nu$
- $\mu$
- $r_c$  (distance units)

The last coefficient is optional. If not specified, the global cutoff given in the `pair_style` command is used. The exponents  $\nu$  and  $\mu$  are positive integers, usually set to 1. There is usually little to be gained by choosing other values of  $\nu$  and  $\mu$  (See discussion in [Wang2020](#))

---

#### Mixing, shift, table, tail correction, restart, rRESPA info:

This pair style does not support the `pair_modify` mixing and table options.

The `pair_modify` tail and shift options are not relevant for this pair style as it goes to zero at the cut-off radius.

This pair style writes its information to `binary restart files`, so `pair_style` and `pair_coeff` commands do not need to be specified in an input script that reads a restart file.

This pair style does not support the use of the `inner`, `middle`, and `outer` keywords of the `run_style respa` command.

---

## 19.266.4 Restrictions

This pair style can only be used if LAMMPS was built with the EXTRA-PAIR package. See the *Build package* doc page for more info.

## 19.266.5 Related commands

*pair\_coeff*

**Default:** none

---

(Wang2020) X. Wang, S. Ramirez-Hinestrosa, J. Dobnikar, and D. Frenkel, Phys. Chem. Chem. Phys. 22, 10624 (2020).

## 19.267 pair\_style yukawa command

Accelerator Variants: *yukawa/gpu*, *yukawa/omp*, *yukawa/kk*

### 19.267.1 Syntax

```
pair_style yukawa kappa cutoff
```

- kappa = screening length (inverse distance units)
- cutoff = global cutoff for Yukawa interactions (distance units)

### 19.267.2 Examples

```
pair_style yukawa 2.0 2.5
pair_coeff 1 1 100.0 2.3
pair_coeff * * 100.0
```

### 19.267.3 Description

Style *yukawa* computes pairwise interactions with the formula

$$E = A \frac{e^{-\kappa r}}{r} \quad r < r_c$$

$r_c$  is the cutoff.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- A (energy\*distance units)
- cutoff (distance units)

The last coefficient is optional. If not specified, the global yukawa cutoff is used.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.267.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the A coefficient and cutoff distance for this pair style can be mixed. A is an energy value mixed like a LJ epsilon. The default mix value is *geometric*. See the “pair\_modify” command for details.

This pair style supports the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.267.5 Restrictions

none

#### 19.267.6 Related commands

*pair\_coeff*

#### 19.267.7 Default

none



## 19.268 pair\_style yukawa/colloid command

Accelerator Variants: *yukawa/colloid/gpu*, *yukawa/colloid/omp*

### 19.268.1 Syntax

```
pair_style yukawa/colloid kappa cutoff
```

- kappa = screening length (inverse distance units)
- cutoff = global cutoff for colloidal Yukawa interactions (distance units)

### 19.268.2 Examples

```
pair_style yukawa/colloid 2.0 2.5
pair_coeff 1 1 100.0 2.3
pair_coeff * * 100.0
```

### 19.268.3 Description

Style *yukawa/colloid* computes pairwise interactions with the formula

$$E = \frac{A}{\kappa} e^{-\kappa(r-(r_i+r_j))} \quad r < r_c$$

where  $r_i$  and  $r_j$  are the radii of the two particles and  $r_c$  is the cutoff.

In contrast to *pair\_style yukawa*, this functional form arises from the Coulombic interaction between two colloid particles, screened due to the presence of an electrolyte, see the book by *Safran* for a derivation in the context of DLVO theory. *Pair\_style yukawa* is a screened Coulombic potential between two point-charges and uses no such approximation.

This potential applies to nearby particle pairs for which the Derjagin approximation holds, meaning  $h \ll r_i + r_j$ , where  $h$  is the surface-to-surface separation of the two particles.

When used in combination with *pair\_style colloid*, the two terms become the so-called DLVO potential, which combines electrostatic repulsion and van der Waals attraction.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- A (energy/distance units)
- cutoff (distance units)

The prefactor A is determined from the relationship between surface charge and surface potential due to the presence of electrolyte. Note that the A for this potential style has different units than the A used in *pair\_style yukawa*. For low surface potentials, i.e. less than about 25 mV, A can be written as:

$$A = 2\pi R\epsilon\epsilon_0\kappa\psi^2$$

where

- R = colloid radius (distance units)

- $\epsilon_0$  = permittivity of free space (charge<sup>2</sup>/energy/distance units)
- $\epsilon$  = relative permittivity of fluid medium (dimensionless)
- $\kappa$  = inverse screening length (1/distance units)
- $\psi$  = surface potential (energy/charge units)

The last coefficient is optional. If not specified, the global yukawa/colloid cutoff is used.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.268.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs I,J and I != J, the A coefficient and cutoff distance for this pair style can be mixed. A is an energy value mixed like a LJ epsilon. The default mix value is *geometric*. See the “pair\_modify” command for details.

This pair style supports the *pair\_modify* shift option for the energy of the pair interaction.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure.

This pair style writes its information to *binary restart files*, so pair\_style and pair\_coeff commands do not need to be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

#### 19.268.5 Restrictions

This style is part of the COLLOID package. It is only enabled if LAMMPS was built with that package. See the *Build package* page for more info.

This pair style requires that atoms be finite-size spheres with a diameter, as defined by the *atom\_style sphere* command.

Per-particle polydispersity is not yet supported by this pair style; per-type polydispersity is allowed. This means all particles of the same type must have the same diameter. Each type can have a different diameter.

## 19.268.6 Related commands

*pair\_coeff*

## 19.268.7 Default

none

(**Safran**) Safran, Statistical Thermodynamics of Surfaces, Interfaces, And Membranes, Westview Press, ISBN: 978-0813340791 (2003).

## 19.269 pair\_style zbl command

Accelerator Variants: *zbl/gpu*, *zbl/kk*, *zbl/omp*

### 19.269.1 Syntax

```
pair_style zbl inner outer
```

- inner = distance where switching function begins
- outer = global cutoff for ZBL interaction

### 19.269.2 Examples

```
pair_style zbl 3.0 4.0
pair_coeff * * 73.0 73.0
pair_coeff 1 1 14.0 14.0
```

### 19.269.3 Description

Style *zbl* computes the Ziegler-Biersack-Littmark (ZBL) screened nuclear repulsion for describing high-energy collisions between atoms. (*Ziegler*). It includes an additional switching function that ramps the energy, force, and curvature smoothly to zero between an inner and outer cutoff. The potential energy due to a pair of atoms at a distance  $r_{ij}$  is given by:

$$E_{ij}^{ZBL} = \frac{1}{4\pi\epsilon_0} \frac{Z_i Z_j e^2}{r_{ij}} \phi(r_{ij}/a) + S(r_{ij})$$

$$a = \frac{0.46850}{Z_i^{0.23} + Z_j^{0.23}}$$

$$\phi(x) = 0.18175e^{-3.19980x} + 0.50986e^{-0.94229x} + 0.28022e^{-0.40290x} + 0.02817e^{-0.20162x}$$

where  $e$  is the electron charge,  $\epsilon_0$  is the electrical permittivity of vacuum, and  $Z_i$  and  $Z_j$  are the nuclear charges of the two atoms. The switching function  $S(r)$  is identical to that used by *pair\_style lj/gromacs*. Here, the inner and outer cutoff are the same for all pairs of atom types.

The following coefficients must be defined for each pair of atom types via the *pair\_coeff* command as in the examples above, or in the LAMMPS data file.

- $Z_i$  (atomic number for first atom type, e.g. 13.0 for aluminum)
- $Z_j$  (ditto for second atom type)

The values of  $Z_i$  and  $Z_j$  are normally equal to the atomic numbers of the two atom types. Thus, the user may optionally specify only the coefficients for each  $i == i$  pair, and rely on the obvious mixing rule for cross interactions (see below). Note that when  $i == i$  it is required that  $Z_i == Z_j$ . When used with *hybrid/overlay* and pairs are assigned to more than one sub-style, the mixing rule is not used and each pair of types interacting with the ZBL sub-style must be included in a `pair_coeff` command.

---

**Note:** The numerical values of the exponential decay constants in the screening function depend on the unit of distance. In the above equation they are given for units of angstroms. LAMMPS will automatically convert these values to the distance unit of the specified LAMMPS *units* setting. The values of Z should always be given as multiples of a proton's charge, e.g. 29.0 for copper.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

#### 19.269.4 Mixing, shift, table, tail correction, restart, rRESPA info

For atom type pairs  $i,j$  and  $i \neq i$ , the  $Z_i$  and  $Z_j$  coefficients can be mixed by taking  $Z_i$  and  $Z_j$  from the values specified for  $i == i$  and  $j == j$  cases. When used with *hybrid/overlay* and pairs are assigned to more than one sub-style, the mixing rule is not used and each pair of types interacting with the ZBL sub-style must be included in a `pair_coeff` command. The *pair\_modify* mix option has no effect on the mixing behavior

The ZBL pair style does not support the *pair\_modify* shift option, since the ZBL interaction is already smoothed to 0.0 at the cutoff.

The *pair\_modify* table option is not relevant for this pair style.

This pair style does not support the *pair\_modify* tail option for adding long-range tail corrections to energy and pressure, since there are no corrections for a potential that goes to 0.0 at the cutoff.

This pair style does not write information to *binary restart files*, so `pair_style` and `pair_coeff` commands must be specified in an input script that reads a restart file.

This pair style can only be used via the *pair* keyword of the *run\_style respa* command. It does not support the *inner*, *middle*, *outer* keywords.

---

## 19.269.5 Restrictions

none

## 19.269.6 Related commands

*pair\_coeff*

## 19.269.7 Default

none

---

(Ziegler) J.F. Ziegler, J. P. Biersack and U. Littmark, “The Stopping and Range of Ions in Matter”, Volume 1, Pergamon, 1985.

## 19.270 pair\_style zero command

### 19.270.1 Syntax

```
pair_style zero cutoff [nocoeff]
```

- zero = style name of this pair style
- cutoff = global cutoff (distance units)
- nocoeff = ignore all pair\_coeff parameters (optional)

### 19.270.2 Examples

```
pair_style zero 10.0
pair_style zero 5.0 nocoeff
pair_coeff * *
pair_coeff 1 2*4 3.0
```

### 19.270.3 Description

Define a global or per-type cutoff length for the purpose of building a neighbor list and acquiring ghost atoms, but do not compute any pairwise forces or energies.

This can be useful for fixes or computes which require a neighbor list to enumerate pairs of atoms within some cutoff distance, but when pairwise forces are not otherwise needed. Examples are the *fix bond/create*, *compute rdf*, *compute voronoi/atom* commands.

Note that the *comm\_modify cutoff* command can be used to insure communication of ghost atoms even when a pair style is not defined, but it will not trigger neighbor list generation.

The optional *nocoeff* flag allows to read data files with a PairCoeff section for any pair style. Similarly, any pair\_coeff commands will only be checked for the atom type numbers and the rest ignored. In this case, only the global cutoff will be used.

The following coefficients must be defined for each pair of atoms types via the *pair\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands, or by mixing as described below:

- cutoff (distance units)

This coefficient is optional. If not specified, the global cutoff specified in the *pair\_style* command is used. If the *pair\_style* has been specified with the optional *nocoeff* flag, then a cutoff pair coefficient is ignored.

---

#### 19.270.4 Mixing, shift, table, tail correction, restart, rRESPA info

The cutoff distance for this pair style can be mixed. The default mix value is *geometric*. See the “*pair\_modify*” command for details.

This pair style does not support the *pair\_modify* shift, table, and tail options.

This pair style writes its information to *binary restart files*, so *pair\_style* and *pair\_coeff* commands do not need to be specified in an input script that reads a restart file.

This pair style supports the use of the *inner*, *middle*, and *outer* keywords of the *run\_style respa* command.

---

#### 19.270.5 Restrictions

none

#### 19.270.6 Related commands

*pair\_style none*

#### 19.270.7 Default

none

## BOND STYLES

### 20.1 bond\_style bpm/rotational command

#### 20.1.1 Syntax

```
bond_style bpm/rotational keyword value attribute1 attribute2 ...
```

- optional keyword = *overlay/pair* or *store/local* or *smooth*

*store/local* values = fix\_ID N attributes ...

\* fix\_ID = ID of associated internal fix to store data

\* N = prepare data for output every this many timesteps

\* attributes = zero or more of the below attributes may be appended

*id1, id2* = IDs of 2 atoms in the bond

*time* = the timestep the bond broke

*x, y, z* = the center of mass position of the 2 atoms when the bond broke.

↪(distance units)

*x/ref, y/ref, z/ref* = the initial center of mass position of the 2 atoms.

↪(distance units)

*overlay/pair* value = none

bonded particles will still interact with pair forces

*smooth* value = yes or no

smooths bond forces near the breaking point

#### 20.1.2 Examples

```
bond_style bpm/rotational
bond_coeff 1 1.0 0.2 0.02 0.02 0.20 0.04 0.04 0.04 0.1 0.02 0.002 0.002

bond_style bpm/rotational store/local myfix 1000 time id1 id2
dump 1 all local 1000 dump.broken f_myfix[1] f_myfix[2] f_myfix[3]
dump_modify 1 write_header no
```

### 20.1.3 Description

The *bpm/rotational* bond style computes forces and torques based on deviations from the initial reference state of the two atoms. The reference state is stored by each bond when it is first computed in the setup of a run. Data is then preserved across run commands and is written to *binary restart files* such that restarting the system will not reset the reference state of a bond.

Forces include a normal and tangential component. The base normal force has a magnitude of

$$f_r = k_r(r - r_0)$$

where  $k_r$  is a stiffness and  $r$  is the current distance and  $r_0$  is the initial distance between the two particles.

A tangential force is applied perpendicular to the normal direction which is proportional to the tangential shear displacement with a stiffness of  $k_s$ . This tangential force also induces a torque. In addition, bending and twisting torques are also applied to particles which are proportional to angular bending and twisting displacements with stiffnesses of  $k_b$  and  $k_t$ , respectively. Details on the calculation of shear displacements and angular displacements can be found in *ref*: '(Wang) < Wang2009 > and (Wang and Mora).

Bonds will break under sufficient stress. A breaking criteria is calculated

$$B = \max\left\{0, \frac{f_r}{f_{r,c}} + \frac{|f_s|}{f_{s,c}} + \frac{|\tau_b|}{\tau_{b,c}} + \frac{|\tau_t|}{\tau_{t,c}}\right\}$$

where  $|f_s|$  is the magnitude of the shear force and  $|\tau_b|$  and  $|\tau_t|$  are the magnitudes of the bending and twisting forces, respectively. The corresponding variables  $f_{r,c}$ ,  $f_{s,c}$ ,  $\tau_{b,c}$ , and  $\tau_{t,c}$  are critical limits to each force or torque. If  $B$  is ever equal to or exceeds one, the bond will break. This is done by setting its type to 0 such that forces and torques are no longer computed.

After computing the base magnitudes of the forces and torques, they can be optionally multiplied by an extra factor  $w$  to smoothly interpolate forces and torques to zero as the bond breaks. This term is calculated as  $w = (1.0 - B^4)$ . This smoothing factor can be added or removed using the *smooth* keyword.

Finally, additional damping forces and torques are applied to the two particles. A force is applied proportional to the difference in the normal velocity of particles using a similar construction as dissipative particle dynamics (*Groot*):

$$F_D = -\gamma_n w (\hat{r} \cdot \vec{v})$$

where  $\gamma_n$  is the damping strength,  $\hat{r}$  is the radial normal vector, and  $\vec{v}$  is the velocity difference between the two particles. Similarly, tangential forces are applied to each atom proportional to the relative differences in sliding velocities with a constant prefactor  $\gamma_s$  (*Wang et al.*) along with their associated torques. The rolling and twisting components of the relative angular velocities of the two atoms are also damped by applying torques with prefactors of  $\gamma_r$  and  $\gamma_t$ , respectively.

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $k_r$  (force/distance units)
- $k_s$  (force/distance units)
- $k_t$  (force\*distance/radians units)
- $k_b$  (force\*distance/radians units)
- $f_{r,c}$  (force units)
- $f_{s,c}$  (force units)
- $\tau_{b,c}$  (force\*distance units)



- $\tau_{t,c}$  (force\*distance units)
- $\gamma_n$  (force/velocity units)
- $\gamma_s$  (force/velocity units)
- $\gamma_r$  (force\*distance/velocity units)
- $\gamma_t$  (force\*distance/velocity units)

By default, pair forces are not calculated between bonded particles. Pair forces can alternatively be overlaid on top of bond forces using the *overlay/pair* keyword. These settings require specific *special\_bonds* settings described in the restrictions. Further details can be found in the *:doc: how to <Howto\_BPM>* page on BPMs.

If the *store/local* keyword is used, this fix will track bonds that break during the simulation. Whenever a bond breaks, data is processed and transferred to an internal fix labeled *fix\_ID*. This allows the local data to be accessed by other LAMMPS commands. Following any optional keyword/value arguments, a list of one or more attributes is specified. These include the IDs of the two atoms in the bond. The other attributes for the two atoms include the timestep during which the bond broke and the current/initial center of mass position of the two atoms.

Data is continuously accumulated over intervals of  $N$  timesteps. At the end of each interval, all of the saved accumulated data is deleted to make room for new data. Individual datum may therefore persist anywhere between 1 to  $N$  timesteps depending on when they are saved. This data can be accessed using the *fix\_ID* and a *dump local* command. To ensure all data is output, the dump frequency should correspond to the same interval of  $N$  timesteps. A dump frequency of an integer multiple of  $N$  can be used to regularly output a sample of the accumulated data.

Note that when unbroken bonds are dumped to a file via the *dump local* command, bonds with type 0 (broken bonds) are not included. The *delete\_bonds* command can also be used to query the status of broken bonds or permanently delete them, e.g.:

```
delete_bonds all stats
delete_bonds all bond 0 remove
```

## 20.1.4 Restart and other info

This bond style writes the reference state of each bond to *binary restart files*. Loading a restart file will properly resume bonds.

The *single()* function of these pair styles returns 0.0 for the energy of a pairwise interaction, since energy is not conserved in these dissipative potentials. It also returns only the normal component of the pairwise interaction force.

The accumulated data is not written to restart files and should be output before a restart file is written to avoid missing data.

The internal fix calculates a local vector or local array depending on the number of input values. The length of the vector or number of rows in the array is the number of recorded, lost interactions. If a single input is specified, a local vector is produced. If two or more inputs are specified, a local array is produced where the number of columns = the number of inputs. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector or array will be floating point values that correspond to the specified attribute.

### 20.1.5 Restrictions

This bond style can only be used if LAMMPS was built with the BPM package. See the *Build package* doc page for more info.

By default if pair interactions are to be disabled, this bond style requires setting

```
special_bonds lj 0 1 1 coul 1 1 1
```

and *newton* must be set to bond off. If the *overlay/pair* option is used, this bond style alternatively requires setting

```
special_bonds lj/coul 1 1 1
```

The *bpm/rotational* style requires *atom style bpm/sphere*.

### 20.1.6 Related commands

*bond\_coeff*, *fix nve/bpm/sphere*

### 20.1.7 Default

The option defaults are *smooth = yes*

---

**(Wang)** Wang, Acta Geotechnica, 4, p 117-127 (2009).

**(Wang and Mora)** Wang, Mora, Advances in Geocomputing, 119, p 183-228 (2009).

**(Groot)** Groot and Warren, J Chem Phys, 107, 4423-35 (1997).

**(Wang et al, 2015)** Wang, Y., Alonso-Marroquin, F., & Guo, W. W. (2015). Rolling and sliding in 3-D discrete element models. Particuology, 23, 49-55.

## 20.2 bond\_style bpm/spring command

### 20.2.1 Syntax

```
bond_style bpm/spring keyword value attribute1 attribute2 ...
```

- optional keyword = *overlay/pair* or *store/local* or *smooth*

*store/local* values = *fix\_ID* *N* *attributes* ...

\* *fix\_ID* = ID of associated internal fix to store data

\* *N* = prepare data for output every this many timesteps

\* *attributes* = zero or more of the below attributes may be appended

*id1*, *id2* = IDs of 2 atoms in the bond

*time* = the timestep the bond broke

*x*, *y*, *z* = the center of mass position of the 2 atoms when the bond broke.↵

↵(distance units)

*x/ref*, *y/ref*, *z/ref* = the initial center of mass position of the 2 atoms.↵

↵(distance units)

`overlay/pair` value = none  
 bonded particles will still interact with pair forces

`smooth` value = yes or no  
 smooths bond forces near the breaking point

## 20.2.2 Examples

```
bond_style bpm/spring
bond_coeff 1 1.0 0.05 0.1

bond_style bpm/spring myfix 1000 time id1 id2
dump 1 all local 1000 dump.broken f_myfix[1] f_myfix[2] f_myfix[3]
dump_modify 1 write_header no
```

## 20.2.3 Description

The *bpm/spring* bond style computes forces and torques based on deviations from the initial reference state of the two atoms. The reference state is stored by each bond when it is first computed in the setup of a run. Data is then preserved across run commands and is written to *binary restart files* such that restarting the system will not reset the reference state of a bond.

This bond style only applies central-body forces which conserve the translational and rotational degrees of freedom of a bonded set of particles. The force has a magnitude of

$$F = k(r - r_0)w$$

where  $k_r$  is a stiffness,  $r$  is the current distance and  $r_0$  is the initial distance between the two particles, and  $w$  is an optional smoothing factor discussed below. Bonds will break at a strain of  $\epsilon_c$ . This is done by setting its type to 0 such that forces are no longer computed.

An additional damping force is applied to the bonded particles. This force is proportional to the difference in the normal velocity of particles using a similar construction as dissipative particle dynamics (*Groot*):

$$F_D = -\gamma w(\hat{r} \bullet \vec{v})$$

where  $\gamma$  is the damping strength,  $\hat{r}$  is the radial normal vector, and  $\vec{v}$  is the velocity difference between the two particles.

The smoothing factor  $w$  can be added or removed using the *smooth* keyword. It is constructed such that forces smoothly go to zero, avoiding discontinuities, as bonds approach the critical strain

$$w = 1.0 - \left( \frac{r - r_0}{r_0 \epsilon_c} \right)^8.$$

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $k$  (force/distance units)
- $\epsilon_c$  (unit less)
- $\gamma$  (force/velocity units)

By default, pair forces are not calculated between bonded particles. Pair forces can alternatively be overlaid on top of bond forces using the *overlay/pair* keyword. These settings require specific *special\_bonds* settings described in the restrictions. Further details can be found in the *:doc: how to <Howto\_BPM>* page on BPMs.

If the *store/local* keyword is used, this fix will track bonds that break during the simulation. Whenever a bond breaks, data is processed and transferred to an internal fix labeled *fix\_ID*. This allows the local data to be accessed by other LAMMPS commands. Following any optional keyword/value arguments, a list of one or more attributes is specified. These include the IDs of the two atoms in the bond. The other attributes for the two atoms include the timestep during which the bond broke and the current/initial center of mass position of the two atoms.

Data is continuously accumulated over intervals of  $N$  timesteps. At the end of each interval, all of the saved accumulated data is deleted to make room for new data. Individual datum may therefore persist anywhere between 1 to  $N$  timesteps depending on when they are saved. This data can be accessed using the *fix\_ID* and a *dump local* command. To ensure all data is output, the dump frequency should correspond to the same interval of  $N$  timesteps. A dump frequency of an integer multiple of  $N$  can be used to regularly output a sample of the accumulated data.

Note that when unbroken bonds are dumped to a file via the *dump local* command, bonds with type 0 (broken bonds) are not included. The *delete\_bonds* command can also be used to query the status of broken bonds or permanently delete them, e.g.:

```
delete_bonds all stats
delete_bonds all bond 0 remove
```

---

## 20.2.4 Restart and other info

This bond style writes the reference state of each bond to *binary restart files*. Loading a restart file will properly resume bonds.

The `single()` function of these pair styles returns 0.0 for the energy of a pairwise interaction, since energy is not conserved in these dissipative potentials.

The accumulated data is not written to restart files and should be output before a restart file is written to avoid missing data.

The internal fix calculates a local vector or local array depending on the number of input values. The length of the vector or number of rows in the array is the number of recorded, lost interactions. If a single input is specified, a local vector is produced. If two or more inputs are specified, a local array is produced where the number of columns = the number of inputs. The vector or array can be accessed by any command that uses local values from a compute as input. See the *Howto output* page for an overview of LAMMPS output options.

The vector or array will be floating point values that correspond to the specified attribute.

## 20.2.5 Restrictions

This bond style can only be used if LAMMPS was built with the BPM package. See the *Build package* doc page for more info.

By default if pair interactions are to be disabled, this bond style requires setting

```
special_bonds lj 0 1 1 coul 1 1 1
```

and *newton* must be set to bond off. If the *overlay/pair* option is used, this bond style alternatively requires setting

```
special_bonds lj/coul 1 1 1
```

## 20.2.6 Related commands

*bond\_coeff*, *pair bpm/spring*

## 20.2.7 Default

The option defaults are *smooth = yes*

---

(Groot) Groot and Warren, J Chem Phys, 107, 4423-35 (1997).

## 20.3 bond\_style class2 command

Accelerator Variants: *class2/omp*, *class2/kk*

### 20.3.1 Syntax

```
bond_style class2
```

### 20.3.2 Examples

```
bond_style class2
bond_coeff 1 1.0 100.0 80.0 80.0
```

### 20.3.3 Description

The *class2* bond style uses the potential

$$E = K_2(r - r_0)^2 + K_3(r - r_0)^3 + K_4(r - r_0)^4$$

where  $r_0$  is the equilibrium bond distance.

See ([Sun](#)) for a description of the COMPASS class2 force field.

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $r_0$  (distance)
- $K_2$  (energy/distance<sup>2</sup>)
- $K_3$  (energy/distance<sup>3</sup>)
- $K_4$  (energy/distance<sup>4</sup>)

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 20.3.4 Restrictions

This bond style can only be used if LAMMPS was built with the CLASS2 package. See the [Build package](#) page for more info.

### 20.3.5 Related commands

*bond\_coeff*, *delete\_bonds*

### 20.3.6 Default

none

---

(Sun) Sun, J Phys Chem B 102, 7338-7364 (1998).

## 20.4 `bond_style fene` command

Accelerator Variants: *fene/intel*, *fene/kk*, *fene/omp*

## 20.5 `bond_style fene/nm` command

### 20.5.1 Syntax

```
bond_style fene  
bond_style fene/nm
```

### 20.5.2 Examples

```
bond_style fene  
bond_coeff 1 30.0 1.5 1.0 1.0  
  
bond_style fene/nm  
bond_coeff 1 2.25344 1.5 1.0 1.12246 2 6
```

### 20.5.3 Description

The *fene* bond style uses the potential

$$E = -0.5KR_0^2 \ln \left[ 1 - \left( \frac{r}{R_0} \right)^2 \right] + 4\epsilon \left[ \left( \frac{\sigma}{r} \right)^{12} - \left( \frac{\sigma}{r} \right)^6 \right] + \epsilon$$

to define a finite extensible nonlinear elastic (FENE) potential (*Kremer*), used for bead-spring polymer models. The first term is attractive, the second Lennard-Jones term is repulsive. The first term extends to  $R_0$ , the maximum extent of the bond. The second term is cutoff at  $2^{\frac{1}{6}}\sigma$ , the minimum of the LJ potential.

The *fene/nm* bond style substitutes the standard LJ potential with the generalized LJ potential in the same form as in pair style *nm/cut*. The bond energy is then given by

$$E = -0.5Kr_0^2 \ln \left[ 1 - \left( \frac{r}{R_0} \right)^2 \right] + \frac{E_0}{(n-m)} \left[ m \left( \frac{r_0}{r} \right)^n - n \left( \frac{r_0}{r} \right)^m \right]$$

Similar to the *fene* style, the generalized Lennard-Jones is cut off at the potential minimum,  $r_0$ , to be repulsive only. The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy/distance<sup>2</sup>)
- $R_0$  (distance)
- $\epsilon$  (energy)
- $\sigma$  (distance)

For the *fene/nm* style, the following coefficients are used. Please note, that the standard LJ potential and thus the regular FENE potential is recovered for ( $n=12$   $m=6$ ) and  $r_0 = 2^{\frac{1}{6}}\sigma$ .

- $K$  (energy/distance<sup>2</sup>)
- $R_0$  (distance)
- $E_0$  (energy)
- $r_0$  (distance)
- $n$  (unitless)
- $m$  (unitless)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

## 20.5.4 Restrictions

The *fene* bond style can only be used if LAMMPS was built with the MOLECULE package; the *fene/nm* bond style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the [Build package](#) page for more info.

You typically should specify *special\_bonds fene* or *special\_bonds lj/coul 0 1 1* to use this bond style. LAMMPS will issue a warning if that's not the case.

## 20.5.5 Related commands

*bond\_coeff*, *delete\_bonds*, *pair style lj/cut*, *pair style nm/cut*.

## 20.5.6 Default

none

---

**(Kremer)** Kremer, Grest, J Chem Phys, 92, 5057 (1990).

## 20.6 bond\_style fene/expand command

Accelerator Variants: *fene/expand/omp*

### 20.6.1 Syntax

```
bond_style fene/expand
```

### 20.6.2 Examples

```
bond_style fene/expand  
bond_coeff 1 30.0 1.5 1.0 1.0 0.5
```

### 20.6.3 Description

The *fene/expand* bond style uses the potential

$$E = -0.5KR_0^2 \ln \left[ 1 - \left( \frac{r - \Delta}{R_0} \right)^2 \right] + 4\epsilon \left[ \left( \frac{\sigma}{r - \Delta} \right)^{12} - \left( \frac{\sigma}{r - \Delta} \right)^6 \right] + \epsilon$$

to define a finite extensible nonlinear elastic (FENE) potential (*Kremer*), used for bead-spring polymer models. The first term is attractive, the second Lennard-Jones term is repulsive.

The *fene/expand* bond style is similar to *fene* except that an extra shift factor of  $\Delta$  (positive or negative) is added to  $r$  to effectively change the bead size of the bonded atoms. The first term now extends to  $R_0 + \Delta$  and the second term is cutoff at  $2^{\frac{1}{6}}\sigma + \Delta$ .

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:



- $K$  (energy/distance<sup>2</sup>)
  - $R_0$  (distance)
  - $\epsilon$  (energy)
  - $\sigma$  (distance)
  - $\Delta$  (distance)
- 

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 20.6.4 Restrictions

This bond style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) page for more info.

You typically should specify *special\_bonds fene* or *special\_bonds lj/coul 0 1 1* to use this bond style. LAMMPS will issue a warning if that's not the case.

## 20.6.5 Related commands

*bond\_coeff*, *delete\_bonds*

## 20.6.6 Default

none

---

(**Kremer**) Kremer, Grest, J Chem Phys, 92, 5057 (1990).

# 20.7 bond\_style gaussian command

## 20.7.1 Syntax

```
bond_style gaussian
```

---

## 20.7.2 Examples

```
bond_style gaussian
bond_coeff 1 300.0 2 0.0128 0.375 3.37 0.0730 0.148 3.63
```

## 20.7.3 Description

The *gaussian* bond style uses the potential:

$$E = -k_B T \ln \left( \sum_{i=1}^n \frac{A_i}{w_i \sqrt{\pi/2}} \exp \left( -\frac{(r - r_i)^2}{w_i^2} \right) \right)$$

This analytical form is a suitable potential for obtaining mesoscale effective force fields which can reproduce target atomistic distributions (*Milano*)

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- T temperature at which the potential was derived
- n (integer >=1)
- A<sub>1</sub> (-)
- w<sub>1</sub> (-)
- r<sub>1</sub> (length)
- ...
- A<sub>n</sub> (-)
- w<sub>n</sub> (-)
- r<sub>n</sub> (length)

## 20.7.4 Restrictions

This bond style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

## 20.7.5 Related commands

*bond\_coeff*

## 20.7.6 Default

none

---

(*Milano*) G. Milano, S. Goudeau, F. Mueller-Plathe, J. Polym. Sci. B Polym. Phys. 43, 871 (2005).

## 20.8 bond\_style gromos command

Accelerator Variants: *gromos/omp*

### 20.8.1 Syntax

```
bond_style gromos
```

### 20.8.2 Examples

```
bond_style gromos  
bond_coeff 5 80.0 1.2
```

### 20.8.3 Description

The *gromos* bond style uses the potential

$$E = K(r^2 - r_0^2)^2$$

where  $r_0$  is the equilibrium bond distance. Note that the usual 1/4 factor is included in  $K$ .

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy/distance<sup>4</sup>)
- $r_0$  (distance)

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 20.8.4 Restrictions

This bond style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* page for more info.

## 20.8.5 Related commands

*bond\_coeff*, *delete\_bonds*

## 20.8.6 Default

none

## 20.9 bond\_style harmonic command

Accelerator Variants: *harmonic/intel*, *harmonic/kk*, *harmonic/omp*

### 20.9.1 Syntax

```
bond_style harmonic
```

### 20.9.2 Examples

```
bond_style harmonic  
bond_coeff 5 80.0 1.2
```

### 20.9.3 Description

The *harmonic* bond style uses the potential

$$E = K(r - r_0)^2$$

where  $r_0$  is the equilibrium bond distance. Note that the usual 1/2 factor is included in  $K$ .

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy/distance<sup>2</sup>)
- $r_0$  (distance)

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

## 20.9.4 Restrictions

This bond style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* page for more info.

## 20.9.5 Related commands

*bond\_coeff*, *delete\_bonds*

## 20.9.6 Default

none

## 20.10 bond\_style harmonic/shift command

Accelerator Variants: *harmonic/shift/omp*

### 20.10.1 Syntax

```
bond_style harmonic/shift
```

### 20.10.2 Examples

```
bond_style harmonic/shift
bond_coeff 5 10.0 0.5 1.0
```

### 20.10.3 Description

The *harmonic/shift* bond style is a shifted harmonic bond that uses the potential

$$E = \frac{U_{\min}}{(r_0 - r_c)^2} [(r - r_0)^2 - (r_c - r_0)^2]$$

where  $r_0$  is the equilibrium bond distance, and  $r_c$  the critical distance. The potential is  $-U_{\min}$  at  $r_0$  and zero at  $r_c$ . The spring constant is  $k = U_{\min}/[2(r_0 - r_c)^2]$ .

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $U_{\min}$  (energy)

- $r_0$  (distance)
  - $r_c$  (distance)
- 

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 20.10.4 Restrictions

This bond style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the [Build package](#) doc page for more info.

## 20.10.5 Related commands

*bond\_coeff*, *delete\_bonds*, *bond\_harmonic*

## 20.10.6 Default

none

## 20.11 bond\_style harmonic/shift/cut command

Accelerator Variants: *harmonic/shift/cut/omp*

### 20.11.1 Syntax

```
bond_style harmonic/shift/cut
```

### 20.11.2 Examples

```
bond_style harmonic/shift/cut  
bond_coeff 5 10.0 0.5 1.0
```

### 20.11.3 Description

The *harmonic/shift/cut* bond style is a shifted harmonic bond that uses the potential

$$E = \frac{U_{\min}}{(r_0 - r_c)^2} [(r - r_0)^2 - (r_c - r_0)^2]$$

where  $r_0$  is the equilibrium bond distance, and  $r_c$  the critical distance. The bond potential is zero for distances  $r > r_c$ . The potential is  $-U_{\min}$  at  $r_0$  and zero at  $r_c$ . The spring constant is  $k = U_{\min}/[2(r_0 - r_c)^2]$ .

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $U_{\min}$  (energy)
- $r_0$  (distance)
- $r_c$  (distance)

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 20.11.4 Restrictions

This bond style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

### 20.11.5 Related commands

*bond\_coeff*, *delete\_bonds*, *bond\_harmonic*, *bond\_style harmonic/shift*

### 20.11.6 Default

none

## 20.12 bond\_style hybrid command

### 20.12.1 Syntax

```
bond_style hybrid style1 style2 ...
```

- style1,style2 = list of one or more bond styles

### 20.12.2 Examples

### 20.12.3 Description

The *hybrid* style enables the use of multiple bond styles in one simulation. A bond style is assigned to each bond type. For example, bonds in a polymer flow (of bond type 1) could be computed with a *fene* potential and bonds in the wall boundary (of bond type 2) could be computed with a *harmonic* potential. The assignment of bond type to style is made via the *bond\_coeff* command or in the data file.

In the *bond\_coeff* commands, the name of a bond style must be added after the bond type, with the remaining coefficients being those appropriate to that style. In the example above, the 2 *bond\_coeff* commands set bonds of bond type 1 to be computed with a *harmonic* potential with coefficients 80.0, 1.2 for  $K, r_0$ . All other bond types (2-N) are computed with a *fene* potential with coefficients 30.0, 1.5, 1.0, 1.0 for  $K, R_0, \epsilon, \sigma$ .

If bond coefficients are specified in the data file read via the *read\_data* command, then the same rule applies. E.g. “harmonic” or “fene” must be added after the bond type, for each line in the “Bond Coeffs” section, e.g.

```
Bond Coeffs
```

```
1 harmonic 80.0 1.2
2 fene 30.0 1.5 1.0 1.0
...
```

A bond style of *none* with no additional coefficients can be used in place of a bond style, either in a input script *bond\_coeff* command or in the data file, if you desire to turn off interactions for specific bond types.

---

### 20.12.4 Restrictions

This bond style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* page for more info.

Unlike other bond styles, the hybrid bond style does not store bond coefficient info for individual sub-styles in a *binary restart files*. Thus when restarting a simulation from a restart file, you need to re-specify *bond\_coeff* commands.



## 20.12.5 Related commands

*bond\_coeff*, *delete\_bonds*

## 20.12.6 Default

none

## 20.13 bond\_style mesocnt command

### 20.13.1 Syntax

```
bond_style mesocnt
```

### 20.13.2 Examples

```
bond_style mesocnt
bond_coeff 1 C 10 10 20.0
bond_coeff 4 custom 800.0 10.0
```

### 20.13.3 Description

New in version 15Sep2022.

The *mesocnt* bond style is a wrapper for the *harmonic* style, and uses the potential

$$E = K(r - r_0)^2$$

where  $r_0$  is the equilibrium bond distance. Note that the usual 1/2 factor is included in  $K$ . The style implements parameterization presets of  $K$  for mesoscopic simulations of carbon nanotubes based on the atomistic simulations of (*Srivastava*).

Other presets can be readily implemented in the future.

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- preset = *C* or *custom*
- additional parameters depending on preset

Preset *C* is for carbon nanotubes, and the additional parameters are:

- chiral index  $n$  (unitless)
- chiral index  $m$  (unitless)
- $r_0$  (distance)

Preset *custom* is simply a direct wrapper for the *harmonic* style, and the additional parameters are:

- $K$  (energy/distance<sup>2</sup>)
- $r_0$  (distance)

### 20.13.4 Restrictions

This bond style can only be used if LAMMPS was built with the MOLECULE and MESONT packages. See the *Build package* page for more info.

### 20.13.5 Related commands

*bond\_coeff*, *delete\_bonds*

### 20.13.6 Default

none

---

(Srivastava) Zhigilei, Wei and Srivastava, Phys. Rev. B 71, 165417 (2005).

## 20.14 bond\_style mm3 command

### 20.14.1 Syntax

```
bond_style mm3
```

### 20.14.2 Examples

```
bond_style mm3
bond_coeff 1 100.0 107.0
```

### 20.14.3 Description

The *mm3* bond style uses the potential that is anharmonic in the bond as defined in (*Allinger*)

$$E = K(r - r_0)^2 [1 - 2.55(r - r_0) + (7/12)2.55^2(r - r_0)^2]$$

where  $r_0$  is the equilibrium value of the bond, and  $K$  is a prefactor. The anharmonic prefactors have units  $\text{angstrom}^{-n}$ :  $-2.55 \text{ angstrom}^{-1}$  and  $(7/12)2.55^2 \text{ angstrom}^{-2}$ . The code takes care of the necessary unit conversion for these factors internally. Note that the MM3 papers contains an error in Eq (1):  $(7/12)2.55$  should be replaced with  $(7/12)2.55^2$

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy/distance<sup>2</sup>)
- $r_0$  (distance)

## 20.14.4 Restrictions

This bond style can only be used if LAMMPS was built with the YAFF package. See the *Build package* doc page for more info.

## 20.14.5 Related commands

*bond\_coeff*

## 20.14.6 Default

none

---

(Allinger) Allinger, Yuh, Lii, JACS, 111(23), 8551-8566 (1989),

## 20.15 bond\_style morse command

Accelerator Variants: *morse/omp*

### 20.15.1 Syntax

```
bond_style morse
```

### 20.15.2 Examples

```
bond_style morse
bond_coeff 5 1.0 2.0 1.2
```

### 20.15.3 Description

The *morse* bond style uses the potential

$$E = D \left[ 1 - e^{-\alpha(r-r_0)} \right]^2$$

where  $r_0$  is the equilibrium bond distance,  $\alpha$  is a stiffness parameter, and  $D$  determines the depth of the potential well.

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $D$  (energy)
- $\alpha$  (inverse distance)
- $r_0$  (distance)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 20.15.4 Restrictions

This bond style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) page for more info.

## 20.15.5 Related commands

*bond\_coeff*, *delete\_bonds*

## 20.15.6 Default

none

## 20.16 bond\_style none command

### 20.16.1 Syntax

```
bond_style none
```

### 20.16.2 Examples

```
bond_style none
```

### 20.16.3 Description

Using a bond style of none means bond forces and energies are not computed, even if pairs of bonded atoms were listed in the data file read by the *read\_data* command.

See the *bond\_style zero* command for a way to calculate bond statistics, but compute no bond interactions.

### 20.16.4 Restrictions

none

### 20.16.5 Related commands

none

*bond\_style zero*

### 20.16.6 Default

none

## 20.17 bond\_style nonlinear command

Accelerator Variants: *nonlinear/omp*

### 20.17.1 Syntax

```
bond_style nonlinear
```

### 20.17.2 Examples

```
bond_style nonlinear
bond_coeff 2 100.0 1.1 1.4
```

### 20.17.3 Description

The *nonlinear* bond style uses the potential

$$E = \frac{\epsilon(r - r_0)^2}{[\lambda^2 - (r - r_0)^2]}$$

to define an anharmonic spring (*Rector*) of equilibrium length  $r_0$  and maximum extension lamda.

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $\epsilon$  (energy)
- $r_0$  (distance)

- $\lambda$  (distance)
- 

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 20.17.4 Restrictions

This bond style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the [Build package](#) page for more info.

## 20.17.5 Related commands

*bond\_coeff*, *delete\_bonds*

## 20.17.6 Default

none

---

(Rector) Rector, Van Swol, Henderson, Molecular Physics, 82, 1009 (1994).

## 20.18 bond\_style oxdna/fene command

## 20.19 bond\_style oxdna2/fene command

## 20.20 bond\_style oxrna2/fene command

### 20.20.1 Syntax

```
bond_style oxdna/fene
```

```
bond_style oxdna2/fene
```

```
bond_style oxrna2/fene
```

## 20.20.2 Examples

```

bond_style oxdna/fene
bond_coeff * 2.0 0.25 0.7525

bond_style oxdna2/fene
bond_coeff * 2.0 0.25 0.7564

bond_style oxrna2/fene
bond_coeff * 2.0 0.25 0.76107

```

## 20.20.3 Description

The *oxdna/fene*, *oxdna2/fene*, and *oxrna2/fene* bond styles use the potential

$$E = -\frac{\epsilon}{2} \ln \left[ 1 - \left( \frac{r - r_0}{\Delta} \right)^2 \right]$$

to define a modified finite extensible nonlinear elastic (FENE) potential (*Ouldridge*) to model the connectivity of the phosphate backbone in the oxDNA/oxRNA force field for coarse-grained modelling of DNA/RNA.

The following coefficients must be defined for the bond type via the *bond\_coeff* command as given in the above example, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $\epsilon$  (energy)
- $\Delta$  (distance)
- $r_0$  (distance)

---

**Note:** The oxDNA bond style has to be used together with the corresponding oxDNA pair styles for excluded volume interaction *oxdna/excv*, stacking *oxdna/stk*, cross-stacking *oxdna/xstk* and coaxial stacking interaction *oxdna/coaxstk* as well as hydrogen-bonding interaction *oxdna/hbond* (see also documentation of *pair\_style oxdna/excv*). For the oxDNA2 (*Snodin*) bond style the analogous pair styles *oxdna2/excv*, *oxdna2/stk*, *oxdna2/xstk*, *oxdna2/coaxstk*, *oxdna2/hbond* and an additional Debye-Hueckel pair style *oxdna2/dh* have to be defined. The same applies to the oxRNA2 (*Sulc1*) styles. The coefficients in the above example have to be kept fixed and cannot be changed without reparameterizing the entire model.

---



---

**Note:** This bond style has to be used with the *atom\_style hybrid bond ellipsoid oxdna* (see documentation of *atom\_style*). The *atom\_style oxdna* stores the 3'-to-5' polarity of the nucleotide strand, which is set through the bond topology in the data file. The first (second) atom in a bond definition is understood to point towards the 3'-end (5'-end) of the strand.

---

**Warning:** If data files are produced with *write\_data*, then the *newton* command should be set to *newton on* or *newton off on*. Otherwise the data files will not have the same 3'-to-5' polarity as the initial data file. This limitation does not apply to binary restart files produced with *write\_restart*.

Example input and data files for DNA and RNA duplexes can be found in examples/PACKAGES/cgdna/examples/oxDNA/, /oxDNA2/ and /oxRNA2/. A simple python setup tool which creates

single straight or helical DNA strands, DNA/RNA duplexes or arrays of DNA/RNA duplexes can be found in [examples/PACKAGES/cgdna/util/](#).

Please cite (*Henrich*) in any publication that uses this implementation. An updated documentation that contains general information on the model, its implementation and performance as well as the structure of the data and input file can be found [here](#).

Please cite also the relevant oxDNA/oxRNA publications. These are (*Ouldridge*) and (*Ouldridge-DPhil*) for oxDNA, (*Snodin*) for oxDNA2, (*Sulc1*) for oxRNA2 and for sequence-specific hydrogen-bonding and stacking interactions (*Sulc2*).

---

## 20.20.4 Restrictions

This bond style can only be used if LAMMPS was built with the CG-DNA package and the MOLECULE and ASPHERE package. See the *Build package* page for more info.

## 20.20.5 Related commands

*pair\_style oxdna/excv*, *pair\_style oxdna2/excv*, *pair\_style oxrna2/excv*, *bond\_coeff*, *atom\_style oxdna*, *fix nve/dotc/langevin*

## 20.20.6 Default

none

---

**(Henrich)** O. Henrich, Y. A. Gutierrez-Fosado, T. Curk, T. E. Ouldridge, Eur. Phys. J. E 41, 57 (2018).

**(Ouldridge-DPhil)** T.E. Ouldridge, Coarse-grained modelling of DNA and DNA self-assembly, DPhil. University of Oxford (2011).

**(Ouldridge)** T.E. Ouldridge, A.A. Louis, J.P.K. Doye, J. Chem. Phys. 134, 085101 (2011).

**(Snodin)** B.E. Snodin, F. Randisi, M. Mosayebi, et al., J. Chem. Phys. 142, 234901 (2015).

**(Sulc1)** P. Sulc, F. Romano, T. E. Ouldridge, et al., J. Chem. Phys. 140, 235102 (2014).

**(Sulc2)** P. Sulc, F. Romano, T.E. Ouldridge, L. Rovigatti, J.P.K. Doye, A.A. Louis, J. Chem. Phys. 137, 135101 (2012).

## 20.21 `bond_style quartic` command

Accelerator Variants: *quartic/omp*



## 20.21.1 Syntax

```
bond_style quartic
```

## 20.21.2 Examples

```
bond_style quartic
bond_coeff 2 1200 -0.55 0.25 1.3 34.6878
```

## 20.21.3 Description

The *quartic* bond style uses the potential

$$E = K(r - R_c)^2(r - R_c - B_1)(r - R_c - B_2) + U_0 + 4\epsilon \left[ \left(\frac{\sigma}{r}\right)^{12} - \left(\frac{\sigma}{r}\right)^6 \right] + \epsilon$$

to define a bond that can be broken as the simulation proceeds (e.g. due to a polymer being stretched). The  $\sigma$  and  $\epsilon$  used in the LJ portion of the formula are both set equal to 1.0 by LAMMPS.

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy/distance<sup>4</sup>)
- $B_1$  (distance)
- $B_2$  (distance)
- $R_c$  (distance)
- $U_0$  (energy)

This potential was constructed to mimic the FENE bond potential for coarse-grained polymer chains. When monomers with  $\sigma = \epsilon = 1.0$  are used, the following choice of parameters gives a quartic potential that looks nearly like the FENE potential:

$$\begin{aligned} K &= 1200 \\ B_1 &= -0.55 \\ B_2 &= 0.25 \\ R_c &= 1.3 \\ U_0 &= 34.6878 \end{aligned}$$

Different parameters can be specified using the *bond\_coeff* command, but you will need to choose them carefully so they form a suitable bond potential.

$R_c$  is the cutoff length at which the bond potential goes smoothly to a local maximum. If a bond length ever becomes  $> R_c$ , LAMMPS “breaks” the bond, which means two things. First, the bond potential is turned off by setting its type to 0, and is no longer computed. Second, a pairwise interaction between the two atoms is turned on, since they are no longer bonded. See the *Howto* page on broken bonds for more information.

LAMMPS does the second task via a computational sleight-of-hand. It subtracts the pairwise interaction as part of the bond computation. When the bond breaks, the subtraction stops. For this to work, the pairwise interaction must always be computed by the *pair\_style* command, whether the bond is broken or not. This means that *special\_bonds* must be set to 1,1,1, as indicated as a restriction below.

Note that when bonds are dumped to a file via the *dump local* command, bonds with type 0 are not included. The *delete\_bonds* command can also be used to query the status of broken bonds or permanently delete them, e.g.:

```
delete_bonds all stats
delete_bonds all bond 0 remove
```

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 20.21.4 Restrictions

This bond style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) page for more info.

The *quartic* style requires that *special\_bonds* parameters be set to 1,1,1. Three- and four-body interactions (angle, dihedral, etc) cannot be used with *quartic* bonds.

## 20.21.5 Related commands

*bond\_coeff*, *delete\_bonds*

## 20.21.6 Default

none

## 20.22 bond\_style special command

### 20.22.1 Syntax

```
bond_style special
```

## 20.22.2 Examples

```
bond_style special
bond_coeff 0.5 0.5
```

## 20.22.3 Description

The *special* bond style can be used to create conceptual bonds which effectively impose weightings on the pairwise Lennard Jones and/or Coulombic interactions between selected pairs of particles in the system. The form of the pairwise interaction will be whatever is computed by the *pair\_style* command defined for the system; this command defines the weightings for its two terms.

This command can thus be useful to apply weightings that cannot be handled by the *special\_bonds* command, such as on 1-5 or 1-6 interactions. Or it can be used to add pairwise forces between one or more pairs of atoms that otherwise would not be include in the *pair\_style* computation.

The potential for this bond style has the form

$$E = w_{LJ}E_{LJ} + w_{Coul}E_{Coul}$$

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $w_{LJ}$  weight (0.0 to 1.0) on pairwise Lennard-Jones interactions
- $w_{Coul}$  weight (0.0 to 1.0) on pairwise Coulombic interactions

Normally this bond style should be used in conjunction with one (or more) other bond styles which compute forces between atoms directly bonded to each other in a molecule. This means the *bond\_style hybrid* command should be used with *bond\_style special* as one of its sub-styles.

Note that the same as for any other bond style, pairs of bonded atoms must be enumerated in the data file read by the *read\_data* command. Thus if this command is used to weight all 1-5 interactions in the system, all the 1-5 pairs of atoms must be listed in the “Bonds” section of the data file.

This bond style imposes strict requirements on settings made with the *special\_bonds* command. These requirements ensure that the new bonds created by this style do not create spurious 1-2, 1-3, or 1-4 interactions within the molecular topology.

Specifically 1-2 interactions must have weights of zero, 1-3 interactions must either have weights of unity or *special\_bonds angle yes* must be used, and 1-4 interactions must have weights of unity or *special\_bonds dihedral yes* must be used.

If this command is used to create bonded interactions between particles that are further apart than usual (e.g. 1-5 or 1-6 interactions), this style may require an increase in the communication cutoff via the *comm\_modify cutoff* command. If LAMMPS cannot find a partner atom in a bond, an error will be issued.

### 20.22.4 Restrictions

This bond style can only be used if LAMMPS was built with the MISC package. See the *Build package* doc page for more info.

This bond style requires the use of a *pair\_style* which computes a pairwise additive interaction and provides the ability to compute interactions for individual pairs of atoms. Manybody potentials are not compatible in general, but also some other pair styles are missing the required functionality and thus will cause an error.

This command is not compatible with long-range Coulombic interactions. If a *kspace\_style* <*kspace\_style*> is declared, an error will be issued.

### 20.22.5 Related commands

*bond\_coeff*, *special\_bonds*

### 20.22.6 Default

none

## 20.23 bond\_style table command

Accelerator Variants: *table/omp*

### 20.23.1 Syntax

```
bond_style table style N
```

- style = *linear* or *spline* = method of interpolation
- N = use N values in table

### 20.23.2 Examples

```
bond_style table linear 1000  
bond_coeff 1 file.table ENTRY1
```

### 20.23.3 Description

Style *table* creates interpolation tables of length *N* from bond potential and force values listed in a file(s) as a function of bond length. The files are read by the *bond\_coeff* command.

The interpolation tables are created by fitting cubic splines to the file values and interpolating energy and force values at each of *N* distances. During a simulation, these tables are used to interpolate energy and force values as needed. The interpolation is done in one of 2 styles: *linear* or *spline*.

For the *linear* style, the bond length is used to find 2 surrounding table values from which an energy or force is computed by linear interpolation.

For the *spline* style, a cubic spline coefficients are computed and stored at each of the  $N$  values in the table. The bond length is used to find the appropriate set of coefficients which are used to evaluate a cubic polynomial which computes the energy or force.

The following coefficients must be defined for each bond type via the *bond\_coeff* command as in the example above.

- filename
- keyword

The filename specifies a file containing tabulated energy and force values. The keyword specifies a section of the file. The format of this file is described below.

The format of a tabulated file is as follows (without the parenthesized comments):

```
# Bond potential for harmonic (one or more comment or blank lines)

HAM                               (keyword is the first text on line)
N 101 FP 0 0 EQ 0.5              (N, FP, EQ parameters)
                                (blank line)
1 0.00 338.0000 1352.0000        (index, bond-length, energy, force)
2 0.01 324.6152 1324.9600
...
101 1.00 338.0000 -1352.0000
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The line can contain additional text, but the initial text must match the argument specified in the *bond\_coeff* command. The next line lists (in any order) one or more parameters for the table. Each parameter is a keyword followed by one or more numeric values.

The parameter “N” is required and its value is the number of table entries that follow. Note that this may be different than the  $N$  specified in the *bond\_style table* command. Let  $N_{table} = N$  in the *bond\_style* command, and  $N_{file} = “N”$  in the tabulated file. What LAMMPS does is a preliminary interpolation by creating splines using the  $N_{file}$  tabulated values as nodal points. It uses these to interpolate as needed to generate energy and force values at  $N_{table}$  different points. The resulting tables of length  $N_{table}$  are then used as described above, when computing energy and force for individual bond lengths. This means that if you want the interpolation tables of length  $N_{table}$  to match exactly what is in the tabulated file (with effectively no preliminary interpolation), you should set  $N_{table} = N_{file}$ .

The “FP” parameter is optional. If used, it is followed by two values *fplo* and *fphi*, which are the derivatives of the force at the innermost and outermost bond lengths. These values are needed by the spline construction routines. If not specified by the “FP” parameter, they are estimated (less accurately) by the first two and last two force values in the table.

The “EQ” parameter is also optional. If used, it is followed by a the equilibrium bond length, which is used, for example, by the *fix shake* command. If not used, the equilibrium bond length is to the distance in the table with the lowest potential energy.

Following a blank line, the next  $N$  lines list the tabulated values. On each line, the first value is the index from 1 to  $N$ , the second value is the bond length  $r$  (in distance units), the third value is the energy (in energy units), and the fourth is the force (in force units). The bond lengths must range from a LO value to a HI value, and increase from one line to the next. If the actual bond length is ever smaller than the LO value or larger than the HI value, then the calculation is aborted with an error, so it is advisable to cover the whole range of possible bond lengths.

Note that one file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified keyword.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 20.23.4 Restart, fix\_modify, output, run start/stop, minimize info

This bond style writes the settings for the “bond\_style table” command to *binary restart files*, so a bond\_style command does not need to be specified in an input script that reads a restart file. However, the coefficient information is not stored in the restart file, since it is tabulated in the potential files. Thus, bond\_coeff commands do need to be specified in the restart input script.

## 20.23.5 Restrictions

This bond style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) page for more info.

## 20.23.6 Related commands

*bond\_coeff*, *delete\_bonds*

## 20.23.7 Default

none

## 20.24 bond\_style zero command

### 20.24.1 Syntax

```
bond_style zero keyword
```

- zero or more keywords may be appended
- keyword = *nocoeff*

## 20.24.2 Examples

```
bond_style zero
bond_style zero nocoeff
bond_coeff *
bond_coeff * 2.14
```

## 20.24.3 Description

Using an bond style of zero means bond forces and energies are not computed, but the geometry of bond pairs is still accessible to other commands.

As an example, the *compute bond/local* command can be used to compute distances for the list of pairs of bond atoms listed in the data file read by the *read\_data* command. If no bond style is defined, this command cannot be used.

The optional *nocoeff* flag allows to read data files with a BondCoeff section for any bond style. Similarly, any *bond\_coeff* commands will only be checked for the bond type number and the rest ignored.

Note that the *bond\_coeff* command must be used for all bond types. If specified, there can be only one value, which is going to be used to assign an equilibrium distance, e.g. for use with *fix shake*.

## 20.24.4 Restrictions

none

## 20.24.5 Related commands

*bond\_style none*

## 20.24.6 Default

none





## ANGLE STYLES

### 21.1 angle\_style amoeba command

#### 21.1.1 Syntax

```
angle_style amoeba
```

#### 21.1.2 Examples

```
angle_style amoeba
angle_coeff * 75.0 -25.0 1.0 0.3 0.02 0.003
angle_coeff * ba 3.6551 24.895 1.0119 1.5228
angle_coeff * ub -7.6 1.5537
```

#### 21.1.3 Description

The *amoeba* angle style uses the potential

$$\begin{aligned}
 E &= E_a + E_{ba} + E_{ub} \\
 E_a &= K_2 (\theta - \theta_0)^2 + K_3 (\theta - \theta_0)^3 + K_4 (\theta - \theta_0)^4 + K_5 (\theta - \theta_0)^5 + K_6 (\theta - \theta_0)^6 \\
 E_{ba} &= N_1 (r_{ij} - r_1)(\theta - \theta_0) + N_2 (r_{jk} - r_2)(\theta - \theta_0) \\
 E_{UB} &= K_{ub} (r_{ik} - r_{ub})^2
 \end{aligned}$$

where  $E_a$  is the angle term,  $E_{ba}$  is a bond-angle term,  $E_{UB}$  is a Urey-Bradley bond term,  $\theta_0$  is the equilibrium angle,  $r_1$  and  $r_2$  are the equilibrium bond lengths, and  $r_{ub}$  is the equilibrium Urey-Bradley bond length.

These formulas match how the Tinker MD code performs its angle calculations for the AMOEBA and HIPPO force fields. See the [Howto amoeba](#) page for more information about the implementation of AMOEBA and HIPPO in LAMMPS.

Note that the  $E_a$  and  $E_{ba}$  formulas are identical to those used for the `angle_style class2/p6` command, however there is no bond-bond cross term formula for  $E_{bb}$ . Additionally, there is a  $E_{UB}$  term for a Urey-Bradley bond. It is effectively a harmonic bond between the I and K atoms of angle IJK, even though that bond is not enumerated in the “Bonds” section of the data file.

There are also two ways that Tinker computes the angle  $\theta$  in the  $E_a$  formula. The first is the standard way of treating IJK as an “in-plane” angle. The second is an “out-of-plane” method which Tinker may use if the center atom J in the angle is bonded to one additional atom in addition to I and K. In this case, all 4 atoms are used to compute the  $E_a$  formula, resulting in forces on all 4 atoms. In the Tinker PRM file, these 2 options are denoted by *angle* versus *anglep*

entries in the “Angle Bending Parameters” section of the PRM force field file. The *pflag* coefficient described below selects between the 2 options.

---

Coefficients for the  $E_a$ ,  $E_{bb}$ , and  $E_{ub}$  formulas must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands.

These are the 8 coefficients for the  $E_a$  formula:

- *pflag* = 0 or 1
- *ubflag* = 0 or 1
- $\theta_0$  (degrees)
- $K_2$  (energy)
- $K_3$  (energy)
- $K_4$  (energy)
- $K_5$  (energy)
- $K_6$  (energy)

A *pflag* value of 0 vs 1 selects between the “in-plane” and “out-of-plane” options described above. *Ubflag* is 1 if there is a Urey-Bradley term associated with this angle type, else it is 0.  $\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence the various  $K$  values are effectively energy per radian<sup>2</sup> or radian<sup>3</sup> or radian<sup>4</sup> or radian<sup>5</sup> or radian<sup>6</sup>.

For the  $E_{ba}$  formula, each line in a *angle\_coeff* command in the input script lists 5 coefficients, the first of which is “ba” to indicate they are BondAngle coefficients. In a data file, these coefficients should be listed under a “BondAngle Coeffs” heading and you must leave out the “ba”, i.e. only list 4 coefficients after the angle type.

- ba
- $N_1$  (energy/distance<sup>2</sup>)
- $N_2$  (energy/distance<sup>2</sup>)
- $r_1$  (distance)
- $r_2$  (distance)

The  $\theta_0$  value in the  $E_{ba}$  formula is not specified, since it is the same value from the  $E_a$  formula.

For the  $E_{ub}$  formula, each line in a *angle\_coeff* command in the input script lists 3 coefficients, the first of which is “ub” to indicate they are UreyBradley coefficients. In a data file, these coefficients should be listed under a “UreyBradley Coeffs” heading and you must leave out the “ub”, i.e. only list 2 coefficients after the angle type.

- ub
  - $K_{ub}$  (energy/distance<sup>2</sup>)
  - $r_{ub}$  (distance)
-

### 21.1.4 Restrictions

This angle style can only be used if LAMMPS was built with the AMOEBA package. See the *Build package* doc page for more info.

### 21.1.5 Related commands

*angle\_coeff*

### 21.1.6 Default

none

## 21.2 angle\_style charmm command

Accelerator Variants: *charmm/intel*, *charmm/kk*, *charmm/omp*

### 21.2.1 Syntax

```
angle_style charmm
```

### 21.2.2 Examples

```
angle_style charmm
angle_coeff 1 300.0 107.0 50.0 3.0
```

### 21.2.3 Description

The *charmm* angle style uses the potential

$$E = K(\theta - \theta_0)^2 + K_{ub}(r - r_{ub})^2$$

with an additional Urey-Bradley term based on the distance  $r$  between the first and third atoms in the angle.  $K$ ,  $\theta_0$ ,  $K_{ub}$ , and  $r_{ub}$  are coefficients defined for each angle type.

See (*MacKerell*) for a description of the CHARMM force field.

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $\theta_0$  (degrees)
- $K_{ub}$  (energy/distance<sup>2</sup>)
- $r_{ub}$  (distance)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence  $K$  is effectively energy per radian<sup>2</sup>.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 21.2.4 Restrictions

This angle style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) doc page for more info.

## 21.2.5 Related commands

*angle\_coeff*

## 21.2.6 Default

none

---

(MacKerell) MacKerell, Bashford, Bellott, Dunbrack, Evanseck, Field, Fischer, Gao, Guo, Ha, et al, J Phys Chem, 102, 3586 (1998).

## 21.3 angle\_style class2 command

Accelerator Variants: *class2/kk*, *class2/omp*

## 21.4 angle\_style class2/p6 command

### 21.4.1 Syntax

```
angle_style class2
```

## 21.4.2 Examples

```

angle_style class2
angle_coeff * 75.0 25.0 0.3 0.002
angle_coeff 1 bb 10.5872 1.0119 1.5228
angle_coeff * ba 3.6551 24.895 1.0119 1.5228

```

## 21.4.3 Description

The *class2* angle style uses the potential

$$\begin{aligned}
 E &= E_a + E_{bb} + E_{ba} \\
 E_a &= K_2(\theta - \theta_0)^2 + K_3(\theta - \theta_0)^3 + K_4(\theta - \theta_0)^4 \\
 E_{bb} &= M(r_{ij} - r_1)(r_{jk} - r_2) \\
 E_{ba} &= N_1(r_{ij} - r_1)(\theta - \theta_0) + N_2(r_{jk} - r_2)(\theta - \theta_0)
 \end{aligned}$$

where  $E_a$  is the angle term,  $E_{bb}$  is a bond-bond term, and  $E_{ba}$  is a bond-angle term.  $\theta_0$  is the equilibrium angle and  $r_1$  and  $r_2$  are the equilibrium bond lengths.

See ([Sun](#)) for a description of the COMPASS *class2* force field.

Coefficients for the  $E_a$ ,  $E_{bb}$ , and  $E_{ba}$  formulas must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands.

These are the 4 coefficients for the  $E_a$  formula:

- $\theta_0$  (degrees)
- $K_2$  (energy)
- $K_3$  (energy)
- $K_4$  (energy)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence the various  $K$  are effectively energy per radian<sup>2</sup> or radian<sup>3</sup> or radian<sup>4</sup>.

For the  $E_{bb}$  formula, each line in a *angle\_coeff* command in the input script lists 4 coefficients, the first of which is “bb” to indicate they are BondBond coefficients. In a data file, these coefficients should be listed under a “BondBond Coeffs” heading and you must leave out the “bb”, i.e. only list 3 coefficients after the angle type.

- bb
- $M$  (energy/distance<sup>2</sup>)
- $r_1$  (distance)
- $r_2$  (distance)

For the  $E_{ba}$  formula, each line in a *angle\_coeff* command in the input script lists 5 coefficients, the first of which is “ba” to indicate they are BondAngle coefficients. In a data file, these coefficients should be listed under a “BondAngle Coeffs” heading and you must leave out the “ba”, i.e. only list 4 coefficients after the angle type.

- ba
- $N_1$  (energy/distance)
- $N_2$  (energy/distance)
- $r_1$  (distance)

- $r_2$  (distance)

The  $\theta_0$  value in the  $E_{ba}$  formula is not specified, since it is the same value from the  $E_a$  formula.

---

**Note:** It is important that the order of the I,J,K atoms in each angle listed in the Angles section of the data file read by the `read_data` command be consistent with the order of the  $r_1$  and  $r_2$  BondBond and BondAngle coefficients. This is because the terms in the formulas for  $E_{bb}$  and  $E_{ba}$  will use the I,J atoms to compute  $r_{ij}$  and the J,K atoms to compute  $r_{jk}$ .

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

The *class2/p6* angle style uses the *class2* potential expanded to sixth order:

$$E_a = K_2 (\theta - \theta_0)^2 + K_3 (\theta - \theta_0)^3 + K_4 (\theta - \theta_0)^4 + K_5 (\theta - \theta_0)^5 + K_6 (\theta - \theta_0)^6$$

In this expanded term 6 coefficients for the  $E_a$  formula need to be set:

- $\theta_0$  (degrees)
- $K_2$  (energy)
- $K_3$  (energy)
- $K_4$  (energy)
- $K_5$  (energy)
- $K_6$  (energy)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence the various  $K$  are effectively energy per radian<sup>2</sup> or radian<sup>3</sup> or radian<sup>4</sup> or radian<sup>5</sup> or radian<sup>6</sup>.

The bond-bond and bond-angle terms remain unchanged.

---

#### 21.4.4 Restrictions

This angle style can only be used if LAMMPS was built with the CLASS2 package. For the *class2/p6* style LAMMPS needs to be built with the MOFFF package. See the [Build package](#) doc page for more info.

## 21.4.5 Related commands

*angle\_coeff*

## 21.4.6 Default

none

(Sun) Sun, J Phys Chem B 102, 7338-7364 (1998).

## 21.5 angle\_style cosine command

Accelerator Variants: *cosine/omp*, *cosine/kk*

### 21.5.1 Syntax

```
angle_style cosine
```

### 21.5.2 Examples

```
angle_style cosine
angle_coeff * 75.0
```

### 21.5.3 Description

The *cosine* angle style uses the potential

$$E = K[1 + \cos(\theta)]$$

where  $K$  is defined for each angle type.

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

## 21.5.4 Restrictions

This angle style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

## 21.5.5 Related commands

*angle\_coeff*

## 21.5.6 Default

none

## 21.6 angle\_style cosine/buck6d command

### 21.6.1 Syntax

```
angle_style cosine/buck6d
```

### 21.6.2 Examples

```
angle_style cosine/buck6d
angle_coeff 1 cosine/buck6d 1.978350 4 180.000000
```

### 21.6.3 Description

The *cosine/buck6d* angle style uses the potential

$$E = K [1 + \cos(n\theta - \theta_0)]$$

where  $K$  is the energy constant,  $n$  is the periodic multiplicity and  $\theta_0$  is the equilibrium angle.

The coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands in the following order:

- $K$  (energy)
- $n$
- $\theta_0$  (degrees)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally.

Additional to the cosine term the *cosine/buck6d* angle style computes the short range (vdW) interaction belonging to the *pair\_style buck6d* between the end atoms of the angle. For this reason this angle style only works in combination with the *pair\_style buck6d* styles and needs the *special\_bonds* 1-3 interactions to be weighted 0.0 to prevent double counting.



## 21.6.4 Restrictions

*cosine/buck6d* can only be used in combination with the *pair\_style buck6d* style and with a *special\_bonds* 0.0 weighting of 1-3 interactions.

This angle style can only be used if LAMMPS was built with the MOFFF package. See the *Build package* doc page for more info.

## 21.6.5 Related commands

*angle\_coeff*

## 21.6.6 Default

none

## 21.7 angle\_style cosine/delta command

Accelerator Variants: *cosine/delta/omp*

### 21.7.1 Syntax

```
angle_style cosine/delta
```

### 21.7.2 Examples

```
angle_style cosine/delta
angle_coeff 2*4 75.0 100.0
```

### 21.7.3 Description

The *cosine/delta* angle style uses the potential

$$E = K[1 - \cos(\theta - \theta_0)]$$

where  $\theta_0$  is the equilibrium value of the angle, and  $K$  is a prefactor. Note that the usual 1/2 factor is included in  $K$ .

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $\theta_0$  (degrees)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages*

page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 21.7.4 Restrictions

This angle style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

## 21.7.5 Related commands

*angle\_coeff*, *angle\_style cosine/squared*

## 21.7.6 Default

none

# 21.8 angle\_style cosine/periodic command

Accelerator Variants: *cosine/periodic/omp*

## 21.8.1 Syntax

```
angle_style cosine/periodic
```

## 21.8.2 Examples

```
angle_style cosine/periodic
angle_coeff * 75.0 1 6
```

### 21.8.3 Description

The *cosine/periodic* angle style uses the following potential, which may be particularly used for organometallic systems where  $n = 4$  might be used for an octahedral complex and  $n = 3$  might be used for a trigonal center:

$$E = \frac{2.0}{n^2} * C [1 - B(-1)^n \cos(n\theta)]$$

where  $C$ ,  $B$  and  $n$  are coefficients defined for each angle type.

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $C$  (energy)
- $B = 1$  or  $-1$
- $n = 1, 2, 3, 4, 5$  or  $6$  for periodicity

Note that the prefactor  $C$  is specified as coefficient and not the overall force constant  $K = \frac{2C}{n^2}$ . When  $B = 1$ , it leads to a minimum for the linear geometry. When  $B = -1$ , it leads to a maximum for the linear geometry.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

### 21.8.4 Restrictions

This angle style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

### 21.8.5 Related commands

*angle\_coeff*

### 21.8.6 Default

none

## 21.9 angle\_style cosine/shift command

Accelerator Variants: *cosine/shift/omp*

### 21.9.1 Syntax

```
angle_style cosine/shift
```

### 21.9.2 Examples

```
angle_style cosine/shift  
angle_coeff * 10.0 45.0
```

### 21.9.3 Description

The *cosine/shift* angle style uses the potential

$$E = -\frac{U_{\min}}{2} [1 + \cos(\theta - \theta_0)]$$

where  $\theta_0$  is the equilibrium angle. The potential is bounded between  $-U_{\min}$  and zero. In the neighborhood of the minimum  $E = -U_{\min} + U_{\min}/4(\theta - \theta_0)^2$  hence the spring constant is  $\frac{U_{\min}}{2}$ .

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $U_{\min}$  (energy)
- $\theta$  (angle)

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 21.9.4 Restrictions

This angle style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

## 21.9.5 Related commands

*angle\_coeff*, *angle\_style cosine/shift/exp*

## 21.9.6 Default

none

## 21.10 angle\_style cosine/shift/exp command

Accelerator Variants: *cosine/shift/exp/omp*

### 21.10.1 Syntax

```
angle_style cosine/shift/exp
```

### 21.10.2 Examples

```
angle_style cosine/shift/exp
angle_coeff * 10.0 45.0 2.0
```

### 21.10.3 Description

The *cosine/shift/exp* angle style uses the potential

$$E = -U_{\min} \frac{e^{-aU(\theta, \theta_0)} - 1}{e^a - 1} \quad \text{with} \quad U(\theta, \theta_0) = -0.5 (1 + \cos(\theta - \theta_0))$$

where  $U_{\min}$ ,  $\theta$ , and  $a$  are defined for each angle type.

The potential is bounded between  $[-U_{\min}, 0]$  and the minimum is located at the angle  $\theta_0$ . The  $a$  parameter can be both positive or negative and is used to control the spring constant at the equilibrium.

The spring constant is given by  $k = A \exp(A) U_{\min} / [2(\exp(a) - 1)]$ . For  $a > 3$ ,  $\frac{k}{U_{\min}} = \frac{a}{2}$  to better than 5% relative error. For negative values of the  $a$  parameter, the spring constant is essentially zero, and anharmonic terms takes over. The potential is furthermore well behaved in the limit  $a \rightarrow 0$ , where it has been implemented to linear order in  $a$  for  $a < 0.001$ . In this limit the potential reduces to the cosineshifted potential.

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $U_{min}$  (energy)
- $\theta$  (angle)

- $A$  (real number)
- 

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 21.10.4 Restrictions

This angle style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the [Build package](#) doc page for more info.

## 21.10.5 Related commands

*angle\_coeff*, *angle\_style cosine/shift*, *dihedral\_style cosine/shift/exp*

## 21.10.6 Default

none

## 21.11 angle\_style cosine/squared command

Accelerator Variants: *cosine/squared/omp*

### 21.11.1 Syntax

```
angle_style cosine/squared
```

### 21.11.2 Examples

```
angle_style cosine/squared
angle_coeff 2*4 75.0 100.0
```

### 21.11.3 Description

The *cosine/squared* angle style uses the potential

$$E = K[\cos(\theta) - \cos(\theta_0)]^2$$

, which is commonly used in the *DREIDING* force field, where  $\theta_0$  is the equilibrium value of the angle, and  $K$  is a prefactor. Note that the usual 1/2 factor is included in  $K$ .

See (*Mayo*) for a description of the DREIDING force field.

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $\theta_0$  (degrees)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 21.11.4 Restrictions

This angle style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

### 21.11.5 Related commands

*angle\_coeff*

### 21.11.6 Default

none

---

(**Mayo**) Mayo, Olfason, Goddard III, J Phys Chem, 94, 8897-8909 (1990).

## 21.12 angle\_style cross command

### 21.12.1 Syntax

```
angle_style cross
```

### 21.12.2 Examples

```
angle_style cross  
angle_coeff 1 200.0 100.0 100.0 1.25 1.25 107.0
```

### 21.12.3 Description

The *cross* angle style uses a potential that couples the bond stretches of a bend with the angle stretch of that bend:

$$E = K_{SS} (r_{12} - r_{12,0}) (r_{32} - r_{32,0}) + K_{BS0} (r_{12} - r_{12,0}) (\theta - \theta_0) + K_{BS1} (r_{32} - r_{32,0}) (\theta - \theta_0)$$

where  $r_{12,0}$  is the rest value of the bond length between atom 1 and 2,  $r_{32,0}$  is the rest value of the bond length between atom 3 and 2, and  $\theta_0$  is the rest value of the angle.  $K_{SS}$  is the force constant of the bond stretch-bond stretch term and  $K_{BS0}$  and  $K_{BS1}$  are the force constants of the bond stretch-angle stretch terms.

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K_{SS}$  (energy/distance<sup>2</sup>)
- $K_{BS0}$  (energy/distance)
- $K_{BS1}$  (energy/distance)
- $r_{12,0}$  (distance)
- $r_{32,0}$  (distance)
- $\theta_0$  (degrees)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence the  $K_{BS0}$  and  $K_{BS1}$  are effectively energy/distance per radian.

### 21.12.4 Restrictions

This angle style can only be used if LAMMPS was built with the YAFF package. See the *Build package* doc page for more info.



## 21.12.5 Related commands

*angle\_coeff*

## 21.12.6 Default

none

## 21.13 angle\_style dipole command

Accelerator Variants: *dipole/omp*

### 21.13.1 Syntax

```
angle_style dipole
```

### 21.13.2 Examples

```
angle_style dipole
angle_coeff 6 2.1 180.0
```

### 21.13.3 Description

The *dipole* angle style is used to control the orientation of a dipolar atom within a molecule (*Orsi*). Specifically, the *dipole* angle style restrains the orientation of a point dipole  $\mu_j$  (embedded in atom  $j$ ) with respect to a reference (bond) vector  $r_{ij}^{\vec{}} = r_i^{\vec{}} - r_j^{\vec{}}$ , where  $i$  is another atom of the same molecule (typically,  $i$  and  $j$  are also covalently bonded).

It is convenient to define an angle gamma between the ‘free’ vector  $\mu_j^{\vec{}}$  and the reference (bond) vector  $r_{ij}^{\vec{}}$ :

$$\cos \gamma = \frac{\mu_j^{\vec{}} \cdot r_{ij}^{\vec{}}}{\mu_j r_{ij}}$$

The *dipole* angle style uses the potential:

$$E = K(\cos \gamma - \cos \gamma_0)^2$$

where  $K$  is a rigidity constant and  $\gamma_0$  is an equilibrium (reference) angle.

The torque on the dipole can be obtained by differentiating the potential using the ‘chain rule’ as in appendix C.3 of (*Allen*):

$$\vec{T}_j = \frac{2K(\cos \gamma - \cos \gamma_0)}{\mu_j r_{ij}} r_{ij}^{\vec{}} \times \mu_j^{\vec{}}$$

Example: if  $\gamma_0$  is set to 0 degrees, the torque generated by the potential will tend to align the dipole along the reference direction defined by the (bond) vector  $r_{ij}^{\vec{}}$  (in other words,  $\mu_j^{\vec{}}$  is restrained to point towards atom  $i$ ).

The dipolar torque  $\vec{T}_j$  must be counterbalanced in order to conserve the local angular momentum. This is achieved via an additional force couple generating a torque equivalent to the opposite of  $\vec{T}_j$ :

$$\begin{aligned} -\vec{T}_j &= r_{ij} \times \vec{F}_i \\ \vec{F}_j &= -\vec{F}_i \end{aligned}$$

where  $\vec{F}_i$  and  $\vec{F}_j$  are applied on atoms  $i$  and  $j$ , respectively.

The following coefficients must be defined for each angle type via the `angle_coeff` command as in the example above, or in the data file or restart files read by the `read_data` or `read_restart` commands:

- $K$  (energy)
  - $\gamma_0$  (degrees)
- 

Styles with a `gpu`, `intel`, `kk`, `omp`, or `opt` suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the `-suffix` *command-line switch* when you invoke LAMMPS, or you can use the `suffix` command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

### 21.13.4 Restrictions

This angle style can only be used if LAMMPS was built with the DIPOLE package. See the [Build package](#) doc page for more info.

---

**Note:** In the “Angles” section of the data file, the atom ID  $j$  defining the direction of the dipole vector to restrain must come before the atom ID of the reference atom  $i$ . A third atom ID  $k$  must also be provided to comply with the requirement of a valid angle definition. This atom ID  $k$  should be chosen to be that of an atom bonded to atom  $i$  to avoid errors with “lost angle atoms” when running in parallel. Since the LAMMPS code checks for valid angle definitions, cannot use the same atom ID of either  $i$  or  $j$  (this was allowed and recommended with older LAMMPS versions).

---

The `newton` command for intramolecular interactions must be “on” (which is the default except when using some accelerator packages).

---

**Note:** This angle style should **NOT** be used with fix shake.

---

### 21.13.5 Related commands

*angle\_coeff*, *angle\_hybrid*

### 21.13.6 Default

none

**(Orsi)** Orsi & Essex, The ELBA force field for coarse-grain modeling of lipid membranes, PloS ONE 6(12): e28637, 2011.

**(Allen)** Allen & Tildesley, Computer Simulation of Liquids, Clarendon Press, Oxford, 1987.

## 21.14 *angle\_style* *fourier* command

Accelerator Variants: *fourier/omp*

### 21.14.1 Syntax

```
angle_style fourier
```

### 21.14.2 Examples

```
angle_style fourier  
angle_coeff 75.0 1.0 1.0 1.0
```

### 21.14.3 Description

The *fourier* angle style uses the potential

$$E = K[C_0 + C_1 \cos(\theta) + C_2 \cos(2\theta)]$$

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $C_0$  (real)
- $C_1$  (real)
- $C_2$  (real)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 21.14.4 Restrictions

This angle style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

### 21.14.5 Related commands

*angle\_coeff*

### 21.14.6 Default

none

## 21.15 angle\_style fourier/simple command

Accelerator Variants: *fourier/simple/omp*

### 21.15.1 Syntax

```
angle_style fourier/simple
```

### 21.15.2 Examples

```
angle_style fourier/simple
angle_coeff 100.0 -1.0 1.0
```

### 21.15.3 Description

The *fourier/simple* angle style uses the potential

$$E = K[1.0 + c \cos(n\theta)]$$

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- *K* (energy)
- *c* (real)

- *n* (real)
- 

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 21.15.4 Restrictions

This angle style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the [Build package](#) doc page for more info.

## 21.15.5 Related commands

*angle\_coeff*

## 21.15.6 Default

none

# 21.16 angle\_style gaussian command

## 21.16.1 Syntax

```
angle_style gaussian
```

## 21.16.2 Examples

```
angle_style gaussian  
angle_coeff 1 300.0 2 0.0128 0.375 80.0 0.0730 0.148 123.0
```

### 21.16.3 Description

The *gaussian* angle style uses the potential:

$$E = -k_B T \ln \left( \sum_{i=1}^n \frac{A_i}{w_i \sqrt{\pi/2}} \exp \left( -\frac{(\theta - \theta_i)^2}{w_i^2} \right) \right)$$

This analytical form is a suitable potential for obtaining mesoscale effective force fields which can reproduce target atomistic distributions (*Milano*) The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- T temperature at which the potential was derived
- *n* (integer >=1)
- *A*<sub>1</sub> (-)
- *w*<sub>1</sub> (-)
- *θ*<sub>1</sub> (degrees)
- ...
- *A*<sub>*n*</sub> (-)
- *w*<sub>*n*</sub> (-)
- *θ*<sub>*n*</sub> (degrees)

### 21.16.4 Restrictions

This angle style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

### 21.16.5 Related commands

*angle\_coeff*

### 21.16.6 Default

none

---

(*Milano*) G. Milano, S. Goudeau, F. Mueller-Plathe, J. Polym. Sci. B Polym. Phys. 43, 871 (2005).

## 21.17 *angle\_style harmonic* command

Accelerator Variants: *harmonic/intel*, *harmonic/kk*, *harmonic/omp*

### 21.17.1 Syntax

```
angle_style harmonic
```

### 21.17.2 Examples

```
angle_style harmonic
angle_coeff 1 300.0 107.0
```

### 21.17.3 Description

The *harmonic* angle style uses the potential

$$E = K(\theta - \theta_0)^2$$

where  $\theta_0$  is the equilibrium value of the angle, and  $K$  is a prefactor. Note that the usual 1/2 factor is included in  $K$ .

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $\theta_0$  (degrees)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence  $K$  is effectively energy per radian<sup>2</sup>.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 21.17.4 Restrictions

This angle style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) doc page for more info.

## 21.17.5 Related commands

*angle\_coeff*

## 21.17.6 Default

none

## 21.18 `angle_style` hybrid command

### 21.18.1 Syntax

```
angle_style hybrid style1 style2 ...
```

- style1,style2 = list of one or more angle styles

### 21.18.2 Examples

```
angle_style hybrid harmonic cosine
angle_coeff 1 harmonic 80.0 30.0
angle_coeff 2* cosine 50.0
```

### 21.18.3 Description

The *hybrid* style enables the use of multiple angle styles in one simulation. An angle style is assigned to each angle type. For example, angles in a polymer flow (of angle type 1) could be computed with a *harmonic* potential and angles in the wall boundary (of angle type 2) could be computed with a *cosine* potential. The assignment of angle type to style is made via the *angle\_coeff* command or in the data file.

In the *angle\_coeff* commands, the name of an angle style must be added after the angle type, with the remaining coefficients being those appropriate to that style. In the example above, the 2 *angle\_coeff* commands set angles of angle type 1 to be computed with a *harmonic* potential with coefficients 80.0, 30.0 for  $K$ ,  $\theta_0$ . All other angle types ( $2 - N$ ) are computed with a *cosine* potential with coefficient 50.0 for  $K$ .

If angle coefficients are specified in the data file read via the *read\_data* command, then the same rule applies. E.g. “harmonic” or “cosine”, must be added after the angle type, for each line in the “Angle Coeffs” section, e.g.

```
Angle Coeffs
1 harmonic 80.0 30.0
2 cosine 50.0
...
```

If *class2* is one of the angle hybrid styles, the same rule holds for specifying additional BondBond (and BondAngle) coefficients either via the input script or in the data file. I.e. *class2* must be added to each line after the angle type. For lines in the BondBond (or BondAngle) section of the data file for angle types that are not *class2*, you must use an angle style of *skip* as a placeholder, e.g.



**BondBond Coeffs**

```
1 skip
2 class2 3.6512 1.0119 1.0119
...
```

Note that it is not necessary to use the angle style *skip* in the input script, since BondBond (or BondAngle) coefficients need not be specified at all for angle types that are not *class2*.

An angle style of *none* with no additional coefficients can be used in place of an angle style, either in a input script *angle\_coeff* command or in the data file, if you desire to turn off interactions for specific angle types.

## 21.18.4 Restrictions

This angle style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

Unlike other angle styles, the hybrid angle style does not store angle coefficient info for individual sub-styles in a *binary restart files*. Thus when restarting a simulation from a restart file, you need to re-specify *angle\_coeff* commands.

## 21.18.5 Related commands

*angle\_coeff*

## 21.18.6 Default

none

## 21.19 angle\_style mesocnt command

### 21.19.1 Syntax

```
angle_style mesocnt
```

### 21.19.2 Examples

```
angle_style mesocnt
angle_coeff 1 buckling C 10 10 20.0
angle_coeff 4 harmonic C 8 4 10.0
angle_coeff 2 buckling custom 400.0 50.0 5.0
angle_coeff 1 harmonic custom 300.0
```

### 21.19.3 Description

New in version 15Sep2022.

The *mesocnt* angle style uses the potential

$$E = K_H \Delta\theta^2, \quad |\Delta\theta| < \Delta\theta_B$$
$$E = K_H \Delta\theta_B^2 + K_B (\Delta\theta - \Delta\theta_B), \quad |\Delta\theta| \geq \Delta\theta_B$$

where  $\Delta\theta = \theta - \pi$  is the bending angle of the nanotube,  $K_H$  and  $K_B$  are prefactors for the harmonic and linear regime respectively and  $\Delta\theta_B$  is the buckling angle. Note that the usual 1/2 factor for the harmonic potential is included in  $K_H$ .

The style implements parameterization presets of  $K_H$ ,  $K_B$  and  $\Delta\theta_B$  for mesoscopic simulations of carbon nanotubes based on the atomistic simulations of (*Srivastava*) and buckling considerations of (*Zhigilei*).

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the examples above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- mode = *buckling* or *harmonic*
- preset = *C* or *custom*
- additional parameters depending on preset

If mode *harmonic* is chosen, the potential is simply harmonic and does not switch to the linear term when the buckling angle is reached. In *buckling* mode, the full piecewise potential is used.

Preset *C* is for carbon nanotubes, and the additional parameters are:

- chiral index  $n$  (unitless)
- chiral index  $m$  (unitless)
- $r_0$  (distance)

Here,  $r_0$  is the equilibrium distance of the bonds included in the angle, see *bond\_style mesocnt*.

In harmonic mode with preset *custom*, the additional parameter is:

- $K_H$  (energy)

Hence, this setting is simply a wrapper for *bond\_style harmonic* with an equilibrium angle of 180 degrees.

In harmonic mode with preset *custom*, the additional parameters are:

- $K_H$  (energy)
- $K_B$  (energy)
- $\Delta\theta_B$  (degrees)

$\Delta\theta_B$  is specified in degrees, but LAMMPS converts it to radians internally; hence  $K_H$  is effectively energy per radian<sup>2</sup> and  $K_B$  is energy per radian.

---

In *buckling* mode, this angle style adds the *buckled* property to all atoms in the simulation, which is an integer flag indicating whether the bending angle at a given atom has exceeded  $\Delta\theta_B$ . It can be accessed as an atomic variable, e.g. for custom dump commands, as *i\_buckled*.

---

**Note:** If the initial state of the simulation contains buckled nanotubes and *pair\_style mesocnt* is used, the *i\_buckled* atomic variable needs to be initialized before the *pair\_style* is defined by doing a *run 0* command straight after the *angle\_style* command. See below for an example.

---

If CNTs are already buckled at the start of the simulation, this script will correctly initialize *i\_buckled*:

```
angle_style mesocnt
angle_coeff 1 buckling C 10 10 20.0

run 0

pair_style mesocnt 60.0
pair_coeff * * C_10_10.mesocnt 1
```

### 21.19.4 Restrictions

This angle style can only be used if LAMMPS was built with the MOLECULE and MESONT packages. See the *Build package* doc page for more info.

### 21.19.5 Related commands

*angle\_coeff*

### 21.19.6 Default

none

---

(Srivastava) Zhigilei, Wei, Srivastava, Phys. Rev. B 71, 165417 (2005).

(Zhigilei) Volkov and Zhigilei, ACS Nano 4, 6187 (2010).

## 21.20 angle\_style mm3 command

### 21.20.1 Syntax

```
angle_style mm3
```

### 21.20.2 Examples

```
angle_style mm3
angle_coeff 1 100.0 107.0
```

### 21.20.3 Description

The *mm3* angle style uses the potential that is anharmonic in the angle as defined in (*Allinger*)

$$E = K(\theta - \theta_0)^2 [1 - 0.014(\theta - \theta_0) + 5.6(10)^{-5}(\theta - \theta_0)^2 - 7.0(10)^{-7}(\theta - \theta_0)^3 + 9(10)^{-10}(\theta - \theta_0)^4]$$

where  $\theta_0$  is the equilibrium value of the angle, and  $K$  is a prefactor. The anharmonic prefactors have units  $\text{deg}^{-n}$ , for example  $-0.014 \text{ deg}^{-1}$ ,  $5.6 \cdot 10^{-5} \text{ deg}^{-2}$ , ...

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $\theta_0$  (degrees)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence  $K$  is effectively energy per radian<sup>2</sup>.

### 21.20.4 Restrictions

This angle style can only be used if LAMMPS was built with the YAFF package. See the *Build package* doc page for more info.

### 21.20.5 Related commands

*angle\_coeff*

### 21.20.6 Default

none

## 21.21 angle\_style none command

### 21.21.1 Syntax

```
angle_style none
```

### 21.21.2 Examples

```
angle_style none
```

### 21.21.3 Description

Using an angle style of none means angle forces and energies are not computed, even if triplets of angle atoms were listed in the data file read by the *read\_data* command.

See the *angle\_style zero* command for a way to calculate angle statistics, but compute no angle interactions.

### 21.21.4 Restrictions

none

### 21.21.5 Related commands

*angle\_style zero*

### 21.21.6 Default

none

## 21.22 angle\_style quartic command

Accelerator Variants: *quartic/omp*

### 21.22.1 Syntax

```
angle_style quartic
```

### 21.22.2 Examples

```
angle_style quartic
angle_coeff 1 129.1948 56.8726 -25.9442 -14.2221
```

### 21.22.3 Description

The *quartic* angle style uses the potential

$$E = K_2(\theta - \theta_0)^2 + K_3(\theta - \theta_0)^3 + K_4(\theta - \theta_0)^4$$

where  $\theta_0$  is the equilibrium value of the angle, and  $K$  is a prefactor. Note that the usual 1/2 factor is included in  $K$ .

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $\theta_0$  (degrees)
- $K_2$  (energy)
- $K_3$  (energy)

- $K_4$  (energy)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence the various  $K$  are effectively energy per radian<sup>2</sup> or radian<sup>3</sup> or radian<sup>4</sup>.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

#### 21.22.4 Restrictions

This angle style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the [Build package](#) doc page for more info.

#### 21.22.5 Related commands

*angle\_coeff*

#### 21.22.6 Default

none

### 21.23 angle\_style spica command

Accelerator Variants: *spica/omp*

#### 21.23.1 Syntax

```
angle_style spica
angle_style spica/omp
```

## 21.23.2 Examples

```
angle_style spica
angle_coeff 1 300.0 107.0
```

## 21.23.3 Description

The *spica* angle style is a combination of the harmonic angle potential,

$$E = K(\theta - \theta_0)^2$$

where  $\theta_0$  is the equilibrium value of the angle and  $K$  a prefactor, with the *repulsive* part of the non-bonded *lj/spica* pair style between the atoms 1 and 3. This angle potential is intended for coarse grained MD simulations with the SPICA (formerly called SDK) parameterization using the *pair\_style lj/spica*. Relative to the pair\_style *lj/spica*, however, the energy is shifted by  $\epsilon$ , to avoid sudden jumps. Note that the usual 1/2 factor is included in  $K$ .

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above:

- $K$  (energy)
- $\theta_0$  (degrees)

$\theta_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence  $K$  is effectively energy per radian<sup>2</sup>.

The required *lj/spica* parameters are extracted automatically from the pair\_style.

Style *sdk*, the original implementation of style *spica*, is available for backward compatibility.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

## 21.23.4 Restrictions

This angle style can only be used if LAMMPS was built with the CG-SPICA package. See the *Build package* doc page for more info.

### 21.23.5 Related commands

*angle\_coeff*, *angle\_style harmonic*, *pair\_style lj/spica*, *pair\_style lj/spica/coul/long*

### 21.23.6 Default

none

## 21.24 angle\_style table command

Accelerator Variants: *table/omp*

### 21.24.1 Syntax

```
angle_style table style N
```

- style = *linear* or *spline* = method of interpolation
- N = use N values in table

### 21.24.2 Examples

```
angle_style table linear 1000  
angle_coeff 3 file.table ENTRY1
```

### 21.24.3 Description

Style *table* creates interpolation tables of length  $N$  from angle potential and derivative values listed in a file(s) as a function of angle. The files are read by the *angle\_coeff* command.

The interpolation tables are created by fitting cubic splines to the file values and interpolating energy and derivative values at each of  $N$  angles. During a simulation, these tables are used to interpolate energy and force values on individual atoms as needed. The interpolation is done in one of 2 styles: *linear* or *spline*.

For the *linear* style, the angle is used to find 2 surrounding table values from which an energy or its derivative is computed by linear interpolation.

For the *spline* style, a cubic spline coefficients are computed and stored at each of the  $N$  values in the table. The angle is used to find the appropriate set of coefficients which are used to evaluate a cubic polynomial which computes the energy or derivative.

The following coefficients must be defined for each angle type via the *angle\_coeff* command as in the example above.

- filename
- keyword

The filename specifies a file containing tabulated energy and derivative values. The keyword specifies a section of the file. The format of this file is described below.

---

The format of a tabulated file is as follows (without the parenthesized comments):



```
# Angle potential for harmonic (one or more comment or blank lines)

HAM                               (keyword is the first text on line)
N 181 FP 0 0 EQ 90.0             (N, FP, EQ parameters)
                                   (blank line)
1 0.0 200.5 2.5                  (index, angle, energy, derivative)
2 1.0 198.0 2.5
...
181 180.0 0.0 0.0
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The line can contain additional text, but the initial text must match the argument specified in the [angle\\_coeff](#) command. The next line lists (in any order) one or more parameters for the table. Each parameter is a keyword followed by one or more numeric values.

The parameter “N” is required and its value is the number of table entries that follow. Note that this may be different than the  $N$  specified in the [angle\\_style table](#) command. Let  $N_{\text{table}} = N$  in the [angle\\_style](#) command, and  $N_{\text{file}} = “N”$  in the tabulated file. What LAMMPS does is a preliminary interpolation by creating splines using the  $N_{\text{file}}$  tabulated values as nodal points. It uses these to interpolate as needed to generate energy and derivative values at  $N_{\text{table}}$  different points. The resulting tables of length  $N_{\text{table}}$  are then used as described above, when computing energy and force for individual angles and their atoms. This means that if you want the interpolation tables of length  $N_{\text{table}}$  to match exactly what is in the tabulated file (with effectively no preliminary interpolation), you should set  $N_{\text{table}} = N_{\text{file}}$ .

The “FP” parameter is optional. If used, it is followed by two values  $f_{\text{plo}}$  and  $f_{\text{phi}}$ , which are the second derivatives at the innermost and outermost angle settings. These values are needed by the spline construction routines. If not specified by the “FP” parameter, they are estimated (less accurately) by the first two and last two derivative values in the table.

The “EQ” parameter is also optional. If used, it is followed by a the equilibrium angle value, which is used, for example, by the [fix shake](#) command. If not used, the equilibrium angle is set to 180.0.

Following a blank line, the next  $N$  lines list the tabulated values. On each line, the first value is the index from 1 to  $N$ , the second value is the angle value (in degrees), the third value is the energy (in energy units), and the fourth is  $-dE/d(\theta)$  (also in energy units). The third term is the energy of the 3-atom configuration for the specified angle. The last term is the derivative of the energy with respect to the angle (in degrees, not radians). Thus the units of the last term are still energy, not force. The angle values must increase from one line to the next. The angle values must also begin with 0.0 and end with 180.0, i.e. span the full range of possible angles.

Note that one file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified keyword.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 21.24.4 Restart, fix\_modify, output, run start/stop, minimize info

This angle style writes the settings for the “angle\_style table” command to *binary restart files*, so a angle\_style command does not need to be specified in an input script that reads a restart file. However, the coefficient information is not stored in the restart file, since it is tabulated in the potential files. Thus, angle\_coeff commands do need to be specified in the restart input script.

### 21.24.5 Restrictions

This angle style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

### 21.24.6 Related commands

*angle\_coeff*

### 21.24.7 Default

none

## 21.25 angle\_style zero command

### 21.25.1 Syntax

```
angle_style zero keyword
```

- zero or more keywords may be appended
- keyword = *nocoeff*

### 21.25.2 Examples

```
angle_style zero  
angle_style zero nocoeff  
angle_coeff *  
angle_coeff * 120.0
```

### 21.25.3 Description

Using an angle style of zero means angle forces and energies are not computed, but the geometry of angle triplets is still accessible to other commands.

As an example, the *compute angle/local* command can be used to compute the theta values for the list of triplets of angle atoms listed in the data file read by the *read\_data* command. If no angle style is defined, this command cannot be used.

The optional *nocoeff* flag allows to read data files with AngleCoeff section for any angle style. Similarly, any *angle\_coeff* commands will only be checked for the angle type number and the rest ignored.

Note that the *angle\_coeff* command must be used for all angle types. If specified, there can be only one value, which is going to be used to assign an equilibrium angle, e.g. for use with *fix shake*.

### 21.25.4 Restrictions

none

### 21.25.5 Related commands

*angle\_style none*

### 21.25.6 Default

none



## DIHEDRAL STYLES

### 22.1 dihedral\_style charmm command

Accelerator Variants: *charmm/intel*, *charmm/kk*, *charmm/omp*

### 22.2 dihedral\_style charmmfsw command

#### 22.2.1 Syntax

```
dihedral_style style
```

- style = *charmm* or *charmmfsw*

#### 22.2.2 Examples

```
dihedral_style charmm
dihedral_style charmmfsw
dihedral_coeff 1 0.2 1 180 1.0
dihedral_coeff 2 1.8 1 0 1.0
dihedral_coeff 1 3.1 2 180 0.5
```

#### 22.2.3 Description

The *charmm* and *charmmfsw* dihedral styles use the potential

$$E = K[1 + \cos(n\phi - d)]$$

See (*MacKerell*) for a description of the CHARMM force field. This dihedral style can also be used for the AMBER force field (see comment on weighting factors below). See (*Cornell*) for a description of the AMBER force field.

---

**Note:** The newer *charmmfsw* style was released in March 2017. We recommend it be used instead of the older *charmm* style when running a simulation with the CHARMM force field, either with long-range Coulombics or a Coulombic cutoff, via the *pair\_style lj/charmmfsw/coul/long* and *pair\_style lj/charmmfsw/coul/charmmfsh* commands respectively. Otherwise the older *charmm* style is fine to use. See the discussion below and more details on the *pair\_style charmm* doc page.

---

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $n$  (integer  $\geq 0$ )
- $d$  (integer value of degrees)
- weighting factor (1.0, 0.5, or 0.0)

The weighting factor is required to correct for double counting pairwise non-bonded Lennard-Jones interactions in cyclic systems or when using the CHARMM dihedral style with non-CHARMM force fields. With the CHARMM dihedral style, interactions between the first and fourth atoms in a dihedral are skipped during the normal non-bonded force computation and instead evaluated as part of the dihedral using special epsilon and sigma values specified with the *pair\_coeff* command of pair styles that contain “lj/charmm” (e.g. *pair\_style lj/charmm/coul/long*). In 6-membered rings, the same 1-4 interaction would be computed twice (once for the clockwise 1-4 pair in dihedral 1-2-3-4 and once in the counterclockwise dihedral 1-6-5-4) and thus the weighting factor has to be 0.5 in this case. In 4-membered or 5-membered rings, the 1-4 dihedral also is counted as a 1-2 or 1-3 interaction when going around the ring in the opposite direction and thus the weighting factor is 0.0, as the 1-2 and 1-3 exclusions take precedence.

Note that this dihedral weighting factor is unrelated to the scaling factor specified by the *special\_bonds* command which applies to all 1-4 interactions in the system. For CHARMM force fields, the *special\_bonds* 1-4 interaction scaling factor should be set to 0.0. Since the corresponding 1-4 non-bonded interactions are computed with the dihedral. This means that if any of the weighting factors defined as dihedral coefficients (fourth coeff above) are non-zero, then you must use a pair style with “lj/charmm” and set the *special\_bonds* 1-4 scaling factor to 0.0 (which is the default). Otherwise 1-4 non-bonded interactions in dipoles will be computed twice.

For simulations using the CHARMM force field with a Coulombic cutoff, the difference between the *charmm* and *charmmfsw* styles is in the computation of the 1-4 non-bond interactions, though only if the distance between the two atoms is within the switching region of the pairwise potential defined by the corresponding CHARMM pair style, i.e. within the outer cutoff specified for the pair style. The *charmmfsw* style should only be used when using the corresponding *pair\_style lj/charmmfsw/coul/charmmfsw* or *pair\_style lj/charmmfsw/coul/long* commands. Use the *charmm* style with the older *pair\_style* commands that have just “charmm” in their style name. See the discussion on the *CHARMM pair\_style* page for details.

Note that for AMBER force fields, which use pair styles with “lj/cut”, the *special\_bonds* 1-4 scaling factor should be set to the AMBER defaults (1/2 and 5/6) and all the dihedral weighting factors (fourth coeff above) must be set to 0.0. In this case, you can use any pair style you wish, since the dihedral does not need any Lennard-Jones parameter information and will not compute any 1-4 non-bonded interactions. Likewise the *charmm* or *charmmfsw* styles are identical in this case since no 1-4 non-bonded interactions are computed.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 22.2.4 Restrictions

When using run\_style *respa*, these dihedral styles must be assigned to the same r-RESPA level as *pair* or *outer*.

When used in combination with CHARMM pair styles, the 1-4 *special\_bonds* scaling factors must be set to 0.0. Otherwise non-bonded contributions for these 1-4 pairs will be computed multiple times.

These dihedral styles can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

## 22.2.5 Related commands

*dihedral\_coeff*

## 22.2.6 Default

none

---

(**Cornell**) Cornell, Cieplak, Bayly, Gould, Merz, Ferguson, Spellmeyer, Fox, Caldwell, Kollman, JACS 117, 5179-5197 (1995).

(**MacKerell**) MacKerell, Bashford, Bellott, Dunbrack, Evanseck, Field, Fischer, Gao, Guo, Ha, et al, J Phys Chem B, 102, 3586 (1998).

## 22.3 dihedral\_style class2 command

Accelerator Variants: *class2/omp*, *class2/kk*

### 22.3.1 Syntax

```
dihedral_style class2
```

### 22.3.2 Examples

```
dihedral_style class2
dihedral_coeff 1 100 75 100 70 80 60
dihedral_coeff * mbt 3.5945 0.1704 -0.5490 1.5228
dihedral_coeff * ebt 0.3417 0.3264 -0.9036 0.1368 0.0 -0.8080 1.0119 1.1010
dihedral_coeff 2 at 0.0 -0.1850 -0.7963 -2.0220 0.0 -0.3991 110.2453 105.1270
dihedral_coeff * aat -13.5271 110.2453 105.1270
dihedral_coeff * bb13 0.0 1.0119 1.1010
```

### 22.3.3 Description

The *class2* dihedral style uses the potential

$$\begin{aligned}
 E &= E_d + E_{mbt} + E_{ebt} + E_{at} + E_{aat} + E_{bb13} \\
 E_d &= \sum_{n=1}^3 K_n [1 - \cos(n\phi - \phi_n)] \\
 E_{mbt} &= (r_{jk} - r_2) [A_1 \cos(\phi) + A_2 \cos(2\phi) + A_3 \cos(3\phi)] \\
 E_{ebt} &= (r_{ij} - r_1) [B_1 \cos(\phi) + B_2 \cos(2\phi) + B_3 \cos(3\phi)] + \\
 &\quad (r_{kl} - r_3) [C_1 \cos(\phi) + C_2 \cos(2\phi) + C_3 \cos(3\phi)] \\
 E_{at} &= (\theta_{ijk} - \theta_1) [D_1 \cos(\phi) + D_2 \cos(2\phi) + D_3 \cos(3\phi)] + \\
 &\quad (\theta_{jkl} - \theta_2) [E_1 \cos(\phi) + E_2 \cos(2\phi) + E_3 \cos(3\phi)] \\
 E_{aat} &= M(\theta_{ijk} - \theta_1)(\theta_{jkl} - \theta_2) \cos(\phi) \\
 E_{bb13} &= N(r_{ij} - r_1)(r_{kl} - r_3)
 \end{aligned}$$

where  $E_d$  is the dihedral term,  $E_{mbt}$  is a middle-bond-torsion term,  $E_{ebt}$  is an end-bond-torsion term,  $E_{at}$  is an angle-torsion term,  $E_{aat}$  is an angle-angle-torsion term, and  $E_{bb13}$  is a bond-bond-13 term.

$\theta_1$  and  $\theta_2$  are equilibrium angles and  $r_1$ ,  $r_2$ , and  $r_3$  are equilibrium bond lengths.

See ([Sun](#)) for a description of the COMPASS *class2* force field.

Coefficients for the  $E_d$ ,  $E_{mbt}$ ,  $E_{ebt}$ ,  $E_{at}$ ,  $E_{aat}$ , and  $E_{bb13}$  formulas must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands.

These are the 6 coefficients for the  $E_d$  formula:

- $K_1$  (energy)
- $\phi_1$  (degrees)
- $K_2$  (energy)
- $\phi_2$  (degrees)
- $K_3$  (energy)
- $\phi_3$  (degrees)

For the  $E_{mbt}$  formula, each line in a *dihedral\_coeff* command in the input script lists 5 coefficients, the first of which is *mbt* to indicate they are MiddleBondTorsion coefficients. In a data file, these coefficients should be listed under a *MiddleBondTorsion Coeffs* heading and you must leave out the *mbt*, i.e. only list 4 coefficients after the dihedral type.

- *mbt*
- $A_1$  (energy/distance)
- $A_2$  (energy/distance)
- $A_3$  (energy/distance)
- $r_2$  (distance)

For the  $E_{ebt}$  formula, each line in a *dihedral\_coeff* command in the input script lists 9 coefficients, the first of which is *ebt* to indicate they are EndBondTorsion coefficients. In a data file, these coefficients should be listed under a *EndBondTorsion Coeffs* heading and you must leave out the *ebt*, i.e. only list 8 coefficients after the dihedral type.

- *ebt*
- $B_1$  (energy/distance)



- $B_2$  (energy/distance)
- $B_3$  (energy/distance)
- $C_1$  (energy/distance)
- $C_2$  (energy/distance)
- $C_3$  (energy/distance)
- $r_1$  (distance)
- $r_3$  (distance)

For the  $E_{at}$  formula, each line in a *dihedral\_coeff* command in the input script lists 9 coefficients, the first of which is *at* to indicate they are AngleTorsion coefficients. In a data file, these coefficients should be listed under a *AngleTorsion Coeffs* heading and you must leave out the *at*, i.e. only list 8 coefficients after the dihedral type.

- *at*
- $D_1$  (energy)
- $D_2$  (energy)
- $D_3$  (energy)
- $E_1$  (energy)
- $E_2$  (energy)
- $E_3$  (energy)
- $\theta_1$  (degrees)
- $\theta_2$  (degrees)

$\theta_1$  and  $\theta_2$  are specified in degrees, but LAMMPS converts them to radians internally; hence the various  $D$  and  $E$  are effectively energy per radian.

For the  $E_{aat}$  formula, each line in a *dihedral\_coeff* command in the input script lists 4 coefficients, the first of which is *aat* to indicate they are AngleAngleTorsion coefficients. In a data file, these coefficients should be listed under a *AngleAngleTorsion Coeffs* heading and you must leave out the *aat*, i.e. only list 3 coefficients after the dihedral type.

- *aat*
- $M$  (energy)
- $\theta_1$  (degrees)
- $\theta_2$  (degrees)

$\theta_1$  and  $\theta_2$  are specified in degrees, but LAMMPS converts them to radians internally; hence  $M$  is effectively energy per radian<sup>2</sup>.

For the  $E_{bb13}$  formula, each line in a *dihedral\_coeff* command in the input script lists 4 coefficients, the first of which is *bb13* to indicate they are BondBond13 coefficients. In a data file, these coefficients should be listed under a *BondBond13 Coeffs* heading and you must leave out the *bb13*, i.e. only list 3 coefficients after the dihedral type.

- *bb13*
- $N$  (energy/distance<sup>2</sup>)
- $r_1$  (distance)
- $r_3$  (distance)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 22.3.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the CLASS2 package. See the [Build package](#) doc page for more info.

### 22.3.5 Related commands

*dihedral\_coeff*

### 22.3.6 Default

none

---

(Sun) Sun, J Phys Chem B 102, 7338-7364 (1998).

## 22.4 dihedral\_style cosine/shift/exp command

Accelerator Variants: *cosine/shift/exp/omp*

### 22.4.1 Syntax

```
dihedral_style cosine/shift/exp
```

### 22.4.2 Examples

```
dihedral_style cosine/shift/exp
dihedral_coeff 1 10.0 45.0 2.0
```

### 22.4.3 Description

The *cosine/shift/exp* dihedral style uses the potential

$$E = -U_{min} \frac{e^{-aU(\theta, \theta_0)} - 1}{e^a - 1} \quad \text{with} \quad U(\theta, \theta_0) = -0.5(1 + \cos(\theta - \theta_0))$$

where  $U_{min}$ ,  $\theta$ , and  $a$  are defined for each dihedral type.

The potential is bounded between  $[-U_{min} : 0]$  and the minimum is located at the angle  $\theta_0$ . The  $a$  parameter can be both positive or negative and is used to control the spring constant at the equilibrium.

The spring constant is given by  $k = ae^a \frac{U_{min}}{2(e^a - 1)}$ . For  $a > 3$  and  $\frac{k}{U_{min}} = \frac{a}{2}$  to better than 5% relative error. For negative values of the  $a$  parameter, the spring constant is essentially zero, and anharmonic terms takes over. The potential is furthermore well behaved in the limit  $a \rightarrow 0$ , where it has been implemented to linear order in  $a$  for  $a < 0.001$ .

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $U_{min}$  (energy)
- $\theta$  (angle)
- $a$  (real number)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

### 22.4.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

### 22.4.5 Related commands

*dihedral\_coeff*, *angle\_style cosine/shift/exp*

## 22.4.6 Default

none

## 22.5 dihedral\_style fourier command

Accelerator Variants: *fourier/intel*, *fourier/omp*

### 22.5.1 Syntax

```
dihedral_style fourier
```

### 22.5.2 Examples

```
dihedral_style fourier
dihedral_coeff 1 3 -0.846200 3 0.0 7.578800 1 0 0.138000 2 -180.0
```

### 22.5.3 Description

The *fourier* dihedral style uses the potential:

$$E = \sum_{i=1,m} K_i [1.0 + \cos(n_i \phi - d_i)]$$

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $m$  (integer  $\geq 1$ )
- $K_1$  (energy)
- $n_1$  (integer  $\geq 0$ )
- $d_1$  (degrees)
- [...]
- $K_m$  (energy)
- $n_m$  (integer  $\geq 0$ )
- $d_m$  (degrees)

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 22.5.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

## 22.5.5 Related commands

*dihedral\_coeff*

## 22.5.6 Default

none

## 22.6 dihedral\_style harmonic command

Accelerator Variants: *harmonic/intel*, *harmonic/kk*, *harmonic/omp*

### 22.6.1 Syntax

```
dihedral_style harmonic
```

### 22.6.2 Examples

```
dihedral_style harmonic
dihedral_coeff 1 80.0 1 2
```

### 22.6.3 Description

The *harmonic* dihedral style uses the potential

$$E = K[1 + d \cos(n\phi)]$$

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- *K* (energy)
- *d* (+1 or -1)
- *n* (integer  $\geq 0$ )

---

**Note:** Here are important points to take note of when defining LAMMPS dihedral coefficients for the harmonic style, so that they are compatible with how harmonic dihedrals are defined by other force fields:

---

- The LAMMPS convention is that the trans position = 180 degrees, while in some force fields trans = 0 degrees.
  - Some force fields reverse the sign convention on  $d$ .
  - Some force fields let  $n$  be positive or negative which corresponds to  $d = 1$  or  $d = -1$  for the harmonic style.
- 

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 22.6.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) doc page for more info.

## 22.6.5 Related commands

*dihedral\_coeff*

## 22.6.6 Default

none

## 22.7 dihedral\_style helix command

Accelerator Variants: *helix/omp*

### 22.7.1 Syntax

```
dihedral_style helix
```

## 22.7.2 Examples

```
dihedral_style helix
dihedral_coeff 1 80.0 100.0 40.0
```

## 22.7.3 Description

The *helix* dihedral style uses the potential

$$E = A[1 - \cos(\theta)] + B[1 + \cos(3\theta)] + C[1 + \cos(\theta + \frac{\pi}{4})]$$

This coarse-grain dihedral potential is described in (*Guo*). For dihedral angles in the helical region, the energy function is represented by a standard potential consisting of three minima, one corresponding to the trans (t) state and the other to gauche states (g+ and g-). The paper describes how the *A*, *B* and, *C* parameters are chosen so as to balance secondary (largely driven by local interactions) and tertiary structure (driven by long-range interactions).

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- *A* (energy)
- *B* (energy)
- *C* (energy)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

## 22.7.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

## 22.7.5 Related commands

*dihedral\_coeff*

## 22.7.6 Default

none

---

(Guo) Guo and Thirumalai, Journal of Molecular Biology, 263, 323-43 (1996).

## 22.8 dihedral\_style hybrid command

### 22.8.1 Syntax

```
dihedral_style hybrid style1 style2 ...
```

- style1,style2 = list of one or more dihedral styles

### 22.8.2 Examples

```
dihedral_style hybrid harmonic helix
dihedral_coeff 1 harmonic 6.0 1 3
dihedral_coeff 2* helix 10 10 10
```

### 22.8.3 Description

The *hybrid* style enables the use of multiple dihedral styles in one simulation. An dihedral style is assigned to each dihedral type. For example, dihedrals in a polymer flow (of dihedral type 1) could be computed with a *harmonic* potential and dihedrals in the wall boundary (of dihedral type 2) could be computed with a *helix* potential. The assignment of dihedral type to style is made via the *dihedral\_coeff* command or in the data file.

In the *dihedral\_coeff* commands, the name of a dihedral style must be added after the dihedral type, with the remaining coefficients being those appropriate to that style. In the example above, the 2 *dihedral\_coeff* commands set dihedrals of dihedral type 1 to be computed with a *harmonic* potential with coefficients 6.0, 1, 3 for K, d, n. All other dihedral types (2-N) are computed with a *helix* potential with coefficients 10, 10, 10 for A, B, C.

If dihedral coefficients are specified in the data file read via the *read\_data* command, then the same rule applies. E.g. “harmonic” or “helix”, must be added after the dihedral type, for each line in the “Dihedral Coeffs” section, e.g.

```
Dihedral Coeffs
1 harmonic 6.0 1 3
2 helix 10 10 10
...
```

If *class2* is one of the dihedral hybrid styles, the same rule holds for specifying additional AngleTorsion (and EndBondTorsion, etc) coefficients either via the input script or in the data file. I.e. *class2* must be added to each line after the dihedral type. For lines in the AngleTorsion (or EndBondTorsion, etc) Coeffs section of the data file for dihedral types that are not *class2*, you must use an dihedral style of *skip* as a placeholder, e.g.



**AngleTorsion Coeffs**

```
1 skip
2 class2 1.0 1.0 1.0 3.0 3.0 3.0 30.0 50.0
...
```

Note that it is not necessary to use the dihedral style *skip* in the input script, since AngleTorsion (or EndBondTorsion, etc) coefficients need not be specified at all for dihedral types that are not *class2*.

A dihedral style of *none* with no additional coefficients can be used in place of a dihedral style, either in a input script dihedral\_coeff command or in the data file, if you desire to turn off interactions for specific dihedral types.

## 22.8.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

Unlike other dihedral styles, the hybrid dihedral style does not store dihedral coefficient info for individual sub-styles in a *binary restart files*. Thus when restarting a simulation from a restart file, you need to re-specify dihedral\_coeff commands.

## 22.8.5 Related commands

*dihedral\_coeff*

## 22.8.6 Default

none

## 22.9 dihedral\_style multi/harmonic command

Accelerator Variants: *multi/harmonic/omp*

### 22.9.1 Syntax

```
dihedral_style multi/harmonic
```

### 22.9.2 Examples

```
dihedral_style multi/harmonic
dihedral_coeff 1 20 20 20 20 20
```

### 22.9.3 Description

The *multi/harmonic* dihedral style uses the potential

$$E = \sum_{n=1,5} A_n \cos^{n-1}(\phi)$$

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $A_1$  (energy)
  - $A_2$  (energy)
  - $A_3$  (energy)
  - $A_4$  (energy)
  - $A_5$  (energy)
- 

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 22.9.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

### 22.9.5 Related commands

*dihedral\_coeff*

### 22.9.6 Default

none

## 22.10 dihedral\_style nharmonic command

Accelerator Variants: *nharmonic/omp*

### 22.10.1 Syntax

```
dihedral_style nharmonic
```

### 22.10.2 Examples

```
dihedral_style nharmonic
dihedral_coeff * 3 10.0 20.0 30.0
```

### 22.10.3 Description

The *nharmonic* dihedral style uses the potential:

$$E = \sum_{n=1,n} A_n \cos^{n-1}(\phi)$$

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $n$  (integer  $\geq 1$ )
- $A_1$  (energy)
- $A_2$  (energy)
- ...
- $A_n$  (energy)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

## 22.10.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

## 22.10.5 Related commands

*dihedral\_coeff*

## 22.10.6 Default

none

## 22.11 dihedral\_style none command

### 22.11.1 Syntax

```
dihedral_style none
```

### 22.11.2 Examples

```
dihedral_style none
```

### 22.11.3 Description

Using a dihedral style of none means dihedral forces and energies are not computed, even if quadruplets of dihedral atoms were listed in the data file read by the *read\_data* command.

See the *dihedral\_style zero* command for a way to calculate dihedral statistics, but compute no dihedral interactions.

### 22.11.4 Restrictions

none

### 22.11.5 Related commands

*dihedral\_style zero*

## 22.11.6 Default

none

## 22.12 dihedral\_style opls command

Accelerator Variants: *opls/intel*, *opls/kk*, *opls/omp*

### 22.12.1 Syntax

```
dihedral_style opls
```

### 22.12.2 Examples

```
dihedral_style opls
dihedral_coeff 1 1.740 -0.157 0.279 0.000 # CT-CT-CT-CT
dihedral_coeff 2 0.000 0.000 0.366 0.000 # CT-CT-CT-HC
dihedral_coeff 3 0.000 0.000 0.318 0.000 # HC-CT-CT-HC
```

### 22.12.3 Description

The *opls* dihedral style uses the potential

$$E = \frac{1}{2}K_1[1 + \cos(\phi)] + \frac{1}{2}K_2[1 - \cos(2\phi)] + \frac{1}{2}K_3[1 + \cos(3\phi)] + \frac{1}{2}K_4[1 - \cos(4\phi)]$$

Note that the usual 1/2 factor is not included in the K values.

This dihedral potential is used in the OPLS force field and is described in ([Watkins](#)).

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K_1$  (energy)
- $K_2$  (energy)
- $K_3$  (energy)
- $K_4$  (energy)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

## 22.12.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

## 22.12.5 Related commands

*dihedral\_coeff*

## 22.12.6 Default

none

---

(**Watkins**) Watkins and Jorgensen, J Phys Chem A, 105, 4118-4125 (2001).

## 22.13 dihedral\_style quadratic command

Accelerator Variants: *quadratic/omp*

### 22.13.1 Syntax

```
dihedral_style quadratic
```

### 22.13.2 Examples

```
dihedral_style quadratic  
dihedral_coeff 100.0 80.0
```

### 22.13.3 Description

The *quadratic* dihedral style uses the potential:

$$E = K(\phi - \phi_0)^2$$

This dihedral potential can be used to keep a dihedral in a predefined value (cis=zero, right-hand convention is used).

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $\phi_0$  (degrees)

$\phi_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence  $K$  is effectively energy per radian<sup>2</sup>.

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

## 22.13.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the [Build package](#) doc page for more info.

## 22.13.5 Related commands

*dihedral\_coeff*

## 22.13.6 Default

none

## 22.14 dihedral\_style spherical command

### 22.14.1 Syntax

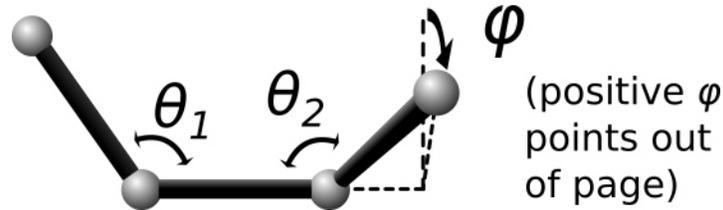
```
dihedral_style spherical
```

### 22.14.2 Examples

```
dihedral_coeff 1 1 286.1 1 124 1 1 90.0 0 1 90.0 0
dihedral_coeff 1 3 69.3 1 93.9 1 1 90 0 1 90 0 &
                49.1 0 0.00 0 1 74.4 1 0 0.00 0 &
                25.2 0 0.00 0 0 0.00 0 1 48.1 1
```

### 22.14.3 Description

The *spherical* dihedral style uses the potential:



$$E(\phi, \theta_1, \theta_2) = \sum_{i=1}^N C_i \Phi_i(\phi) \Theta_{1i}(\theta_1) \Theta_{2i}(\theta_2)$$

$$\Phi_i(\phi) = u_i - \cos((\phi - a_i)K_i)$$

$$\Theta_{1i}(\theta_1) = v_i - \cos((\theta_1 - b_i)L_i)$$

$$\Theta_{2i}(\theta_2) = w_i - \cos((\theta_2 - c_i)M_i)$$

For this dihedral style, the energy can be any function that combines the 4-body dihedral-angle ( $\phi$ ) and the two 3-body bond-angles ( $\theta_1$ ,  $\theta_2$ ). For this reason, there is usually no need to define 3-body “angle” forces separately for the atoms participating in these interactions. It is probably more efficient to incorporate 3-body angle forces into the dihedral interaction even if it requires adding additional terms to the expansion (as was done in the second example). A careful choice of parameters can prevent singularities that occur with traditional force-fields whenever theta1 or theta2 approach 0 or 180 degrees.

The last example above corresponds to an interaction with a single energy minima located near  $\phi = 93.9$ ,  $\theta_1 = 74.4$ ,  $\theta_2 = 48.1$  degrees, and it remains numerically stable at all angles ( $\phi$ ,  $\theta_1$ ,  $\theta_2$ ). In this example, the coefficients 49.1, and 25.2 can be physically interpreted as the harmonic spring constants for theta1 and theta2 around their minima. The coefficient 69.3 is the harmonic spring constant for phi after division by  $\sin(74.4) \cdot \sin(48.1)$  (the minima positions for theta1 and theta2).

The following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above, or in the Dihedral Coeffs section of a data file read by the *read\_data* command:

- $n$  (integer  $\geq 1$ )
- $C_1$  (energy)
- $K_1$  (typically an integer)
- $a_1$  (degrees)
- $u_1$  (typically 0.0 or 1.0)
- $L_1$  (typically an integer)
- $b_1$  (degrees, typically 0.0 or 90.0)
- $v_1$  (typically 0.0 or 1.0)
- $M_1$  (typically an integer)
- $c_1$  (degrees, typically 0.0 or 90.0)
- $w_1$  (typically 0.0 or 1.0)
- [...]
- $C_n$  (energy)
- $K_n$  (typically an integer)
- $a_n$  (degrees)



- $u_n$  (typically 0.0 or 1.0)
  - $L_n$  (typically an integer)
  - $b_n$  (degrees, typically 0.0 or 90.0)
  - $v_n$  (typically 0.0 or 1.0)
  - $M_n$  (typically an integer)
  - $c_n$  (degrees, typically 0.0 or 90.0)
  - $w_n$  (typically 0.0 or 1.0)
- 

## 22.14.4 Restrictions

This dihedral style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

## 22.14.5 Related commands

*dihedral\_coeff*

## 22.14.6 Default

none

## 22.15 dihedral\_style table command

Accelerator Variants: *table/omp*

## 22.16 dihedral\_style table/cut command

### 22.16.1 Syntax

<code>dihedral_style style interpolation Ntable</code>
--

- style = *table* or *table/cut*
- interpolation = *linear* or *spline* = method of interpolation
- Ntable = size of the internal lookup table

## 22.16.2 Examples

```

dihedral_style table spline 400
dihedral_style table linear 1000
dihedral_coeff 1 file.table DIH_TABLE1
dihedral_coeff 2 file.table DIH_TABLE2

dihedral_style table/cut spline 400
dihedral_style table/cut linear 1000
dihedral_coeff 1 aat 1.0 177 180 file.table DIH_TABLE1
dihedral_coeff 2 aat 0.5 170 180 file.table DIH_TABLE2

```

## 22.16.3 Description

The *table* and *table/cut* dihedral styles create interpolation tables of length *Ntable* from dihedral potential and derivative values listed in a file(s) as a function of the dihedral angle “phi”. The files are read by the *dihedral\_coeff* command. For dihedral style *table/cut* additionally an analytic cutoff that is quadratic in the bond-angle (theta) is applied in order to regularize the dihedral interaction.

The interpolation tables are created by fitting cubic splines to the file values and interpolating energy and derivative values at each of *Ntable* dihedral angles. During a simulation, these tables are used to interpolate energy and force values on individual atoms as needed. The interpolation is done in one of 2 styles: *linear* or *spline*.

For the *linear* style, the dihedral angle (phi) is used to find 2 surrounding table values from which an energy or its derivative is computed by linear interpolation.

For the *spline* style, cubic spline coefficients are computed and stored at each of the *Ntable* evenly-spaced values in the interpolated table. For a given dihedral angle (phi), the appropriate coefficients are chosen from this list, and a cubic polynomial is used to compute the energy and the derivative at this angle.

For dihedral style *table* the following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above.

- filename
- keyword

The filename specifies a file containing tabulated energy and derivative values. The keyword specifies which section of the file to read. The format of this file is the same for both dihedral styles and described below.

For dihedral style *table/cut* the following coefficients must be defined for each dihedral type via the *dihedral\_coeff* command as in the example above.

- style (aat)
- cutoff prefactor
- cutoff angle1
- cutoff angle2
- filename
- keyword

The cutoff dihedral style uses a tabulated dihedral interaction with a cutoff function:

$$\begin{aligned}
 f(\theta) &= K & \theta < \theta_1 \\
 f(\theta) &= K \left( 1 - \frac{(\theta - \theta_1)^2}{(\theta_2 - \theta_1)^2} \right) & \theta_1 < \theta < \theta_2
 \end{aligned}$$

The cutoff specifies an prefactor to the cutoff function. While this value would ordinarily equal 1 there may be situations where the value should change.

The cutoff  $\theta_1$  specifies the angle (in degrees) below which the dihedral interaction is unmodified, i.e. the cutoff function is 1.

The cutoff function is applied between  $\theta_1$  and  $\theta_2$ , which is the angle at which the cutoff function drops to zero. The value of zero effectively “turns off” the dihedral interaction.

The filename specifies a file containing tabulated energy and derivative values. The keyword specifies which section of the file to read. The format of this file is the same for both dihedral styles and described below.

The format of a tabulated file is as follows (without the parenthesized comments). It can begin with one or more comment or blank lines.

```
# Table of the potential and its negative derivative

DIH_TABLE1                (keyword is the first text on line)
N 30 DEGREES              (N, NOF, DEGREES, RADIANS, CHECKU/F)
                          (blank line)
1 -168.0 -1.40351172223 0.0423346818422
2 -156.0 -1.70447981034 0.00811786522531
3 -144.0 -1.62956100432 -0.0184129719987
...
30 180.0 -0.707106781187 0.0719306095245

# Example 2: table of the potential. Forces omitted

DIH_TABLE2
N 30 NOF CHECKU testU.dat CHECKF testF.dat

1 -168.0 -1.40351172223
2 -156.0 -1.70447981034
3 -144.0 -1.62956100432
...
30 180.0 -0.707106781187
```

A section begins with a non-blank line whose first character is not a “#”; blank lines or lines starting with “#” can be used as comments between sections. The first line begins with a keyword which identifies the section. The line can contain additional text, but the initial text must match the argument specified in the *dihedral\_coeff* command. The next line lists (in any order) one or more parameters for the table. Each parameter is a keyword followed by one or more numeric values.

Following a blank line, the next N lines list the tabulated values. On each line, the first value is the index from 1 to N, the second value is the angle value, the third value is the energy (in energy units), and the fourth is -dE/d(phi) also in energy units). The third term is the energy of the 4-atom configuration for the specified angle. The fourth term (when present) is the negative derivative of the energy with respect to the angle (in degrees, or radians depending on whether the user selected DEGREES or RADIANS). Thus the units of the last term are still energy, not force. The dihedral angle values must increase from one line to the next.

Dihedral table splines are cyclic. There is no discontinuity at 180 degrees (or at any other angle). Although in the examples above, the angles range from -180 to 180 degrees, in general, the first angle in the list can have any value (positive, zero, or negative). However the *range* of angles represented in the table must be *strictly* less than 360 degrees (2pi radians) to avoid angle overlap. (You may not supply entries in the table for both 180 and -180, for example.) If the user’s table covers only a narrow range of dihedral angles, strange numerical behavior can occur in the large remaining gap.

**Parameters:**

The parameter “N” is required and its value is the number of table entries that follow. Note that this may be different than the N specified in the *dihedral\_style table* command. Let *Ntable* is the number of table entries requested *dihedral\_style* command, and let *Nfile* be the parameter following “N” in the tabulated file (“30” in the sparse example above). What LAMMPS does is a preliminary interpolation by creating splines using the *Nfile* tabulated values as nodal points. It uses these to interpolate as needed to generate energy and derivative values at *Ntable* different points (which are evenly spaced over a 360 degree range, even if the angles in the file are not). The resulting tables of length *Ntable* are then used as described above, when computing energy and force for individual dihedral angles and their atoms. This means that if you want the interpolation tables of length *Ntable* to match exactly what is in the tabulated file (with effectively nopreliminary interpolation), you should set  $Ntable = Nfile$ . To insure the nodal points in the user’s file are aligned with the interpolated table entries, the angles in the table should be integer multiples of  $360/Ntable$  degrees, or  $2\pi/Ntable$  radians (depending on your choice of angle units).

The optional “NOF” keyword allows the user to omit the forces (negative energy derivatives) from the table file (normally located in the fourth column). In their place, forces will be calculated automatically by differentiating the potential energy function indicated by the third column of the table (using either linear or spline interpolation).

The optional “DEGREES” keyword allows the user to specify angles in degrees instead of radians (default).

The optional “RADIANS” keyword allows the user to specify angles in radians instead of degrees. (Note: This changes the way the forces are scaled in the fourth column of the data file.)

The optional “CHECKU” keyword is followed by a filename. This allows the user to save all of the *Ntable* different entries in the interpolated energy table to a file to make sure that the interpolated function agrees with the user’s expectations. (Note: You can temporarily increase the *Ntable* parameter to a high value for this purpose. “*Ntable*” is explained above.)

The optional “CHECKF” keyword is analogous to the “CHECKU” keyword. It is followed by a filename, and it allows the user to check the interpolated force table. This option is available even if the user selected the “NOF” option.

Note that one file can contain many sections, each with a tabulated potential. LAMMPS reads the file section by section until it finds one that matches the specified keyword.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

#### 22.16.4 Restart, fix\_modify, output, run start/stop, minimize info

These dihedral styles write the settings for the “*dihedral\_style table*” or “*dihedral\_style table/cut*” command to *binary restart files*, so a *dihedral\_style* command does not need to be specified in an input script that reads a restart file. However, the coefficient information loaded from the table file(s) is not stored in the restart file, since it is tabulated in the potential files. Thus, suitable *dihedral\_coeff* commands do need to be specified in the restart input script after reading the restart file.

## 22.16.5 Restrictions

The *table* dihedral style can only be used if LAMMPS was built with the MOLECULE package. The *table/cut* dihedral style can only be used if LAMMPS was built with the EXTRA-MOLECULE package. See the *Build package* doc page for more info.

## 22.16.6 Related commands

*dihedral\_coeff*

## 22.16.7 Default

none

## 22.17 dihedral\_style zero command

### 22.17.1 Syntax

```
dihedral_style zero keyword
```

- zero or more keywords may be appended
- keyword = *nocoeff*

### 22.17.2 Examples

```
dihedral_style zero
dihedral_style zero nocoeff
dihedral_coeff *
```

### 22.17.3 Description

Using a dihedral style of zero means dihedral forces and energies are not computed, but the geometry of dihedral quadruplets is still accessible to other commands.

As an example, the *compute dihedral/local* command can be used to compute the theta values for the list of quadruplets of dihedral atoms listed in the data file read by the *read\_data* command. If no dihedral style is defined, this command cannot be used.

The optional *nocoeff* flag allows to read data files with a DihedralCoeff section for any dihedral style. Similarly, any *dihedral\_coeff* commands will only be checked for the dihedral type number and the rest ignored.

Note that the *dihedral\_coeff* command must be used for all dihedral types, though no additional values are specified.

### 22.17.4 Restrictions

none

### 22.17.5 Related commands

none

*dihedral\_style none*

### 22.17.6 Default

none

## IMPROPER STYLES

### 23.1 `improper_style` harmonic command

#### 23.1.1 Syntax

```
improper_style amoeba
```

#### 23.1.2 Examples

```
improper_style amoeba  
improper_coeff 1 49.6
```

#### 23.1.3 Description

The *amoeba* improper style uses the potential

$$E = K(\chi)^2$$

where  $\chi$  is the improper angle and  $K$  is a prefactor. Note that the usual 1/2 factor is included in  $K$ .

This formula seems like a simplified version of the formula for the *improper\_style harmonic* command with  $\chi_0 = 0.0$ . However the computation of the angle  $\chi$  is done differently to match how the Tinker MD code computes its out-of-plane improper for the AMOEBA and HIPPO force fields. See the *Howto amoeba* doc page for more information about the implementation of AMOEBA and HIPPO in LAMMPS.

If the 4 atoms in an improper quadruplet (listed in the data file read by the *read\_data* command are ordered I,J,K,L then atoms I,K,L are considered to lie in a plane and atom J is out-of-plane. The angle  $\chi_0$  is computed as the Allinger angle which is defined as the angle between the plane of I,K,L, and the vector from atom I to atom J.

The following coefficient must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)

Note that the angle  $\chi$  is computed in radians; hence  $K$  is effectively energy per radian<sup>2</sup>.

### 23.1.4 Restrictions

This improper style can only be used if LAMMPS was built with the AMOEBA package. See the *Build package* doc page for more info.

### 23.1.5 Related commands

*improper\_coeff*, *improper\_harmonic* :doc:<*improper\_harmonic*>

### 23.1.6 Default

none

## 23.2 improper\_style class2 command

Accelerator Variants: *class2/omp*, *class2/kk*

### 23.2.1 Syntax

```
improper_style class2
```

### 23.2.2 Examples

```
improper_style class2
improper_coeff 1 100.0 0
improper_coeff * aa 0.0 0.0 0.0 115.06 130.01 115.06
```

### 23.2.3 Description

The *class2* improper style uses the potential

$$E = E_i + E_{aa}$$
$$E_i = K \left[ \frac{\chi_{ijkl} + \chi_{kjli} + \chi_{ljik}}{3} - \chi_0 \right]^2$$
$$E_{aa} = M_1(\theta_{ijk} - \theta_1)(\theta_{kjl} - \theta_3) +$$
$$M_2(\theta_{ijk} - \theta_1)(\theta_{ijl} - \theta_2) +$$
$$M_3(\theta_{ijl} - \theta_2)(\theta_{kjl} - \theta_3)$$

where  $E_i$  is the improper term and  $E_{aa}$  is an angle-angle term. The 3  $\chi$  terms in  $E_i$  are an average over 3 out-of-plane angles.

The 4 atoms in an improper quadruplet (listed in the data file read by the *read\_data* command) are ordered I,J,K,L.  $\chi_{ijkl}$  refers to the angle between the plane of I,J,K and the plane of J,K,L, and the bond JK lies in both planes. Similarly for  $\chi_{kjli}$  and  $\chi_{ljik}$ . Note that atom J appears in the common bonds (JI, JK, JL) of all 3 X terms. Thus J (the second atom in the quadruplet) is the atom of symmetry in the 3  $\chi$  angles.



The subscripts on the various  $\theta$ s refer to different combinations of 3 atoms (I,J,K,L) used to form a particular angle. E.g.  $\theta_{ijl}$  is the angle formed by atoms I,J,L with J in the middle.  $\theta_1, \theta_2, \theta_3$  are the equilibrium positions of those angles. Again, atom J (the second atom in the quadruplet) is the atom of symmetry in the theta angles, since it is always the center atom.

Since atom J is the atom of symmetry, normally the bonds J-I, J-K, J-L would exist for an improper to be defined between the 4 atoms, but this is not required.

See ([Sun](#)) for a description of the COMPASS class2 force field.

Coefficients for the  $E_i$  and  $E_{aa}$  formulas must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands.

These are the 2 coefficients for the  $E_i$  formula:

- $K$  (energy)
- $\chi_0$  (degrees)

$\chi_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence  $K$  is effectively energy per radian<sup>2</sup>.

For the  $E_{aa}$  formula, each line in a *improper\_coeff* command in the input script lists 7 coefficients, the first of which is *aa* to indicate they are AngleAngle coefficients. In a data file, these coefficients should be listed under a *AngleAngle Coeffs* heading and you must leave out the *aa*, i.e. only list 6 coefficients after the improper type.

- *aa*
- $M_1$  (energy)
- $M_2$  (energy)
- $M_3$  (energy)
- $\theta_1$  (degrees)
- $\theta_2$  (degrees)
- $\theta_3$  (degrees)

The  $\theta$  values are specified in degrees, but LAMMPS converts them to radians internally; hence the various  $M$  are effectively energy per radian<sup>2</sup>.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 23.2.4 Restrictions

This improper style can only be used if LAMMPS was built with the CLASS2 package. See the *Build package* doc page for more info.

### 23.2.5 Related commands

*improper\_coeff*

### 23.2.6 Default

none

---

(Sun) Sun, J Phys Chem B 102, 7338-7364 (1998).

## 23.3 improper\_style cossq command

Accelerator Variants: *cossq/omp*

### 23.3.1 Syntax

```
improper_style cossq
```

### 23.3.2 Examples

```
improper_style cossq
improper_coeff 1 4.0 0.0
```

### 23.3.3 Description

The *cossq* improper style uses the potential

$$E = \frac{1}{2}K \cos^2(\chi - \chi_0)$$

where  $\chi$  is the improper angle,  $\chi_0$  is its equilibrium value, and  $K$  is a prefactor.

If the 4 atoms in an improper quadruplet (listed in the data file read by the *read\_data* command) are ordered I,J,K,L then  $\chi$  is the angle between the plane of I,J,K and the plane of J,K,L. Alternatively, you can think of atoms J,K,L as being in a plane, and atom I above the plane, and  $\chi$  as a measure of how far out-of-plane I is with respect to the other 3 atoms.

Note that defining 4 atoms to interact in this way, does not mean that bonds necessarily exist between I-J, J-K, or K-L, as they would in a linear dihedral. Normally, the bonds I-J, I-K, I-L would exist for an improper to be defined between the 4 atoms.

The following coefficients must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
  - $\chi_0$  (degrees)
- 

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 23.3.4 Restrictions

This improper style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) doc page for more info.

### 23.3.5 Related commands

*improper\_coeff*

### 23.3.6 Default

none

## 23.4 improper\_style cvff command

Accelerator Variants: *cvff/intel*, *cvff/omp*

### 23.4.1 Syntax

```
improper_style cvff
```

### 23.4.2 Examples

```
improper_style cvff
improper_coeff 1 80.0 -1 4
```

### 23.4.3 Description

The *cvff* improper style uses the potential

$$E = K[1 + d \cos(n\phi)]$$

where phi is the improper dihedral angle.

If the 4 atoms in an improper quadruplet (listed in the data file read by the *read\_data* command) are ordered I,J,K,L then the improper dihedral angle is between the plane of I,J,K and the plane of J,K,L. Note that because this is effectively a dihedral angle, the formula for this improper style is the same as for *dihedral\_style harmonic*.

Note that defining 4 atoms to interact in this way, does not mean that bonds necessarily exist between I-J, J-K, or K-L, as they would in a linear dihedral. Normally, the bonds I-J, I-K, I-L would exist for an improper to be defined between the 4 atoms.

The following coefficients must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- *K* (energy)
- *d* (+1 or -1)
- *n* (0,1,2,3,4,6)

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 23.4.4 Restrictions

This improper style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

### 23.4.5 Related commands

*improper\_coeff*

### 23.4.6 Default

none

## 23.5 improper\_style distance command

### 23.5.1 Syntax

```
improper_style distance
```

### 23.5.2 Examples

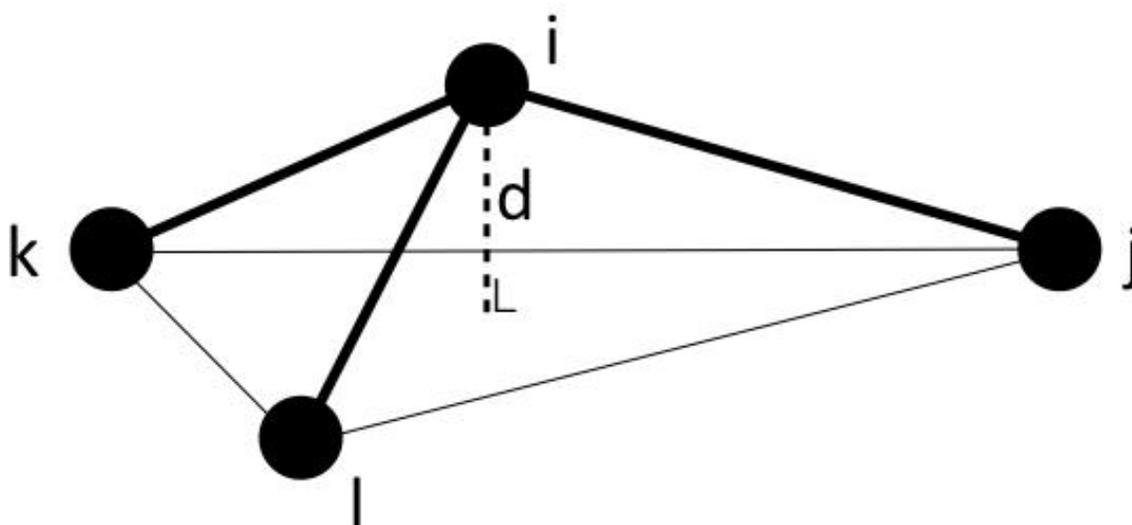
```
improper_style distance
improper_coeff 1 80.0 100.0
```

### 23.5.3 Description

The *distance* improper style uses the potential

$$E = K_2d^2 + K_4d^4$$

where  $d$  is the distance between the central atom and the plane formed by the other three atoms. If the 4 atoms in an improper quadruplet (listed in the data file read by the *read\_data* command) are ordered I,J,K,L then the I-atom is assumed to be the central atom.



Note that defining 4 atoms to interact in this way, does not mean that bonds necessarily exist between I-J, J-K, or K-L, as they would in a linear dihedral. Normally, the bonds I-J, I-K, I-L would exist for an improper to be defined between the 4 atoms.

The following coefficients must be defined for each improper type via the `improper_coeff` command as in the example above, or in the data file or restart files read by the `read_data` or `read_restart` commands:

- $K_2$  (energy/distance<sup>2</sup>)
  - $K_4$  (energy/distance<sup>4</sup>)
- 

### 23.5.4 Restrictions

This improper style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

### 23.5.5 Related commands

*improper\_coeff*

### 23.5.6 Default

none

## 23.6 improper\_style distharm command

### 23.6.1 Syntax

```
improper_style distharm
```

### 23.6.2 Examples

```
improper_style distharm
improper_coeff 1 25.0 0.5
```

### 23.6.3 Description

The *distharm* improper style uses the potential

$$E = K(d - d_0)^2$$

where  $d$  is the oriented distance between the central atom and the plane formed by the other three atoms. If the 4 atoms in an improper quadruplet (listed in the data file read by the *read\_data* command) are ordered I,J,K,L then the L-atom is assumed to be the central atom. Note that this is different from the convention used in the `improper_style` distance. The distance  $d$  is oriented and can take on negative values. This may lead to unwanted behavior if  $d_0$  is not equal to zero.

The following coefficients must be defined for each improper type via the `improper_coeff` command as in the example above, or in the data file or restart files read by the `read_data` or `read_restart` commands:

- $K$  (energy/distance<sup>2</sup>)

- $d_0$  (distance)

### 23.6.4 Restrictions

This improper style can only be used if LAMMPS was built with the YAFF package. See the *Build package* doc page for more info.

### 23.6.5 Related commands

*improper\_coeff*

### 23.6.6 Default

none

## 23.7 improper\_style fourier command

Accelerator Variants: *fourier/omp*

### 23.7.1 Syntax

```
improper_style fourier
```

### 23.7.2 Examples

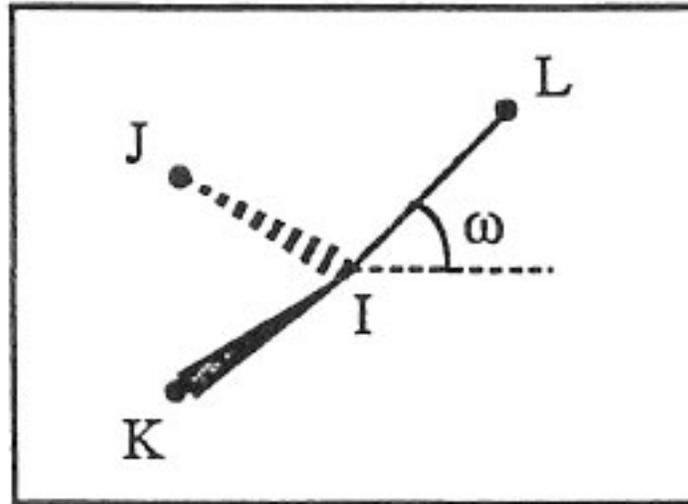
```
improper_style fourier
improper_coeff 1 100.0 0.0 1.0 0.5 1
```

### 23.7.3 Description

The *fourier* improper style uses the following potential:

$$E = K[C_0 + C_1 \cos(\omega) + C_2 \cos(2\omega)]$$

where K is the force constant, C0, C1, C2 are dimensionless coefficients, and omega is the angle between the IL axis and the IJK plane:



If all parameter (see below) is not zero, the all the three possible angles will taken in account.

The following coefficients must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $C_0$  (unitless)
- $C_1$  (unitless)
- $C_2$  (unitless)
- all (0 or 1, optional)

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---



### 23.7.4 Restrictions

This angle style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

### 23.7.5 Related commands

*improper\_coeff*

### 23.7.6 Default

none

## 23.8 improper\_style harmonic command

Accelerator Variants: *harmonic/intel*, *harmonic/kk*, *harmonic/omp*

### 23.8.1 Syntax

```
improper_style harmonic
```

### 23.8.2 Examples

```
improper_style harmonic
improper_coeff 1 100.0 0
```

### 23.8.3 Description

The *harmonic* improper style uses the potential

$$E = K(\chi - \chi_0)^2$$

where  $\chi$  is the improper angle,  $\chi_0$  is its equilibrium value, and  $K$  is a prefactor. Note that the usual 1/2 factor is included in  $K$ .

If the 4 atoms in an improper quadruplet (listed in the data file read by the *read\_data* command) are ordered I,J,K,L then  $\chi$  is the angle between the plane of I,J,K and the plane of J,K,L. Alternatively, you can think of atoms J,K,L as being in a plane, and atom I above the plane, and  $\chi$  as a measure of how far out-of-plane I is with respect to the other 3 atoms.

Note that defining 4 atoms to interact in this way, does not mean that bonds necessarily exist between I-J, J-K, or K-L, as they would in a linear dihedral. Normally, the bonds I-J, I-K, I-L would exist for an improper to be defined between the 4 atoms.

The following coefficients must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)

- $\chi_0$  (degrees)

$\chi_0$  is specified in degrees, but LAMMPS converts it to radians internally; hence  $K$  is effectively energy per radian<sup>2</sup>.

---

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

## 23.8.4 Restrictions

This improper style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) doc page for more info.

## 23.8.5 Related commands

*improper\_coeff*

## 23.8.6 Default

none

# 23.9 improper\_style hybrid command

## 23.9.1 Syntax

```
improper_style hybrid style1 style2 ...
```

- style1,style2 = list of one or more improper styles

## 23.9.2 Examples

```
improper_style hybrid harmonic cvff
improper_coeff 1 harmonic 120.0 30
improper_coeff 2 cvff 20.0 -1 2
```

### 23.9.3 Description

The *hybrid* style enables the use of multiple improper styles in one simulation. An improper style is assigned to each improper type. For example, improvers in a polymer flow (of improper type 1) could be computed with a *harmonic* potential and improvers in the wall boundary (of improper type 2) could be computed with a *cvff* potential. The assignment of improper type to style is made via the *improper\_coeff* command or in the data file.

In the *improper\_coeff* command, the first coefficient sets the improper style and the remaining coefficients are those appropriate to that style. In the example above, the 2 *improper\_coeff* commands would set improvers of improper type 1 to be computed with a *harmonic* potential with coefficients 120.0, 30 for  $K$ ,  $\chi_0$ . Improper type 2 would be computed with a *cvff* potential with coefficients 20.0, -1, 2 for  $K$ ,  $d$ , and  $n$ , respectively.

If improper coefficients are specified in the data file read via the *read\_data* command, then the same rule applies. E.g. “harmonic” or “cvff”, must be added after the improper type, for each line in the “Improper Coeffs” section, e.g.

```
Improper Coeffs
1 harmonic 120.0 30
2 cvff 20.0 -1 2
...
```

If *class2* is one of the improper hybrid styles, the same rule holds for specifying additional AngleAngle coefficients either via the input script or in the data file. I.e. *class2* must be added to each line after the improper type. For lines in the AngleAngle Coeffs section of the data file for dihedral types that are not *class2*, you must use an improper style of *skip* as a placeholder, e.g.

```
AngleAngle Coeffs
1 skip
2 class2 0.0 0.0 0.0 115.06 130.01 115.06
...
```

Note that it is not necessary to use the improper style *skip* in the input script, since AngleAngle coefficients need not be specified at all for improper types that are not *class2*.

An improper style of *none* can be specified as the second argument to the *improper\_coeff* command, if you desire to turn off certain improper types.

### 23.9.4 Restrictions

This improper style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

Unlike other improper styles, the hybrid improper style does not store improper coefficient info for individual sub-styles in a *binary restart files*. Thus when restarting a simulation from a restart file, you need to re-specify *improper\_coeff* commands.

## 23.9.5 Related commands

*improper\_coeff*

## 23.9.6 Default

none

## 23.10 improper\_style inversion/harmonic command

### 23.10.1 Syntax

```
improper_style inversion/harmonic
```

### 23.10.2 Examples

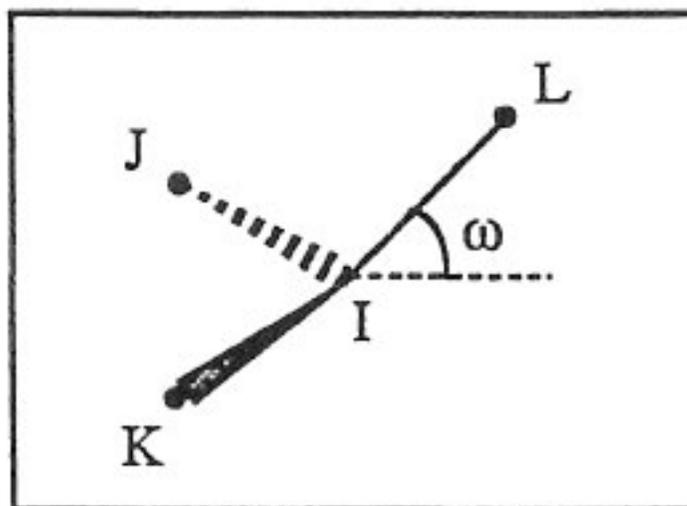
```
improper_style inversion/harmonic
improper_coeff 1 18.776340 0.000000
```

### 23.10.3 Description

The *inversion/harmonic* improper style follows the Wilson-Decius out-of-plane angle definition and uses an harmonic potential:

$$E = K (\omega - \omega_0)^2$$

where  $K$  is the force constant and  $\omega$  is the angle evaluated for all three axis-plane combinations centered around the atom I. For the IL axis and the IJK plane  $\omega$  looks as follows:



Note that the *inversion/harmonic* angle term evaluation differs to the *improper\_umbrella* due to the cyclic evaluation of all possible angles  $\omega$ .

The following coefficients must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $\omega_0$  (degrees)

If  $\omega_0 = 0$  the potential term has a single minimum for the planar structure. Otherwise it has two minima at  $\pm \omega_0$ , with a barrier in between.

---

## 23.10.4 Restrictions

This improper style can only be used if LAMMPS was built with the MOFFF package. See the *Build package* doc page for more info.

## 23.10.5 Related commands

*improper\_coeff*

## 23.10.6 Default

none

## 23.11 improper\_style none command

### 23.11.1 Syntax

```
improper_style none
```

### 23.11.2 Examples

```
improper_style none
```

### 23.11.3 Description

Using an improper style of none means improper forces and energies are not computed, even if quadruplets of improper atoms were listed in the data file read by the *read\_data* command.

See the *improper\_style zero* command for a way to calculate improper statistics, but compute no improper interactions.

### 23.11.4 Restrictions

none

### 23.11.5 Related commands

*improper\_style zero*

### 23.11.6 Default

none

## 23.12 `improper_style ring` command

Accelerator Variants: *ring/omp*

### 23.12.1 Syntax

```
improper_style ring
```

### 23.12.2 Examples

```
improper_style ring  
improper_coeff 1 8000 70.5
```

### 23.12.3 Description

The *ring* improper style uses the potential

$$E = \frac{1}{6}K (\Delta_{ijl} + \Delta_{ijk} + \Delta_{kjl})^6$$
$$\Delta_{ijl} = \cos \theta_{ijl} - \cos \theta_0$$
$$\Delta_{ijk} = \cos \theta_{ijk} - \cos \theta_0$$
$$\Delta_{kjl} = \cos \theta_{kjl} - \cos \theta_0$$

where  $K$  is a prefactor,  $\theta$  is the angle formed by the atoms specified by (i,j,k,l) indices and  $\theta_0$  its equilibrium value.

If the 4 atoms in an improper quadruplet (listed in the data file read by the *read\_data* command) are ordered i,j,k,l then  $\theta_{ijl}$  is the angle between atoms i,j and l,  $\theta_{ijk}$  is the angle between atoms i,j and k,  $\theta_{kjl}$  is the angle between atoms j,k, and l.

The “ring” improper style implements the improper potential introduced by Destree et al., in Equation (9) of (*Destree*). This potential does not affect small amplitude vibrations but is used in an ad-hoc way to prevent the onset of accidentally large amplitude fluctuations leading to the occurrence of a planar conformation of the three bonds i-j, j-k and j-l, an intermediate conformation toward the chiral inversion of a methine carbon. In the “Improvers” section of data file four atoms: i, j, k and l are specified with i,j and l lying on the backbone of the chain and k specifying the chirality of j.

The following coefficients must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
  - $\theta_0$  (degrees)
- 

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the [Accelerator packages](#) page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the [Build package](#) page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix* *command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the [Accelerator packages](#) page for more instructions on how to use the accelerated styles effectively.

---

### 23.12.4 Restrictions

This improper style can only be used if LAMMPS was built with the MOLECULE package. See the [Build package](#) doc page for more info.

### 23.12.5 Related commands

*improper\_coeff*

(Destree) M. Destree, F. Laupretre, A. Lyulin, and J.-P. Ryckaert, J Chem Phys, 112, 9632 (2000).

## 23.13 improper\_style sqdistharm command

### 23.13.1 Syntax

```
improper_style sqdistharm
```

### 23.13.2 Examples

```
improper_style sqdistharm
improper_coeff 1 50.0 0.1
```

### 23.13.3 Description

The *sqdistharm* improper style uses the potential

$$E = K(d^2 - d_0^2)^2$$

where  $d$  is the distance between the central atom and the plane formed by the other three atoms. If the 4 atoms in an improper quadruplet (listed in the data file read by the *read\_data* command) are ordered I,J,K,L then the L-atom is assumed to be the central atom. Note that this is different from the convention used in the *improper\_style* distance.

The following coefficients must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy/distance<sup>4</sup>)
- $d_0^2$  (distance<sup>2</sup>)

Note that  $d_0^2$  (in units distance<sup>2</sup>) has be provided and not  $d_0$ .

---

### 23.13.4 Restrictions

This improper style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

### 23.13.5 Related commands

*improper\_coeff*

### 23.13.6 Default

none

## 23.14 improper\_style umbrella command

Accelerator Variants: *umbrella/omp*

### 23.14.1 Syntax

```
improper_style umbrella
```



## 23.14.2 Examples

```
improper_style umbrella
improper_coeff 1 100.0 180.0
```

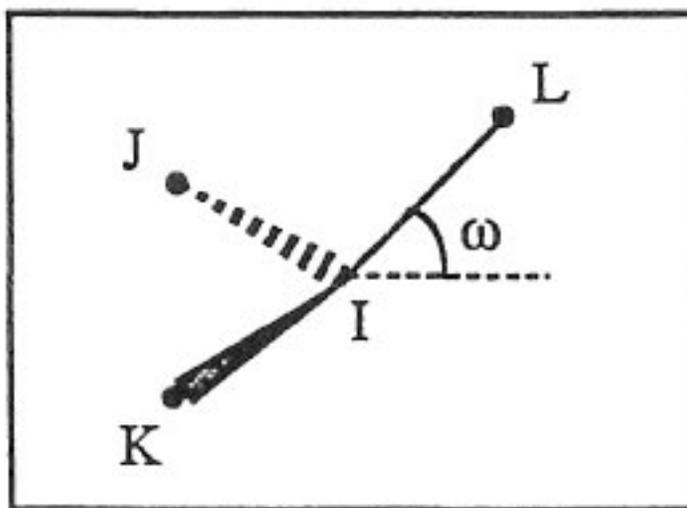
## 23.14.3 Description

The *umbrella* improper style uses the following potential, which is commonly referred to as a classic inversion and used in the *DREIDING* force field:

$$E = \frac{1}{2}K \left( \frac{1}{\sin \omega_0} \right)^2 (\cos \omega - \cos \omega_0)^2 \quad \omega_0 \neq 0^\circ$$

$$E = K(1 - \cos \omega) \quad \omega_0 = 0^\circ$$

where  $K$  is the force constant and  $\omega$  is the angle between the IL axis and the IJK plane:



If  $\omega_0 = 0$  the potential term has a minimum for the planar structure. Otherwise it has two minima at  $\omega + / - \omega_0$ , with a barrier in between.

See (*Mayo*) for a description of the DREIDING force field.

The following coefficients must be defined for each improper type via the *improper\_coeff* command as in the example above, or in the data file or restart files read by the *read\_data* or *read\_restart* commands:

- $K$  (energy)
- $\omega_0$  (degrees)

Styles with a *gpu*, *intel*, *kk*, *omp*, or *opt* suffix are functionally the same as the corresponding style without the suffix. They have been optimized to run faster, depending on your available hardware, as discussed on the *Accelerator packages* page. The accelerated styles take the same arguments and should produce the same results, except for round-off and precision issues.

These accelerated styles are part of the GPU, INTEL, KOKKOS, OPENMP, and OPT packages, respectively. They are only enabled if LAMMPS was built with those packages. See the *Build package* page for more info.

You can specify the accelerated styles explicitly in your input script by including their suffix, or you can use the *-suffix command-line switch* when you invoke LAMMPS, or you can use the *suffix* command in your input script.

See the *Accelerator packages* page for more instructions on how to use the accelerated styles effectively.

---

### 23.14.4 Restrictions

This improper style can only be used if LAMMPS was built with the MOLECULE package. See the *Build package* doc page for more info.

### 23.14.5 Related commands

*improper\_coeff*

### 23.14.6 Default

none

---

(Mayo) Mayo, Olfason, Goddard III, J Phys Chem, 94, 8897-8909 (1990),

## 23.15 improper\_style zero command

### 23.15.1 Syntax

```
improper_style zero [nocoeff]
```

### 23.15.2 Examples

```
improper_style zero  
improper_style zero nocoeff  
improper_coeff *
```

### 23.15.3 Description

Using an improper style of zero means improper forces and energies are not computed, but the geometry of improper quadruplets is still accessible to other commands.

As an example, the *compute improper/local* command can be used to compute the chi values for the list of quadruplets of improper atoms listed in the data file read by the *read\_data* command. If no improper style is defined, this command cannot be used.

The optional *nocoeff* flag allows to read data files with a *ImproperCoeff* section for any improper style. Similarly, any *improper\_coeff* commands will only be checked for the improper type number and the rest ignored.

Note that the *improper\_coeff* command must be used for all improper types, though no additional values are specified.

### 23.15.4 Restrictions

none

### 23.15.5 Related commands

none

*improper\_style none*

### 23.15.6 Default

none



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CHAPTER  
**TWENTYFOUR**

---

**DUMP STYLES**



## BIBLIOGRAPHY

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## **Part IV**

# **Indices and tables**





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### Web Browser Compatibility

The HTML version of the manual makes use of advanced features present in “modern” web browsers. This can lead to incompatibilities with older web browsers (released more than 4 years ago) and specific vendor browsers (e.g. Internet Explorer on Windows; Microsoft Edge works well though) where parts of the pages are not rendered as expected (e.g. the layout is broken or mathematical expressions not typeset). In that case we recommend to install/use a different/newer web browser or use the [PDF version of the manual](#).

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## PYTHON MODULE INDEX

|

`lammps`, 558

`lammps.formats`, 586



## Symbols

- `_LMP_DATATYPE_CONST` (C++ *enum*), 507
  - `_LMP_DATATYPE_CONST::LAMMPS_DOUBLE` (C++ *enumerator*), 507
  - `_LMP_DATATYPE_CONST::LAMMPS_DOUBLE_2D` (C++ *enumerator*), 508
  - `_LMP_DATATYPE_CONST::LAMMPS_INT` (C++ *enumerator*), 507
  - `_LMP_DATATYPE_CONST::LAMMPS_INT64` (C++ *enumerator*), 508
  - `_LMP_DATATYPE_CONST::LAMMPS_INT64_2D` (C++ *enumerator*), 508
  - `_LMP_DATATYPE_CONST::LAMMPS_INT_2D` (C++ *enumerator*), 507
  - `_LMP_DATATYPE_CONST::LAMMPS_STRING` (C++ *enumerator*), 508
  - `_LMP_STYLE_CONST` (C++ *enum*), 508
  - `_LMP_STYLE_CONST::LMP_STYLE_ATOM` (C++ *enumerator*), 508
  - `_LMP_STYLE_CONST::LMP_STYLE_GLOBAL` (C++ *enumerator*), 508
  - `_LMP_STYLE_CONST::LMP_STYLE_LOCAL` (C++ *enumerator*), 508
  - `_LMP_TYPE_CONST` (C++ *enum*), 508
  - `_LMP_TYPE_CONST::LMP_SIZE_COLS` (C++ *enumerator*), 509
  - `_LMP_TYPE_CONST::LMP_SIZE_ROWS` (C++ *enumerator*), 509
  - `_LMP_TYPE_CONST::LMP_SIZE_VECTOR` (C++ *enumerator*), 508
  - `_LMP_TYPE_CONST::LMP_TYPE_ARRAY` (C++ *enumerator*), 508
  - `_LMP_TYPE_CONST::LMP_TYPE_SCALAR` (C++ *enumerator*), 508
  - `_LMP_TYPE_CONST::LMP_TYPE_VECTOR` (C++ *enumerator*), 508
  - `__version__` (in module *lammps*), 558
- A**
- `accelerator_config` (*lammps.lammps* property), 567
  - `angle_coeff`, 715
  - `angle_style`, 716
  - `angle_style amoeba`, 2421
  - `angle_style charmm`, 2423
  - `angle_style charmm/intel`, 2423
  - `angle_style charmm/kk`, 2423
  - `angle_style charmm/omp`, 2423
  - `angle_style class2`, 2424
  - `angle_style class2/kk`, 2424
  - `angle_style class2/omp`, 2424
  - `angle_style class2/p6`, 2424
  - `angle_style cosine`, 2427
  - `angle_style cosine/buck6d`, 2428
  - `angle_style cosine/delta`, 2429
  - `angle_style cosine/delta/omp`, 2429
  - `angle_style cosine/kk`, 2427
  - `angle_style cosine/omp`, 2427
  - `angle_style cosine/periodic`, 2430
  - `angle_style cosine/periodic/omp`, 2430
  - `angle_style cosine/shift`, 2431
  - `angle_style cosine/shift/exp`, 2433
  - `angle_style cosine/shift/exp/omp`, 2433
  - `angle_style cosine/shift/omp`, 2431
  - `angle_style cosine/squared`, 2434
  - `angle_style cosine/squared/omp`, 2434
  - `angle_style cross`, 2435
  - `angle_style dipole`, 2437
  - `angle_style dipole/omp`, 2437
  - `angle_style fourier`, 2439
  - `angle_style fourier/omp`, 2439
  - `angle_style fourier/simple`, 2440
  - `angle_style fourier/simple/omp`, 2440
  - `angle_style gaussian`, 2441
  - `angle_style harmonic`, 2442
  - `angle_style harmonic/intel`, 2442
  - `angle_style harmonic/kk`, 2442
  - `angle_style harmonic/omp`, 2442
  - `angle_style hybrid`, 2444
  - `angle_style mesocnt`, 2445
  - `angle_style mm3`, 2447
  - `angle_style none`, 2448
  - `angle_style quartic`, 2449
  - `angle_style quartic/omp`, 2449
  - `angle_style spica`, 2450

angle\_style spica/omp, 2450  
angle\_style table, 2452  
angle\_style table/omp, 2452  
angle\_style zero, 2454  
angular\_momentum (*lammps.Atom property*), 581  
Atom (*class in lammps*), 580  
Atom2D (*class in lammps*), 581  
atom\_modify, 718  
atom\_style, 721  
AtomList (*class in lammps*), 580  
atoms (*lammps.PyLammps property*), 579  
available\_ids() (*lammps.lammps method*), 568  
available\_plugins() (*lammps.lammps method*), 569  
available\_styles() (*lammps.lammps method*), 568

## B

balance, 726  
bond\_coeff, 733  
bond\_style, 735  
bond\_style bpm/rotational, 2387  
bond\_style bpm/spring, 2390  
bond\_style class2, 2393  
bond\_style class2/kk, 2393  
bond\_style class2/omp, 2393  
bond\_style fene, 2394  
bond\_style fene/expand, 2396  
bond\_style fene/expand/omp, 2396  
bond\_style fene/intel, 2394  
bond\_style fene/kk, 2394  
bond\_style fene/nm, 2394  
bond\_style fene/omp, 2394  
bond\_style gaussian, 2397  
bond\_style gromos, 2398  
bond\_style gromos/omp, 2398  
bond\_style harmonic, 2400  
bond\_style harmonic/intel, 2400  
bond\_style harmonic/kk, 2400  
bond\_style harmonic/omp, 2400  
bond\_style harmonic/shift, 2401  
bond\_style harmonic/shift/cut, 2402  
bond\_style harmonic/shift/cut/omp, 2402  
bond\_style harmonic/shift/omp, 2401  
bond\_style hybrid, 2403  
bond\_style mesocnt, 2405  
bond\_style mm3, 2406  
bond\_style morse, 2407  
bond\_style morse/omp, 2407  
bond\_style none, 2408  
bond\_style nonlinear, 2409  
bond\_style nonlinear/omp, 2409  
bond\_style oxdna/fene, 2410  
bond\_style oxdna2/fene, 2410  
bond\_style oxrna2/fene, 2410  
bond\_style quartic, 2412

bond\_style quartic/omp, 2412  
bond\_style special, 2414  
bond\_style table, 2416  
bond\_style table/omp, 2416  
bond\_style zero, 2418  
bond\_write, 737  
boundary, 738  
box, 740

## C

change\_box, 741  
charge (*lammps.Atom property*), 581  
clear, 746  
clear\_cmd\_history() (*lammps.PyLammps method*), 578  
close() (*fortran subroutine*), 532  
close() (*lammps.lammps method*), 559  
close() (*lammps.PyLammps method*), 577  
comm\_modify, 746  
comm\_style, 749  
command() (*fortran subroutine*), 532  
command() (*lammps.lammps method*), 560  
command() (*lammps.PyLammps method*), 578  
commands\_list() (*fortran subroutine*), 533  
commands\_list() (*lammps.lammps method*), 560  
commands\_string() (*fortran subroutine*), 533  
commands\_string() (*lammps.lammps method*), 560  
communication (*lammps.PyLammps property*), 579  
compute, 750  
compute ackland/atom, 1721  
compute adf, 1722  
compute aggregate/atom, 1756  
compute angle, 1725  
compute angle/local, 1726  
compute angmom/chunk, 1728  
compute ave/sphere/atom, 1730  
compute ave/sphere/atom/kk, 1730  
compute basal/atom, 1732  
compute body/local, 1733  
compute bond, 1735  
compute bond/local, 1736  
compute born/matrix, 1739  
compute centro/atom, 1742  
compute centroid/stress/atom, 1915  
compute chunk/atom, 1744  
compute chunk/spread/atom, 1752  
compute cluster/atom, 1756  
compute cna/atom, 1758  
compute cnp/atom, 1759  
compute com, 1761  
compute com/chunk, 1762  
compute contact/atom, 1764  
compute coord/atom, 1765  
compute coord/atom/kk, 1765



compute damage/atom, 1767  
compute dihedral, 1768  
compute dihedral/local, 1769  
compute dilatation/atom, 1771  
compute dipole, 1772  
compute dipole/chunk, 1773  
compute displace/atom, 1775  
compute dpd, 1777  
compute dpd/atom, 1779  
compute edpd/temp/atom, 1780  
compute efield/atom, 1781  
compute entropy/atom, 1782  
compute erotate/asphere, 1784  
compute erotate/rigid, 1786  
compute erotate/sphere, 1787  
compute erotate/sphere/atom, 1788  
compute event/displace, 1789  
compute fabric, 1790  
compute fep, 1792  
compute fep/ta, 1796  
compute force/tally, 1923  
compute fragment/atom, 1756  
compute global/atom, 1798  
compute group/group, 1801  
compute gyration, 1803  
compute gyration/chunk, 1805  
compute gyration/shape, 1807  
compute gyration/shape/chunk, 1808  
compute heat/flux, 1810  
compute heat/flux/tally, 1923  
compute heat/flux/virial/tally, 1923  
compute hexorder/atom, 1813  
compute hma, 1815  
compute improper, 1818  
compute improper/local, 1819  
compute inertia/chunk, 1821  
compute ke, 1822  
compute ke/atom, 1823  
compute ke/atom/eff, 1824  
compute ke/eff, 1825  
compute ke/rigid, 1827  
compute mesont, 1828  
compute mliap, 1829  
compute momentum, 1831  
compute msd, 1832  
compute msd/chunk, 1834  
compute msd/nongauss, 1836  
compute nbond/atom, 1837  
compute omega/chunk, 1838  
compute orientorder/atom, 1839  
compute orientorder/atom/kk, 1839  
compute pair, 1842  
compute pair/local, 1844  
compute pe, 1846  
compute pe/atom, 1848  
compute pe/mol/tally, 1923  
compute pe/tally, 1923  
compute plasticity/atom, 1849  
compute pressure, 1851  
compute pressure/uef, 1853  
compute property/atom, 1854  
compute property/chunk, 1857  
compute property/local, 1859  
compute ptm/atom, 1862  
compute rdf, 1864  
compute reduce, 1867  
compute reduce/chunk, 1870  
compute reduce/region, 1867  
compute rigid/local, 1873  
compute saed, 1876  
compute slice, 1879  
compute smd/contact/radius, 1881  
compute smd/damage, 1882  
compute smd/hourglass/error, 1883  
compute smd/internal/energy, 1884  
compute smd/plastic/strain, 1885  
compute smd/plastic/strain/rate, 1886  
compute smd/rho, 1887  
compute smd/tlsph/defgrad, 1888  
compute smd/tlsph/dt, 1889  
compute smd/tlsph/num/neighs, 1890  
compute smd/tlsph/shape, 1891  
compute smd/tlsph/strain, 1892  
compute smd/tlsph/strain/rate, 1893  
compute smd/tlsph/stress, 1894  
compute smd/triangle/vertices, 1895  
compute smd/ulsph/effm, 1896  
compute smd/ulsph/num/neighs, 1897  
compute smd/ulsph/strain, 1898  
compute smd/ulsph/strain/rate, 1899  
compute smd/ulsph/stress, 1900  
compute smd/vol, 1901  
compute sna/atom, 1902  
compute sna/grid, 1902  
compute sna/grid/local, 1902  
compute snad/atom, 1902  
compute snap, 1902  
compute snav/atom, 1902  
compute sph/e/atom, 1910  
compute sph/rho/atom, 1911  
compute sph/t/atom, 1912  
compute spin, 1913  
compute stress/atom, 1915  
compute stress/cartesian, 1921  
compute stress/cylinder, 1921  
compute stress/mop, 1918  
compute stress/mop/profile, 1918  
compute stress/spherical, 1921

compute stress/tally, 1923  
compute tdpd/cc/atom, 1926  
compute temp, 1927  
compute temp/asphere, 1929  
compute temp/body, 1931  
compute temp/chunk, 1933  
compute temp/com, 1936  
compute temp/cs, 1938  
compute temp/deform, 1940  
compute temp/deform/eff, 1942  
compute temp/deform/kk, 1940  
compute temp/drude, 1943  
compute temp/eff, 1944  
compute temp/kk, 1927  
compute temp/partial, 1946  
compute temp/profile, 1948  
compute temp/ramp, 1951  
compute temp/region, 1952  
compute temp/region/eff, 1954  
compute temp/rotate, 1955  
compute temp/sphere, 1957  
compute temp/uef, 1959  
compute ti, 1960  
compute torque/chunk, 1962  
compute vacf, 1963  
compute vcm/chunk, 1965  
compute viscosity/cos, 1966  
compute voronoi/atom, 1968  
compute xrd, 1971  
compute\_modify, 757  
computes (*lammps.PyLammps property*), 579  
create\_atoms, 758  
create\_atoms() (*lammps.lammps method*), 565  
create\_bonds, 765  
create\_box, 768

## D

decode\_image\_flags() (*lammps.lammps method*), 565  
delete\_atoms, 770  
delete\_bonds, 773  
dielectric, 775  
dihedral\_coeff, 776  
dihedral\_style, 778  
dihedral\_style charmm, 2457  
dihedral\_style charmm/intel, 2457  
dihedral\_style charmm/kk, 2457  
dihedral\_style charmm/omp, 2457  
dihedral\_style charmmfsw, 2457  
dihedral\_style class2, 2459  
dihedral\_style class2/kk, 2459  
dihedral\_style class2/omp, 2459  
dihedral\_style cosine/shift/exp, 2462  
dihedral\_style cosine/shift/exp/omp, 2462  
dihedral\_style fourier, 2464  
dihedral\_style fourier/intel, 2464  
dihedral\_style fourier/omp, 2464  
dihedral\_style harmonic, 2465  
dihedral\_style harmonic/intel, 2465  
dihedral\_style harmonic/kk, 2465  
dihedral\_style harmonic/omp, 2465  
dihedral\_style helix, 2466  
dihedral\_style helix/omp, 2466  
dihedral\_style hybrid, 2468  
dihedral\_style multi/harmonic, 2469  
dihedral\_style multi/harmonic/omp, 2469  
dihedral\_style nharmonic, 2470  
dihedral\_style nharmonic/omp, 2470  
dihedral\_style none, 2472  
dihedral\_style opls, 2473  
dihedral\_style opls/intel, 2473  
dihedral\_style opls/kk, 2473  
dihedral\_style opls/omp, 2473  
dihedral\_style quadratic, 2474  
dihedral\_style quadratic/omp, 2474  
dihedral\_style spherical, 2475  
dihedral\_style table, 2477  
dihedral\_style table/cut, 2477  
dihedral\_style table/omp, 2477  
dihedral\_style zero, 2481  
dimension, 780  
displace\_atoms, 781  
dump, 783  
dump atom, 783  
dump atom/adios, 794  
dump atom/gz, 783  
dump atom/mpiio, 783  
dump atom/zstd, 783  
dump cfg, 783  
dump cfg/gz, 783  
dump cfg/mpiio, 783  
dump cfg/uef, 795  
dump cfg/zstd, 783  
dump custom, 783  
dump custom/adios, 794  
dump custom/gz, 783  
dump custom/mpiio, 783  
dump custom/zstd, 783  
dump dcd, 783  
dump h5md, 796  
dump image, 798  
dump local, 783  
dump local/gz, 783  
dump local/zstd, 783  
dump molfile, 825  
dump movie, 798  
dump netcdf, 827  
dump netcdf/mpiio, 827  
dump vtk, 829

dump xtc, 783  
 dump xyz, 783  
 dump xyz/gz, 783  
 dump xyz/mpio, 783  
 dump xyz/zstd, 783  
 dump yaml, 783  
 dump\_modify, 814  
 dumps (*lammps.PyLammps property*), 579  
 dynamical\_matrix, 831  
 dynamical\_matrix/kk, 831

## E

echo, 833  
 enable\_cmd\_history (*lammps.PyLammps property*), 578  
 encode\_image\_flags() (*lammps.lammps method*), 564  
 eval() (*lammps.PyLammps method*), 580  
 extract\_atom() (*lammps.lammps method*), 562  
 extract\_atom() (*lammps.numpy\_wrapper.numpy\_wrapper method*), 574  
 extract\_atom\_datatype() (*lammps.lammps method*), 562  
 extract\_box() (*lammps.lammps method*), 560  
 extract\_compute() (*lammps.lammps method*), 563  
 extract\_compute() (*lammps.numpy\_wrapper.numpy\_wrapper method*), 574  
 extract\_fix() (*lammps.lammps method*), 563  
 extract\_fix() (*lammps.numpy\_wrapper.numpy\_wrapper method*), 574  
 extract\_global() (*lammps.lammps method*), 561  
 extract\_global\_datatype() (*lammps.lammps method*), 561  
 extract\_setting() (*lammps.lammps method*), 561  
 extract\_variable() (*lammps.lammps method*), 563  
 extract\_variable() (*lammps.numpy\_wrapper.numpy\_wrapper method*), 575

## F

file() (*fortran subroutine*), 532  
 file() (*lammps.lammps method*), 560  
 file() (*lammps.PyLammps method*), 578  
 finalize() (*lammps.lammps method*), 559  
 find() (*lammps.NeighList method*), 585  
 find() (*lammps.numpy\_wrapper.NumPyNeighList method*), 585  
 find\_compute\_neighlist() (*lammps.lammps method*), 573  
 find\_fix\_neighlist() (*lammps.lammps method*), 573  
 find\_pair\_neighlist() (*lammps.lammps method*), 572  
 fix, 834  
 fix accelerate/cos, 1127  
 fix acks2/reaxff, 1128  
 fix acks2/reaxff/kk, 1128  
 fix adapt, 1130  
 fix adapt/fep, 1136  
 fix addforce, 1140  
 fix addtorque, 1143  
 fix amoeba/bitorsion, 1144  
 fix amoeba/pitorsion, 1146  
 fix append/atoms, 1149  
 fix atc, 1151  
 fix atom/swap, 1157  
 fix ave/atom, 1159  
 fix ave/chunk, 1162  
 fix ave/correlate, 1169  
 fix ave/correlate/long, 1174  
 fix ave/histo, 1176  
 fix ave/histo/weight, 1176  
 fix ave/time, 1181  
 fix aveforce, 1186  
 fix balance, 1188  
 fix bocs, 1194  
 fix bond/break, 1196  
 fix bond/create, 1198  
 fix bond/create/angle, 1198  
 fix bond/react, 1201  
 fix bond/swap, 1210  
 fix box/relax, 1214  
 fix brownian, 1219  
 fix brownian/asphere, 1219  
 fix brownian/sphere, 1219  
 fix charge/regulation, 1222  
 fix cmap, 1226  
 fix colvars, 1228  
 fix controller, 1231  
 fix damping/cundall, 1234  
 fix deform, 1236  
 fix deform/kk, 1236  
 fix deposit, 1244  
 fix dpd/energy, 1248  
 fix dpd/energy/kk, 1248  
 fix drag, 1251  
 fix drude, 1252  
 fix drude/transform/direct, 1253  
 fix drude/transform/inverse, 1253  
 fix dt/reset, 1256  
 fix edpd/source, 1249  
 fix efield, 1258  
 fix ehex, 1260  
 fix electrode/conp, 1263  
 fix electrode/conp/intel, 1263  
 fix electrode/conq, 1263  
 fix electrode/conq/intel, 1263  
 fix electrode/thermo, 1263  
 fix electrode/thermo/intel, 1263  
 fix electron/stopping, 1266  
 fix electron/stopping/fit, 1266

fix enforce2d, 1269  
fix enforce2d/kk, 1269  
fix eos/cv, 1270  
fix eos/table, 1271  
fix eos/table/rx, 1273  
fix eos/table/rx/kk, 1273  
fix evaporate, 1276  
fix external, 1278  
fix ffl, 1280  
fix filter/corotate, 1282  
fix flow/gauss, 1283  
fix freeze, 1286  
fix freeze/kk, 1286  
fix gcmc, 1287  
fix gld, 1293  
fix gle, 1296  
fix gravity, 1298  
fix gravity/kk, 1298  
fix gravity/omp, 1298  
fix grem, 1301  
fix halt, 1302  
fix heat, 1305  
fix hyper/global, 1307  
fix hyper/local, 1310  
fix imd, 1317  
fix indent, 1320  
fix ipi, 1323  
fix langevin, 1324  
fix langevin/drude, 1329  
fix langevin/eff, 1333  
fix langevin/kk, 1324  
fix langevin/spin, 1335  
fix latte, 1336  
fix lb/fluid, 1339  
fix lb/momentum, 1345  
fix lb/viscous, 1347  
fix lineforce, 1348  
fix manifoldforce, 1349  
fix mdi/qm, 1350  
fix meso/move, 1354  
fix mol/swap, 1422  
fix momentum, 1424  
fix momentum/chunk, 1424  
fix momentum/kk, 1424  
fix move, 1426  
fix mscg, 1430  
fix msst, 1432  
fix mvv/dpd, 1435  
fix mvv/edpd, 1435  
fix mvv/tdpd, 1435  
fix neb, 1437  
fix neb/spin, 1440  
fix nph, 1441  
fix nph/asphere, 1456  
fix nph/asphere/omp, 1456  
fix nph/body, 1459  
fix nph/eff, 1450  
fix nph/kk, 1441  
fix nph/omp, 1441  
fix nph/sphere, 1461  
fix nph/sphere/omp, 1461  
fix nphug, 1463  
fix nphug/omp, 1463  
fix npt, 1441  
fix npt/asphere, 1466  
fix npt/asphere/omp, 1466  
fix npt/body, 1469  
fix npt/cauchy, 1471  
fix npt/eff, 1450  
fix npt/gpu, 1441  
fix npt/intel, 1441  
fix npt/kk, 1441  
fix npt/omp, 1441  
fix npt/sphere, 1479  
fix npt/sphere/omp, 1479  
fix npt/uef, 1453  
fix numdiff, 1481  
fix numdiff/virial, 1483  
fix nve, 1485  
fix nve/asphere, 1486  
fix nve/asphere/gpu, 1486  
fix nve/asphere/intel, 1486  
fix nve/asphere/noforce, 1487  
fix nve/awpmd, 1489  
fix nve/body, 1490  
fix nve/bpm/sphere, 1491  
fix nve/dot, 1492  
fix nve/dotc/langevin, 1493  
fix nve/eff, 1495  
fix nve/gpu, 1485  
fix nve/intel, 1485  
fix nve/kk, 1485  
fix nve/limit, 1496  
fix nve/line, 1498  
fix nve/manifold/rattle, 1499  
fix nve/noforce, 1500  
fix nve/omp, 1485  
fix nve/sphere, 1502  
fix nve/sphere/kk, 1502  
fix nve/sphere/omp, 1502  
fix nve/spin, 1503  
fix nve/tri, 1505  
fix nvk, 1506  
fix nvt, 1441  
fix nvt/asphere, 1507  
fix nvt/asphere/omp, 1507  
fix nvt/body, 1509  
fix nvt/eff, 1450

fix nvt/gpu, 1441  
fix nvt/intel, 1441  
fix nvt/kk, 1441  
fix nvt/manifold/rattle, 1511  
fix nvt/omp, 1441  
fix nvt/sllod, 1512  
fix nvt/sllod/eff, 1515  
fix nvt/sllod/intel, 1512  
fix nvt/sllod/kk, 1512  
fix nvt/sllod/omp, 1512  
fix nvt/sphere, 1516  
fix nvt/sphere/omp, 1516  
fix nvt/uef, 1453  
fix oneway, 1519  
fix orient/bcc, 1520  
fix orient/eco, 1523  
fix orient/fcc, 1520  
fix pafi, 1525  
fix pair, 1527  
fix phonon, 1529  
fix pimd, 1532  
fix planeforce, 1535  
fix plumed, 1536  
fix poems, 1538  
fix polarize/bem/gmres, 1540  
fix polarize/bem/icc, 1540  
fix polarize/functional, 1540  
fix pour, 1543  
fix precession/spin, 1546  
fix press/berendsen, 1549  
fix print, 1553  
fix propel/self, 1555  
fix property/atom, 1557  
fix property/atom/kk, 1557  
fix python/invoke, 1561  
fix python/move, 1563  
fix qbmsst, 1564  
fix qeq/comb, 1572  
fix qeq/comb/omp, 1572  
fix qeq/dynamic, 1568  
fix qeq/fire, 1568  
fix qeq/point, 1568  
fix qeq/reaxff, 1574  
fix qeq/reaxff/kk, 1574  
fix qeq/reaxff/omp, 1574  
fix qeq/shielded, 1568  
fix qeq/slater, 1568  
fix qmmm, 1576  
fix qtb, 1577  
fix rattle, 1616  
fix reaxff/bonds, 1580  
fix reaxff/bonds/kk, 1580  
fix reaxff/species, 1582  
fix reaxff/species/kk, 1582  
fix recenter, 1585  
fix restrain, 1587  
fix rhok, 1591  
fix rigid, 1592  
fix rigid/meso, 1603  
fix rigid/nph, 1592  
fix rigid/nph/omp, 1592  
fix rigid/nph/small, 1592  
fix rigid/npt, 1592  
fix rigid/npt/omp, 1592  
fix rigid/npt/small, 1592  
fix rigid/nve, 1592  
fix rigid/nve/omp, 1592  
fix rigid/nve/small, 1592  
fix rigid/nvt, 1592  
fix rigid/nvt/omp, 1592  
fix rigid/nvt/small, 1592  
fix rigid/omp, 1592  
fix rigid/small, 1592  
fix rigid/small/omp, 1592  
fix rx, 1608  
fix rx/kk, 1608  
fix saed/vtk, 1611  
fix setforce, 1614  
fix setforce/kk, 1614  
fix setforce/spin, 1614  
fix shake, 1616  
fix shake/kk, 1616  
fix shardlow, 1620  
fix shardlow/kk, 1620  
fix smd, 1622  
fix smd/adjust\_dt, 1624  
fix smd/integrate\_tlsph, 1625  
fix smd/integrate\_ulsph, 1626  
fix smd/move\_tri\_surf, 1627  
fix smd/setvel, 1629  
fix smd/wall\_surface, 1630  
fix sph, 1632  
fix sph/stationary, 1633  
fix spring, 1634  
fix spring/chunk, 1636  
fix spring/rg, 1637  
fix spring/self, 1639  
fix srd, 1640  
fix store/force, 1645  
fix store/state, 1647  
fix tdpd/source, 1249  
fix temp/berendsen, 1649  
fix temp/csld, 1651  
fix temp/csvr, 1651  
fix temp/rescale, 1654  
fix temp/rescale/eff, 1656  
fix tfmc, 1658  
fix tgnpt/drude, 1660

fix tgnvt/drude, 1660  
 fix thermal/conductivity, 1664  
 fix ti/spring, 1666  
 fix tmd, 1669  
 fix ttm, 1671  
 fix ttm/grid, 1671  
 fix ttm/mod, 1671  
 fix tune/kspace, 1676  
 fix vector, 1678  
 fix viscosity, 1680  
 fix viscous, 1682  
 fix viscous/sphere, 1684  
 fix wall/body/polygon, 1691  
 fix wall/body/polyhedron, 1693  
 fix wall/colloid, 1686  
 fix wall/ees, 1695  
 fix wall/gran, 1697  
 fix wall/gran/region, 1701  
 fix wall/harmonic, 1686  
 fix wall/lj1043, 1686  
 fix wall/lj126, 1686  
 fix wall/lj93, 1686  
 fix wall/lj93/kk, 1686  
 fix wall/morse, 1686  
 fix wall/piston, 1705  
 fix wall/reflect, 1706  
 fix wall/reflect/kk, 1706  
 fix wall/reflect/stochastic, 1709  
 fix wall/region, 1711  
 fix wall/region/ees, 1695  
 fix wall/srd, 1714  
 fix widom, 1717  
 fix\_external\_get\_force() (lammps.lammps  
     method), 570  
 fix\_external\_get\_force() (lammps.numpy\_wrapper.numpy\_wrapper  
     method), 575  
 fix\_external\_set\_energy\_global() (lammps.lammps method), 570  
 fix\_external\_set\_energy\_peratom() (lammps.lammps method), 570  
 fix\_external\_set\_energy\_peratom() (lammps.numpy\_wrapper.numpy\_wrapper  
     method), 576  
 fix\_external\_set\_vector() (lammps.lammps  
     method), 571  
 fix\_external\_set\_vector\_length() (lammps.lammps method), 571  
 fix\_external\_set\_virial\_global() (lammps.lammps method), 570  
 fix\_external\_set\_virial\_peratom() (lammps.lammps method), 571  
 fix\_external\_set\_virial\_peratom() (lammps.numpy\_wrapper.numpy\_wrapper  
     method), 576  
 fix\_modify, 843  
 fix\_modify AtC add\_molecule, 1357  
 fix\_modify AtC add\_species, 1358  
 fix\_modify AtC atom\_element\_map, 1359  
 fix\_modify AtC atom\_weight, 1360  
 fix\_modify AtC atomic\_charge, 1361  
 fix\_modify AtC boundary\_type, 1364  
 fix\_modify AtC boundary\_dynamics, 1362  
 fix\_modify AtC boundary\_faceset, 1363  
 fix\_modify AtC computes, 1376  
 fix\_modify AtC consistent\_fe\_initialization,  
     1364  
 fix\_modify AtC control localized\_lambda, 1365  
 fix\_modify AtC control lumped\_lambda\_solve,  
     1387  
 fix\_modify AtC control mask\_direction, 1388  
 fix\_modify AtC control momentum, 1366  
 fix\_modify AtC control thermal, 1367  
 fix\_modify AtC decomposition, 1369  
 fix\_modify AtC equilibrium\_start, 1370  
 fix\_modify AtC extrinsic  
     electron\_integration, 1370  
 fix\_modify AtC extrinsic exchange, 1371  
 fix\_modify AtC fe\_md\_boundary, 1372  
 fix\_modify AtC fields, 1377  
 fix\_modify AtC filter, 1416  
 fix\_modify AtC filter scale, 1373  
 fix\_modify AtC filter type, 1374  
 fix\_modify AtC fix, 1375  
 fix\_modify AtC fix\_flux, 1375  
 fix\_modify AtC gradients, 1379  
 fix\_modify AtC initial, 1384  
 fix\_modify AtC internal\_element\_set, 1385  
 fix\_modify AtC internal\_quadrature, 1386  
 fix\_modify AtC kernel, 1380  
 fix\_modify AtC kernel\_bandwidth, 1387  
 fix\_modify AtC mass\_matrix, 1389  
 fix\_modify AtC material, 1390  
 fix\_modify AtC mesh add\_to\_nodeset, 1390  
 fix\_modify AtC mesh create, 1391  
 fix\_modify AtC mesh create\_elementset, 1392  
 fix\_modify AtC mesh create\_faceset box, 1393  
 fix\_modify AtC mesh create\_faceset plane,  
     1394  
 fix\_modify AtC mesh create\_nodeset, 1395  
 fix\_modify AtC mesh delete\_elements, 1396  
 fix\_modify AtC mesh nodeset\_to\_elementset,  
     1397  
 fix\_modify AtC mesh output, 1397  
 fix\_modify AtC mesh quadrature, 1398  
 fix\_modify AtC mesh read, 1399  
 fix\_modify AtC mesh write, 1400  
 fix\_modify AtC on\_the\_fly, 1382

- fix\_modify AtC output, 1401  
 fix\_modify AtC output boundary\_integral, 1402  
 fix\_modify AtC output contour\_integral, 1403  
 fix\_modify AtC output nodeset, 1404  
 fix\_modify AtC output volume\_integral, 1405  
 fix\_modify AtC pair\_interactions, 1405  
 fix\_modify AtC poisson\_solver, 1406  
 fix\_modify AtC rates, 1382  
 fix\_modify AtC read\_restart, 1407  
 fix\_modify AtC remove\_molecule, 1408  
 fix\_modify AtC remove\_source, 1409  
 fix\_modify AtC remove\_species, 1409  
 fix\_modify AtC reset\_atomic\_reference\_positions, 1410  
 fix\_modify AtC reset\_time, 1411  
 fix\_modify AtC sample\_frequency, 1412  
 fix\_modify AtC set reference\_potential\_energy, 1412  
 fix\_modify AtC source, 1413  
 fix\_modify AtC source\_integration, 1414  
 fix\_modify AtC temperature\_definition, 1415  
 fix\_modify AtC time\_integration, 1417  
 fix\_modify AtC track\_displacement, 1418  
 fix\_modify AtC unfix, 1419  
 fix\_modify AtC unfix\_flux, 1419  
 fix\_modify AtC write\_atom\_weights, 1420  
 fix\_modify AtC write\_restart, 1421  
 fixes (*lammps.PyLammps property*), 579  
 flush\_buffers() (*lammps.lammps method*), 564  
 force (*lammps.Atom property*), 581  
 force (*lammps.Atom2D property*), 582  
 force\_timeout() (*lammps.lammps method*), 566
- ## G
- gather\_bonds() (*lammps.lammps method*), 564  
 gather\_bonds() (*lammps.numpy\_wrapper.numpy\_wrapper method*), 575  
 get() (*lammps.NeighList method*), 584  
 get() (*lammps.numpy\_wrapper.NumPyNeighList method*), 585  
 get\_gpu\_device\_info() (*lammps.lammps method*), 567  
 get\_mpi\_comm() (*lammps.lammps method*), 559  
 get\_natoms() (*lammps.lammps method*), 560  
 get\_neighlist() (*lammps.lammps method*), 572  
 get\_neighlist() (*lammps.numpy\_wrapper.numpy\_wrapper method*), 576  
 get\_neighlist\_element\_neighbors() (*lammps.lammps method*), 572  
 get\_neighlist\_element\_neighbors() (*lammps.numpy\_wrapper.numpy\_wrapper method*), 576  
 get\_neighlist\_size() (*lammps.lammps method*), 572  
 get\_os\_info() (*lammps.lammps method*), 559  
 get\_thermo() (*lammps.lammps method*), 561  
 group, 846  
 group2ndx, 850  
 groups (*lammps.PyLammps property*), 579
- ## H
- has\_exceptions (*lammps.lammps property*), 566  
 has\_ffmpeg\_support (*lammps.lammps property*), 567  
 has\_gpu\_device (*lammps.lammps property*), 567  
 has\_gzip\_support (*lammps.lammps property*), 566  
 has\_id() (*lammps.lammps method*), 568  
 has\_jpeg\_support (*lammps.lammps property*), 567  
 has\_mpi\_support (*lammps.lammps property*), 566  
 has\_png\_support (*lammps.lammps property*), 566  
 has\_style() (*lammps.lammps method*), 568  
 hyper, 851
- ## I
- id (*lammps.Atom property*), 580  
 if, 854  
 image() (*lammps.IPyLammps method*), 582  
 improper\_coeff, 857  
 improper\_style, 859  
 improper\_style amoeba, 2483  
 improper\_style class2, 2484  
 improper\_style class2/kk, 2484  
 improper\_style class2/omp, 2484  
 improper\_style cossq, 2486  
 improper\_style cossq/omp, 2486  
 improper\_style cvff, 2487  
 improper\_style cvff/intel, 2487  
 improper\_style cvff/omp, 2487  
 improper\_style distance, 2489  
 improper\_style distharm, 2490  
 improper\_style fourier, 2491  
 improper\_style fourier/omp, 2491  
 improper\_style harmonic, 2493  
 improper\_style harmonic/intel, 2493  
 improper\_style harmonic/kk, 2493  
 improper\_style harmonic/omp, 2493  
 improper\_style hybrid, 2494  
 improper\_style inversion/harmonic, 2496  
 improper\_style none, 2497  
 improper\_style ring, 2498  
 improper\_style ring/omp, 2498  
 improper\_style sqdistharm, 2499  
 improper\_style umbrella, 2500  
 improper\_style umbrella/omp, 2500  
 improper\_style zero, 2502  
 include, 860  
 info, 861  
 installed\_packages (*lammps.lammps property*), 568  
 IPyLammps (*class in lammps*), 582  
 is\_running (*lammps.lammps property*), 566

## J

jump, 863

## K

kim\_commands, 865  
 kspace\_modify, 884  
 kspace\_style ewald, 890  
 kspace\_style ewald/dipole, 890  
 kspace\_style ewald/dipole/spin, 890  
 kspace\_style ewald/disp, 890  
 kspace\_style ewald/disp/dipole, 890  
 kspace\_style ewald/electrode, 890  
 kspace\_style ewald/omp, 890  
 kspace\_style msm, 890  
 kspace\_style msm/cg, 890  
 kspace\_style msm/cg/omp, 890  
 kspace\_style msm/dielectric, 890  
 kspace\_style msm/omp, 890  
 kspace\_style pppm, 890  
 kspace\_style pppm/cg, 890  
 kspace\_style pppm/cg/omp, 890  
 kspace\_style pppm/dielectric, 890  
 kspace\_style pppm/dipole, 890  
 kspace\_style pppm/dipole/spin, 890  
 kspace\_style pppm/disp, 890  
 kspace\_style pppm/disp/dielectric, 890  
 kspace\_style pppm/disp/intel, 890  
 kspace\_style pppm/disp/omp, 890  
 kspace\_style pppm/disp/tip4p, 890  
 kspace\_style pppm/disp/tip4p/omp, 890  
 kspace\_style pppm/electrode, 890  
 kspace\_style pppm/electrode/intel, 890  
 kspace\_style pppm/gpu, 890  
 kspace\_style pppm/intel, 890  
 kspace\_style pppm/kk, 890  
 kspace\_style pppm/omp, 890  
 kspace\_style pppm/stagger, 890  
 kspace\_style pppm/tip4p, 890  
 kspace\_style pppm/tip4p/omp, 890  
 kspace\_style scafacos, 890

## L

label, 898  
 labelmap, 899  
 lammps  
   module, 558  
 lammps (class in lammps), 558  
 lammps (fortran type), 532  
 lammps() (fortran function), 532  
 lammps.formats  
   module, 586  
 lammps\_close (C++ function), 489  
 lammps\_command (C++ function), 492

lammps\_commands\_list (C++ function), 492  
 lammps\_commands\_string (C++ function), 492  
 lammps\_config\_accelerator (C++ function), 518  
 lammps\_config\_has\_exceptions (C++ function), 517  
 lammps\_config\_has\_ffmpeg\_support (C++ function), 517  
 lammps\_config\_has\_gzip\_support (C++ function), 516  
 lammps\_config\_has\_jpeg\_support (C++ function), 517  
 lammps\_config\_has\_mpi\_support (C++ function), 516  
 lammps\_config\_has\_package (C++ function), 517  
 lammps\_config\_has\_png\_support (C++ function), 516  
 lammps\_config\_package\_count (C++ function), 518  
 lammps\_config\_package\_name (C++ function), 518  
 lammps\_create\_atoms (C++ function), 511  
 lammps\_decode\_image\_flags (C++ function), 522  
 lammps\_encode\_image\_flags (C++ function), 522  
 lammps\_extract\_atom (C++ function), 503  
 lammps\_extract\_atom\_datatype (C++ function), 503  
 lammps\_extract\_box (C++ function), 495  
 lammps\_extract\_compute (C++ function), 504  
 lammps\_extract\_fix (C++ function), 505  
 lammps\_extract\_global (C++ function), 498  
 lammps\_extract\_global\_datatype (C++ function), 498  
 lammps\_extract\_setting (C++ function), 496  
 lammps\_extract\_variable (C++ function), 506  
 lammps\_file (C++ function), 491  
 lammps\_find\_compute\_neighlist (C++ function), 512  
 lammps\_find\_fix\_neighlist (C++ function), 513  
 lammps\_find\_pair\_neighlist (C++ function), 513  
 lammps\_fix\_external\_set\_energy\_global (C++ function), 523  
 lammps\_fix\_external\_set\_energy\_peratom (C++ function), 524  
 lammps\_fix\_external\_set\_vector (C++ function), 526  
 lammps\_fix\_external\_set\_vector\_length (C++ function), 525  
 lammps\_fix\_external\_set\_virial\_global (C++ function), 524  
 lammps\_fix\_external\_set\_virial\_peratom (C++ function), 525  
 lammps\_flush\_buffers (C++ function), 526  
 lammps\_force\_timeout (C++ function), 527  
 lammps\_free (C++ function), 526  
 lammps\_gather (C++ function), 511  
 lammps\_gather\_atoms (C++ function), 509  
 lammps\_gather\_atoms\_concat (C++ function), 509  
 lammps\_gather\_atoms\_subset (C++ function), 509



- lammps\_gather\_bonds (C++ function), 510  
 lammps\_gather\_concat (C++ function), 511  
 lammps\_gather\_subset (C++ function), 511  
 lammps\_get\_gpu\_device\_info (C++ function), 519  
 lammps\_get\_last\_error\_message (C++ function), 527  
 lammps\_get\_mpi\_comm (C++ function), 496  
 lammps\_get\_natoms (C++ function), 494  
 lammps\_get\_os\_info (C++ function), 516  
 lammps\_get\_thermo (C++ function), 494  
 lammps\_has\_error (C++ function), 527  
 lammps\_has\_gpu\_device (C++ function), 519  
 lammps\_has\_id (C++ function), 520  
 lammps\_has\_style (C++ function), 519  
 lammps\_id\_count (C++ function), 520  
 lammps\_id\_name (C++ function), 521  
 lammps\_is\_running (C++ function), 527  
 lammps\_kokkos\_finalize (C++ function), 490  
 lammps\_memory\_usage (C++ function), 496  
 lammps\_mpi\_finalize (C++ function), 489  
 lammps\_mpi\_init (C++ function), 489  
 lammps\_neighlist\_element\_neighbors (C++ function), 514  
 lammps\_neighlist\_num\_elements (C++ function), 513  
 LAMMPS\_NS::ArgInfo (C++ class), 699  
 LAMMPS\_NS::ArgInfo::ArgInfo (C++ function), 701  
 LAMMPS\_NS::ArgInfo::ArgTypes (C++ enum), 699  
 LAMMPS\_NS::ArgInfo::ArgTypes::BIN1D (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::BIN2D (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::BIN3D (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::BINCYLINDER (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::BINSPHERE (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::COMPUTE (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::DENSITY\_MASS (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::DENSITY\_NUMBER (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::DNAME (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::ERROR (C++ enumerator), 699  
 LAMMPS\_NS::ArgInfo::ArgTypes::F (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::FIX (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::INAME (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::KEYWORD (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::MASS (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::MOLECULE (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::NONE (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::TEMPERATURE (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::TYPE (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::UNKNOWN (C++ enumerator), 699  
 LAMMPS\_NS::ArgInfo::ArgTypes::V (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::VARIABLE (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::ArgTypes::X (C++ enumerator), 700  
 LAMMPS\_NS::ArgInfo::copy\_name (C++ function), 701  
 LAMMPS\_NS::ArgInfo::get\_dim (C++ function), 701  
 LAMMPS\_NS::ArgInfo::get\_index1 (C++ function), 701  
 LAMMPS\_NS::ArgInfo::get\_index2 (C++ function), 701  
 LAMMPS\_NS::ArgInfo::get\_name (C++ function), 701  
 LAMMPS\_NS::ArgInfo::get\_type (C++ function), 701  
 LAMMPS\_NS::Atom (C++ class), 665  
 LAMMPS\_NS::Atom::add\_custom (C++ function), 666  
 LAMMPS\_NS::Atom::Atom (C++ function), 666  
 LAMMPS\_NS::Atom::extract (C++ function), 666  
 LAMMPS\_NS::Atom::extract\_datatype (C++ function), 668  
 LAMMPS\_NS::Atom::PerAtom (C++ struct), 668  
 LAMMPS\_NS::Atom::remove\_custom (C++ function), 666  
 LAMMPS\_NS::Input (C++ class), 668  
 LAMMPS\_NS::Input::file (C++ function), 669  
 LAMMPS\_NS::Input::Input (C++ function), 668  
 LAMMPS\_NS::Input::one (C++ function), 669  
 LAMMPS\_NS::InvalidFloatException (C++ class), 698  
 LAMMPS\_NS::InvalidFloatException::InvalidFloatException (C++ function), 699  
 LAMMPS\_NS::InvalidIntegerException (C++ class), 698  
 LAMMPS\_NS::InvalidIntegerException::InvalidIntegerException (C++ function), 698  
 LAMMPS\_NS::LAMMPS (C++ class), 664  
 LAMMPS\_NS::LAMMPS::~LAMMPS (C++ function), 664  
 LAMMPS\_NS::LAMMPS::is\_installed\_pkg (C++ function), 665

- LAMMPS\_NS::LAMMPS::LAMMPS (C++ function), 664
- LAMMPS\_NS::LAMMPS::match\_style (C++ function), 664
- LAMMPS\_NS::MathSpecial::cube (C++ function), 694
- LAMMPS\_NS::MathSpecial::exp2\_x86 (C++ function), 692
- LAMMPS\_NS::MathSpecial::expmsq (C++ function), 693
- LAMMPS\_NS::MathSpecial::factorial (C++ function), 692
- LAMMPS\_NS::MathSpecial::fm\_exp (C++ function), 693
- LAMMPS\_NS::MathSpecial::my\_erfcx (C++ function), 693
- LAMMPS\_NS::MathSpecial::powint (C++ function), 694
- LAMMPS\_NS::MathSpecial::powsign (C++ function), 694
- LAMMPS\_NS::MathSpecial::powsinxx (C++ function), 694
- LAMMPS\_NS::MathSpecial::square (C++ function), 693
- LAMMPS\_NS::MyPage (C++ class), 707
- LAMMPS\_NS::MyPage::get (C++ function), 708
- LAMMPS\_NS::MyPage::init (C++ function), 708
- LAMMPS\_NS::MyPage::MyPage (C++ function), 708
- LAMMPS\_NS::MyPage::reset (C++ function), 708
- LAMMPS\_NS::MyPage::size (C++ function), 708
- LAMMPS\_NS::MyPage::status (C++ function), 709
- LAMMPS\_NS::MyPage::vget (C++ function), 708
- LAMMPS\_NS::MyPage::vgot (C++ function), 708
- LAMMPS\_NS::MyPoolChunk (C++ class), 709
- LAMMPS\_NS::MyPoolChunk::~MyPoolChunk (C++ function), 709
- LAMMPS\_NS::MyPoolChunk::get (C++ function), 709
- LAMMPS\_NS::MyPoolChunk::MyPoolChunk (C++ function), 709
- LAMMPS\_NS::MyPoolChunk::put (C++ function), 710
- LAMMPS\_NS::MyPoolChunk::size (C++ function), 710
- LAMMPS\_NS::MyPoolChunk::status (C++ function), 710
- LAMMPS\_NS::platform::chdir (C++ function), 673
- LAMMPS\_NS::platform::compiler\_info (C++ function), 670
- LAMMPS\_NS::platform::compress\_info (C++ function), 671
- LAMMPS\_NS::platform::compressed\_read (C++ function), 676
- LAMMPS\_NS::platform::compressed\_write (C++ function), 677
- LAMMPS\_NS::platform::cputime (C++ function), 669
- LAMMPS\_NS::platform::current\_directory (C++ function), 672
- LAMMPS\_NS::platform::cxx\_standard (C++ function), 670
- LAMMPS\_NS::platform::dlclose (C++ function), 676
- LAMMPS\_NS::platform::dlerror (C++ function), 676
- LAMMPS\_NS::platform::dlopen (C++ function), 676
- LAMMPS\_NS::platform::dlsym (C++ function), 676
- LAMMPS\_NS::platform::END\_OF\_FILE (C++ member), 674
- LAMMPS\_NS::platform::file\_is\_readable (C++ function), 672
- LAMMPS\_NS::platform::filepathsep (C++ member), 671
- LAMMPS\_NS::platform::find\_exe\_path (C++ function), 675
- LAMMPS\_NS::platform::fseek (C++ function), 674
- LAMMPS\_NS::platform::ftell (C++ function), 674
- LAMMPS\_NS::platform::ftruncate (C++ function), 674
- LAMMPS\_NS::platform::guesspath (C++ function), 671
- LAMMPS\_NS::platform::has\_compress\_extension (C++ function), 676
- LAMMPS\_NS::platform::is\_console (C++ function), 672
- LAMMPS\_NS::platform::list\_directory (C++ function), 672
- LAMMPS\_NS::platform::list\_pathenv (C++ function), 675
- LAMMPS\_NS::platform::mkdir (C++ function), 673
- LAMMPS\_NS::platform::mpi\_info (C++ function), 670
- LAMMPS\_NS::platform::mpi\_vendor (C++ function), 670
- LAMMPS\_NS::platform::openmp\_standard (C++ function), 670
- LAMMPS\_NS::platform::os\_info (C++ function), 670
- LAMMPS\_NS::platform::path\_basename (C++ function), 672
- LAMMPS\_NS::platform::path\_is\_directory (C++ function), 672
- LAMMPS\_NS::platform::path\_join (C++ function), 672
- LAMMPS\_NS::platform::pathvarsep (C++ member), 671
- LAMMPS\_NS::platform::pclose (C++ function), 674
- LAMMPS\_NS::platform::popen (C++ function), 674
- LAMMPS\_NS::platform::putenv (C++ function), 675
- LAMMPS\_NS::platform::rmdir (C++ function), 673
- LAMMPS\_NS::platform::unlink (C++ function), 673
- LAMMPS\_NS::platform::unsetenv (C++ function), 675
- LAMMPS\_NS::platform::usleep (C++ function), 670
- LAMMPS\_NS::platform::walltime (C++ function), 669
- LAMMPS\_NS::Pointers (C++ class), 665

LAMMPS\_NS::PotentialFileReader (C++ class), 704  
 LAMMPS\_NS::PotentialFileReader::~~PotentialFileReader (C++ function), 705  
 LAMMPS\_NS::PotentialFileReader::ignore\_comments (C++ function), 705  
 LAMMPS\_NS::PotentialFileReader::next\_bigint (C++ function), 706  
 LAMMPS\_NS::PotentialFileReader::next\_double (C++ function), 706  
 LAMMPS\_NS::PotentialFileReader::next\_dvector (C++ function), 705  
 LAMMPS\_NS::PotentialFileReader::next\_int (C++ function), 706  
 LAMMPS\_NS::PotentialFileReader::next\_line (C++ function), 705  
 LAMMPS\_NS::PotentialFileReader::next\_string (C++ function), 706  
 LAMMPS\_NS::PotentialFileReader::next\_tagint (C++ function), 706  
 LAMMPS\_NS::PotentialFileReader::next\_values (C++ function), 706  
 LAMMPS\_NS::PotentialFileReader::PotentialFileReader (C++ function), 705  
 LAMMPS\_NS::PotentialFileReader::rewind (C++ function), 705  
 LAMMPS\_NS::PotentialFileReader::skip\_line (C++ function), 705  
 LAMMPS\_NS::TextFileReader (C++ class), 703  
 LAMMPS\_NS::TextFileReader::~~TextFileReader (C++ function), 703  
 LAMMPS\_NS::TextFileReader::ignore\_comments (C++ member), 704  
 LAMMPS\_NS::TextFileReader::next\_dvector (C++ function), 704  
 LAMMPS\_NS::TextFileReader::next\_line (C++ function), 703  
 LAMMPS\_NS::TextFileReader::next\_values (C++ function), 704  
 LAMMPS\_NS::TextFileReader::rewind (C++ function), 703  
 LAMMPS\_NS::TextFileReader::set\_bufsize (C++ function), 703  
 LAMMPS\_NS::TextFileReader::skip\_line (C++ function), 703  
 LAMMPS\_NS::TextFileReader::TextFileReader (C++ function), 703  
 LAMMPS\_NS::Tokenizer (C++ class), 695  
 LAMMPS\_NS::Tokenizer::as\_vector (C++ function), 696  
 LAMMPS\_NS::Tokenizer::contains (C++ function), 696  
 LAMMPS\_NS::Tokenizer::count (C++ function), 696  
 LAMMPS\_NS::Tokenizer::has\_next (C++ function), 696  
 LAMMPS\_NS::Tokenizer::next (C++ function), 696  
 LAMMPS\_NS::Tokenizer::reset (C++ function), 696  
 LAMMPS\_NS::Tokenizer::skip (C++ function), 696  
 LAMMPS\_NS::Tokenizer::Tokenizer (C++ function), 695  
 LAMMPS\_NS::TokenizerException (C++ class), 696  
 LAMMPS\_NS::TokenizerException::TokenizerException (C++ function), 697  
 LAMMPS\_NS::TokenizerException::what (C++ function), 697  
 LAMMPS\_NS::ubuf (C++ union), 711  
 LAMMPS\_NS::ubuf::d (C++ member), 712  
 LAMMPS\_NS::ubuf::i (C++ member), 712  
 LAMMPS\_NS::ubuf::ubuf (C++ function), 712  
 LAMMPS\_NS::utils::binary\_search (C++ function), 692  
 LAMMPS\_NS::utils::bnumeric (C++ function), 680  
 LAMMPS\_NS::utils::bounds (C++ function), 688  
 LAMMPS\_NS::utils::check\_packages\_for\_style (C++ function), 691  
 LAMMPS\_NS::utils::count\_words (C++ function), 684  
 LAMMPS\_NS::utils::current\_date (C++ function), 691  
 LAMMPS\_NS::utils::date2num (C++ function), 691  
 LAMMPS\_NS::utils::errorurl (C++ function), 690  
 LAMMPS\_NS::utils::expand\_args (C++ function), 688  
 LAMMPS\_NS::utils::expand\_type (C++ function), 689  
 LAMMPS\_NS::utils::fgets\_trunc (C++ function), 678  
 LAMMPS\_NS::utils::flush\_buffers (C++ function), 690  
 LAMMPS\_NS::utils::get\_conversion\_factor (C++ function), 687  
 LAMMPS\_NS::utils::get\_potential\_date (C++ function), 687  
 LAMMPS\_NS::utils::get\_potential\_file\_path (C++ function), 687  
 LAMMPS\_NS::utils::get\_potential\_units (C++ function), 687  
 LAMMPS\_NS::utils::get\_supported\_conversions (C++ function), 687  
 LAMMPS\_NS::utils::getsyserror (C++ function), 690  
 LAMMPS\_NS::utils::has\_utf8 (C++ function), 683  
 LAMMPS\_NS::utils::inumeric (C++ function), 679, 680  
 LAMMPS\_NS::utils::is\_double (C++ function), 686  
 LAMMPS\_NS::utils::is\_id (C++ function), 686  
 LAMMPS\_NS::utils::is\_integer (C++ function), 686  
 LAMMPS\_NS::utils::is\_type (C++ function), 686  
 LAMMPS\_NS::utils::join\_words (C++ function), 684

- LAMMPS\_NS::utils::logical (C++ function), 681  
 LAMMPS\_NS::utils::logmesg (C++ function), 690  
 LAMMPS\_NS::utils::lowercase (C++ function), 682  
 LAMMPS\_NS::utils::merge\_sort (C++ function), 692  
 LAMMPS\_NS::utils::missing\_cmd\_args (C++ function), 690  
 LAMMPS\_NS::utils::numeric (C++ function), 679  
 LAMMPS\_NS::utils::open\_potential (C++ function), 687  
 LAMMPS\_NS::utils::read\_lines\_from\_file (C++ function), 678  
 LAMMPS\_NS::utils::sfgets (C++ function), 677  
 LAMMPS\_NS::utils::sfread (C++ function), 678  
 LAMMPS\_NS::utils::split\_lines (C++ function), 685  
 LAMMPS\_NS::utils::split\_words (C++ function), 685  
 LAMMPS\_NS::utils::star\_subst (C++ function), 683  
 LAMMPS\_NS::utils::strdup (C++ function), 682  
 LAMMPS\_NS::utils::strfind (C++ function), 685  
 LAMMPS\_NS::utils::strmatch (C++ function), 685  
 LAMMPS\_NS::utils::timespec2seconds (C++ function), 691  
 LAMMPS\_NS::utils::tnumeric (C++ function), 681  
 LAMMPS\_NS::utils::trim (C++ function), 682  
 LAMMPS\_NS::utils::trim\_and\_count\_words (C++ function), 684  
 LAMMPS\_NS::utils::trim\_comment (C++ function), 683  
 LAMMPS\_NS::utils::uppercase (C++ function), 682  
 LAMMPS\_NS::utils::utf8\_subst (C++ function), 683  
 LAMMPS\_NS::ValueTokenizer (C++ class), 697  
 LAMMPS\_NS::ValueTokenizer::contains (C++ function), 698  
 LAMMPS\_NS::ValueTokenizer::count (C++ function), 698  
 LAMMPS\_NS::ValueTokenizer::has\_next (C++ function), 698  
 LAMMPS\_NS::ValueTokenizer::next\_bigint (C++ function), 697  
 LAMMPS\_NS::ValueTokenizer::next\_double (C++ function), 698  
 LAMMPS\_NS::ValueTokenizer::next\_int (C++ function), 697  
 LAMMPS\_NS::ValueTokenizer::next\_string (C++ function), 697  
 LAMMPS\_NS::ValueTokenizer::next\_tagint (C++ function), 697  
 LAMMPS\_NS::ValueTokenizer::skip (C++ function), 698  
 LAMMPS\_NS::ValueTokenizer::ValueTokenizer (C++ function), 697  
 lammps\_open (C++ function), 487  
 lammps\_open\_fortran (C++ function), 488  
 lammps\_open\_no\_mpi (C++ function), 488  
 lammps\_python\_finalize (C++ function), 490  
 lammps\_reset\_box (C++ function), 495  
 lammps\_scatter (C++ function), 511  
 lammps\_scatter\_atoms (C++ function), 509  
 lammps\_scatter\_atoms\_subset (C++ function), 509  
 lammps\_scatter\_subset (C++ function), 511  
 lammps\_set\_fix\_external\_callback (C++ function), 523  
 lammps\_set\_variable (C++ function), 507  
 lammps\_style\_count (C++ function), 519  
 lammps\_style\_name (C++ function), 520  
 lammps\_version (C++ function), 515  
 last\_run (lammps.PyLammps property), 578  
 lattice, 900  
 lmp\_print() (lammps.PyLammps method), 580  
 log, 904
- ## M
- mass, 905  
 mass (lammps.Atom property), 580  
 MathEigen::jacobi3 (C++ function), 710, 711  
 mdi, 906  
 min\_modify, 911  
 min\_style, 916  
 min\_style spin, 914  
 minimize, 918  
 module  
   lammps, 558  
   lammps.formats, 586  
 mol (lammps.Atom property), 580  
 molecule, 923
- ## N
- ndx2group, 850  
 neb, 931  
 neb/spin, 936  
 neigh\_modify, 941  
 neighbor, 945  
 NeighList (class in lammps), 584  
 newton, 946  
 next, 948  
 numpy (lammps.lammps property), 559  
 numpy\_wrapper (class in lammps.numpy\_wrapper), 573  
 NumPyNeighList (class in lammps.numpy\_wrapper), 585
- ## O
- omega (lammps.Atom property), 581
- ## P
- package, 950  
 pair\_coeff, 959  
 pair\_modify, 961

pair\_style, 966  
 pair\_style adp, 1975  
 pair\_style adp/kk, 1975  
 pair\_style adp/omp, 1975  
 pair\_style agni, 1977  
 pair\_style agni/omp, 1977  
 pair\_style airebo, 1979  
 pair\_style airebo/intel, 1979  
 pair\_style airebo/morse, 1979  
 pair\_style airebo/morse/intel, 1979  
 pair\_style airebo/morse/omp, 1979  
 pair\_style airebo/omp, 1979  
 pair\_style amoeba, 1983  
 pair\_style atm, 1986  
 pair\_style awpmd/cut, 1989  
 pair\_style beck, 1991  
 pair\_style beck/gpu, 1991  
 pair\_style beck/omp, 1991  
 pair\_style body/nparticle, 1993  
 pair\_style body/rounded/polygon, 1995  
 pair\_style body/rounded/polyhedron, 1997  
 pair\_style bop, 2000  
 pair\_style born, 2006  
 pair\_style born/coul/dsf, 2006  
 pair\_style born/coul/dsf/cs, 2049  
 pair\_style born/coul/long, 2006  
 pair\_style born/coul/long/cs, 2049  
 pair\_style born/coul/long/cs/gpu, 2049  
 pair\_style born/coul/long/gpu, 2006  
 pair\_style born/coul/long/omp, 2006  
 pair\_style born/coul/msm, 2006  
 pair\_style born/coul/msm/omp, 2006  
 pair\_style born/coul/wolf, 2006  
 pair\_style born/coul/wolf/cs, 2049  
 pair\_style born/coul/wolf/cs/gpu, 2049  
 pair\_style born/coul/wolf/gpu, 2006  
 pair\_style born/coul/wolf/omp, 2006  
 pair\_style born/gpu, 2006  
 pair\_style born/omp, 2006  
 pair\_style bpm/spring, 2009  
 pair\_style brownian, 2011  
 pair\_style brownian/omp, 2011  
 pair\_style brownian/poly, 2011  
 pair\_style brownian/poly/omp, 2011  
 pair\_style buck, 2013  
 pair\_style buck/coul/cut, 2013  
 pair\_style buck/coul/cut/gpu, 2013  
 pair\_style buck/coul/cut/intel, 2013  
 pair\_style buck/coul/cut/kk, 2013  
 pair\_style buck/coul/cut/omp, 2013  
 pair\_style buck/coul/long, 2013  
 pair\_style buck/coul/long/cs, 2049  
 pair\_style buck/coul/long/gpu, 2013  
 pair\_style buck/coul/long/intel, 2013  
 pair\_style buck/coul/long/kk, 2013  
 pair\_style buck/coul/long/omp, 2013  
 pair\_style buck/coul/msm, 2013  
 pair\_style buck/coul/msm/omp, 2013  
 pair\_style buck/gpu, 2013  
 pair\_style buck/intel, 2013  
 pair\_style buck/kk, 2013  
 pair\_style buck/long/coul/long, 2018  
 pair\_style buck/long/coul/long/omp, 2018  
 pair\_style buck/mdf, 2201  
 pair\_style buck/omp, 2013  
 pair\_style buck6d/coul/gauss/dsf, 2016  
 pair\_style buck6d/coul/gauss/long, 2016  
 pair\_style colloid, 2028  
 pair\_style colloid/gpu, 2028  
 pair\_style colloid/omp, 2028  
 pair\_style comb, 2031  
 pair\_style comb/omp, 2031  
 pair\_style comb3, 2031  
 pair\_style cosine/squared, 2034  
 pair\_style coul/cut, 2036  
 pair\_style coul/cut/dielectric, 2052  
 pair\_style coul/cut/global, 2036  
 pair\_style coul/cut/global/omp, 2036  
 pair\_style coul/cut/gpu, 2036  
 pair\_style coul/cut/kk, 2036  
 pair\_style coul/cut/omp, 2036  
 pair\_style coul/cut/soft, 2097  
 pair\_style coul/cut/soft/omp, 2097  
 pair\_style coul/debye, 2036  
 pair\_style coul/debye/gpu, 2036  
 pair\_style coul/debye/kk, 2036  
 pair\_style coul/debye/omp, 2036  
 pair\_style coul/diel, 2042  
 pair\_style coul/diel/omp, 2042  
 pair\_style coul/dsf, 2036  
 pair\_style coul/dsf/gpu, 2036  
 pair\_style coul/dsf/kk, 2036  
 pair\_style coul/dsf/omp, 2036  
 pair\_style coul/exclude, 2036  
 pair\_style coul/long, 2036  
 pair\_style coul/long/cs, 2049  
 pair\_style coul/long/cs/gpu, 2049  
 pair\_style coul/long/dielectric, 2052  
 pair\_style coul/long/gpu, 2036  
 pair\_style coul/long/kk, 2036  
 pair\_style coul/long/omp, 2036  
 pair\_style coul/long/soft, 2097  
 pair\_style coul/long/soft/omp, 2097  
 pair\_style coul/msm, 2036  
 pair\_style coul/msm/omp, 2036  
 pair\_style coul/shield, 2044  
 pair\_style coul/slatter, 2045  
 pair\_style coul/slatter/cut, 2045

pair\_style coul/slater/long, 2045  
pair\_style coul/streitz, 2036  
pair\_style coul/tt, 2047  
pair\_style coul/wolf, 2036  
pair\_style coul/wolf/cs, 2049  
pair\_style coul/wolf/kk, 2036  
pair\_style coul/wolf/omp, 2036  
pair\_style dpd, 2060  
pair\_style dpd/ext, 2063  
pair\_style dpd/ext/kk, 2063  
pair\_style dpd/ext/tstat, 2063  
pair\_style dpd/ext/tstat/kk, 2063  
pair\_style dpd/fdt, 2066  
pair\_style dpd/fdt/energy, 2066  
pair\_style dpd/fdt/energy/kk, 2066  
pair\_style dpd/gpu, 2060  
pair\_style dpd/intel, 2060  
pair\_style dpd/kk, 2060  
pair\_style dpd/omp, 2060  
pair\_style dpd/tstat, 2060  
pair\_style dpd/tstat/gpu, 2060  
pair\_style dpd/tstat/kk, 2060  
pair\_style dpd/tstat/omp, 2060  
pair\_style drip, 2069  
pair\_style dsmc, 2071  
pair\_style e3b, 2073  
pair\_style eam, 2076  
pair\_style eam/alloy, 2076  
pair\_style eam/alloy/gpu, 2076  
pair\_style eam/alloy/intel, 2076  
pair\_style eam/alloy/kk, 2076  
pair\_style eam/alloy/omp, 2076  
pair\_style eam/alloy/opt, 2076  
pair\_style eam/cd, 2076  
pair\_style eam/cd/old, 2076  
pair\_style eam/fs, 2076  
pair\_style eam/fs/gpu, 2076  
pair\_style eam/fs/intel, 2076  
pair\_style eam/fs/kk, 2076  
pair\_style eam/fs/omp, 2076  
pair\_style eam/fs/opt, 2076  
pair\_style eam/gpu, 2076  
pair\_style eam/he, 2076  
pair\_style eam/intel, 2076  
pair\_style eam/kk, 2076  
pair\_style eam/omp, 2076  
pair\_style eam/opt, 2076  
pair\_style edip, 2083  
pair\_style edip/multi, 2083  
pair\_style edip/omp, 2083  
pair\_style edpd, 2218  
pair\_style eff/cut, 2086  
pair\_style eim, 2090  
pair\_style eim/omp, 2090  
pair\_style exp6/rx, 2093  
pair\_style exp6/rx/kk, 2093  
pair\_style extep, 2096  
pair\_style gauss, 2104  
pair\_style gauss/cut, 2104  
pair\_style gauss/cut/omp, 2104  
pair\_style gauss/gpu, 2104  
pair\_style gauss/omp, 2104  
pair\_style gayberne, 2106  
pair\_style gayberne/gpu, 2106  
pair\_style gayberne/intel, 2106  
pair\_style gayberne/omp, 2106  
pair\_style gran/hertz/history, 2109  
pair\_style gran/hertz/history/omp, 2109  
pair\_style gran/hooke, 2109  
pair\_style gran/hooke/history, 2109  
pair\_style gran/hooke/history/kk, 2109  
pair\_style gran/hooke/history/omp, 2109  
pair\_style gran/hooke/omp, 2109  
pair\_style granular, 2114  
pair\_style gw, 2126  
pair\_style gw/zbl, 2126  
pair\_style harmonic/cut, 2128  
pair\_style harmonic/cut/omp, 2128  
pair\_style hbond/dreiding/lj, 2130  
pair\_style hbond/dreiding/lj/omp, 2130  
pair\_style hbond/dreiding/morse, 2130  
pair\_style hbond/dreiding/morse/omp, 2130  
pair\_style hdnp, 2134  
pair\_style hippo, 1983  
pair\_style hybrid, 2137  
pair\_style hybrid/kk, 2137  
pair\_style hybrid/overlay, 2137  
pair\_style hybrid/overlay/kk, 2137  
pair\_style hybrid/scaled, 2137  
pair\_style ilp/graphene/hbn, 2143  
pair\_style ilp/graphene/hbn/opt, 2143  
pair\_style ilp/tmd, 2146  
pair\_style ilp/tmd/opt, 2146  
pair\_style kim, 2149  
pair\_style kolmogorov/crespi/full, 2150  
pair\_style kolmogorov/crespi/z, 2153  
pair\_style lcbop, 2154  
pair\_style lebedeva/z, 2156  
pair\_style lennard/mdf, 2201  
pair\_style line/lj, 2157  
pair\_style list, 2159  
pair\_style lj/charmm/coul/charmm, 2021  
pair\_style lj/charmm/coul/charmm/gpu, 2021  
pair\_style lj/charmm/coul/charmm/implicit,  
2021  
pair\_style lj/charmm/coul/charmm/implicit/kk,  
2021

pair\_style lj/charmm/coul/charmm/implicit/omp, 2021  
 pair\_style lj/charmm/coul/charmm/intel, 2021  
 pair\_style lj/charmm/coul/charmm/kk, 2021  
 pair\_style lj/charmm/coul/charmm/omp, 2021  
 pair\_style lj/charmm/coul/long, 2021  
 pair\_style lj/charmm/coul/long/gpu, 2021  
 pair\_style lj/charmm/coul/long/intel, 2021  
 pair\_style lj/charmm/coul/long/kk, 2021  
 pair\_style lj/charmm/coul/long/omp, 2021  
 pair\_style lj/charmm/coul/long/opt, 2021  
 pair\_style lj/charmm/coul/long/soft, 2097  
 pair\_style lj/charmm/coul/long/soft/omp, 2097  
 pair\_style lj/charmm/coul/msm, 2021  
 pair\_style lj/charmm/coul/msm/omp, 2021  
 pair\_style lj/charmmfsw/coul/charmmfsh, 2021  
 pair\_style lj/charmmfsw/coul/long, 2021  
 pair\_style lj/class2, 2025  
 pair\_style lj/class2/coul/cut, 2025  
 pair\_style lj/class2/coul/cut/kk, 2025  
 pair\_style lj/class2/coul/cut/omp, 2025  
 pair\_style lj/class2/coul/cut/soft, 2097  
 pair\_style lj/class2/coul/long, 2025  
 pair\_style lj/class2/coul/long/cs, 2049  
 pair\_style lj/class2/coul/long/gpu, 2025  
 pair\_style lj/class2/coul/long/kk, 2025  
 pair\_style lj/class2/coul/long/omp, 2025  
 pair\_style lj/class2/coul/long/soft, 2097  
 pair\_style lj/class2/gpu, 2025  
 pair\_style lj/class2/kk, 2025  
 pair\_style lj/class2/omp, 2025  
 pair\_style lj/class2/soft, 2097  
 pair\_style lj/cubic, 2166  
 pair\_style lj/cubic/gpu, 2166  
 pair\_style lj/cubic/omp, 2166  
 pair\_style lj/cut, 2162  
 pair\_style lj/cut/coul/cut, 2168  
 pair\_style lj/cut/coul/cut/dielectric, 2052  
 pair\_style lj/cut/coul/cut/dielectric/omp, 2052  
 pair\_style lj/cut/coul/cut/gpu, 2168  
 pair\_style lj/cut/coul/cut/kk, 2168  
 pair\_style lj/cut/coul/cut/omp, 2168  
 pair\_style lj/cut/coul/cut/soft, 2097  
 pair\_style lj/cut/coul/cut/soft/omp, 2097  
 pair\_style lj/cut/coul/debye, 2168  
 pair\_style lj/cut/coul/debye/dielectric, 2052  
 pair\_style lj/cut/coul/debye/dielectric/omp, 2052  
 pair\_style lj/cut/coul/debye/gpu, 2168  
 pair\_style lj/cut/coul/debye/kk, 2168  
 pair\_style lj/cut/coul/debye/omp, 2168  
 pair\_style lj/cut/coul/dsf, 2168  
 pair\_style lj/cut/coul/dsf/gpu, 2168  
 pair\_style lj/cut/coul/dsf/kk, 2168  
 pair\_style lj/cut/coul/dsf/omp, 2168  
 pair\_style lj/cut/coul/long, 2168  
 pair\_style lj/cut/coul/long/cs, 2049  
 pair\_style lj/cut/coul/long/dielectric, 2052  
 pair\_style lj/cut/coul/long/dielectric/omp, 2052  
 pair\_style lj/cut/coul/long/gpu, 2168  
 pair\_style lj/cut/coul/long/intel, 2168  
 pair\_style lj/cut/coul/long/kk, 2168  
 pair\_style lj/cut/coul/long/omp, 2168  
 pair\_style lj/cut/coul/long/opt, 2168  
 pair\_style lj/cut/coul/long/soft, 2097  
 pair\_style lj/cut/coul/long/soft/omp, 2097  
 pair\_style lj/cut/coul/msm, 2168  
 pair\_style lj/cut/coul/msm/dielectric, 2052  
 pair\_style lj/cut/coul/msm/gpu, 2168  
 pair\_style lj/cut/coul/msm/omp, 2168  
 pair\_style lj/cut/coul/wolf, 2168  
 pair\_style lj/cut/coul/wolf/omp, 2168  
 pair\_style lj/cut/dipole/cut, 2055  
 pair\_style lj/cut/dipole/cut/gpu, 2055  
 pair\_style lj/cut/dipole/cut/omp, 2055  
 pair\_style lj/cut/dipole/long, 2055  
 pair\_style lj/cut/dipole/long/gpu, 2055  
 pair\_style lj/cut/gpu, 2162  
 pair\_style lj/cut/intel, 2162  
 pair\_style lj/cut/kk, 2162  
 pair\_style lj/cut/omp, 2162  
 pair\_style lj/cut/opt, 2162  
 pair\_style lj/cut/soft, 2097  
 pair\_style lj/cut/soft/omp, 2097  
 pair\_style lj/cut/thole/long, 2360  
 pair\_style lj/cut/thole/long/omp, 2360  
 pair\_style lj/cut/tip4p/cut, 2172  
 pair\_style lj/cut/tip4p/cut/omp, 2172  
 pair\_style lj/cut/tip4p/long, 2172  
 pair\_style lj/cut/tip4p/long/gpu, 2172  
 pair\_style lj/cut/tip4p/long/omp, 2172  
 pair\_style lj/cut/tip4p/long/opt, 2172  
 pair\_style lj/cut/tip4p/long/soft, 2097  
 pair\_style lj/cut/tip4p/long/soft/omp, 2097  
 pair\_style lj/expand, 2175  
 pair\_style lj/expand/coul/long, 2175  
 pair\_style lj/expand/coul/long/gpu, 2175  
 pair\_style lj/expand/gpu, 2175  
 pair\_style lj/expand/kk, 2175  
 pair\_style lj/expand/omp, 2175  
 pair\_style lj/gromacs, 2124  
 pair\_style lj/gromacs/coul/gromacs, 2124  
 pair\_style lj/gromacs/coul/gromacs/kk, 2124  
 pair\_style lj/gromacs/coul/gromacs/omp, 2124  
 pair\_style lj/gromacs/gpu, 2124  
 pair\_style lj/gromacs/kk, 2124

pair\_style lj/gromacs/omp, 2124  
pair\_style lj/long/coul/long, 2177  
pair\_style lj/long/coul/long/dielectric, 2052  
pair\_style lj/long/coul/long/intel, 2177  
pair\_style lj/long/coul/long/omp, 2177  
pair\_style lj/long/coul/long/opt, 2177  
pair\_style lj/long/dipole/long, 2055  
pair\_style lj/long/tip4p/long, 2177  
pair\_style lj/long/tip4p/long/omp, 2177  
pair\_style lj/mdf, 2201  
pair\_style lj/relres, 2181  
pair\_style lj/relres/omp, 2181  
pair\_style lj/sf/dipole/sf, 2055  
pair\_style lj/sf/dipole/sf/gpu, 2055  
pair\_style lj/sf/dipole/sf/omp, 2055  
pair\_style lj/smooth, 2185  
pair\_style lj/smooth/gpu, 2185  
pair\_style lj/smooth/linear, 2187  
pair\_style lj/smooth/linear/omp, 2187  
pair\_style lj/smooth/omp, 2185  
pair\_style lj/spica, 2317  
pair\_style lj/spica/coul/long, 2317  
pair\_style lj/spica/coul/long/gpu, 2317  
pair\_style lj/spica/coul/long/omp, 2317  
pair\_style lj/spica/coul/msm, 2317  
pair\_style lj/spica/coul/msm/omp, 2317  
pair\_style lj/spica/gpu, 2317  
pair\_style lj/spica/kk, 2317  
pair\_style lj/spica/omp, 2317  
pair\_style lj/switch3/coulgauss/long, 2189  
pair\_style lj96/cut, 2164  
pair\_style lj96/cut/gpu, 2164  
pair\_style lj96/cut/omp, 2164  
pair\_style local/density, 2191  
pair\_style lubricate, 2194  
pair\_style lubricate/omp, 2194  
pair\_style lubricate/poly, 2194  
pair\_style lubricate/poly/omp, 2194  
pair\_style lubricateU, 2198  
pair\_style lubricateU/poly, 2198  
pair\_style mdpd, 2218  
pair\_style mdpd/rhosum, 2218  
pair\_style meam, 2204  
pair\_style meam/kk, 2204  
pair\_style meam/spline, 2210  
pair\_style meam/spline/omp, 2210  
pair\_style meam/sw/spline, 2212  
pair\_style mesocnt, 2214  
pair\_style mesocnt/viscous, 2214  
pair\_style mesont/tpm, 2224  
pair\_style mgpt, 2227  
pair\_style mie/cut, 2229  
pair\_style mie/cut/gpu, 2229  
pair\_style mliap, 2231  
pair\_style mm3/switch3/coulgauss/long, 2189  
pair\_style momb, 2234  
pair\_style morse, 2235  
pair\_style morse/gpu, 2235  
pair\_style morse/kk, 2235  
pair\_style morse/omp, 2235  
pair\_style morse/opt, 2235  
pair\_style morse/smooth/linear, 2235  
pair\_style morse/smooth/linear/omp, 2235  
pair\_style morse/soft, 2097  
pair\_style multi/lucy, 2238  
pair\_style multi/lucy/rx, 2241  
pair\_style multi/lucy/rx/kk, 2241  
pair\_style nb3b/harmonic, 2244  
pair\_style nm/cut, 2246  
pair\_style nm/cut/coul/cut, 2246  
pair\_style nm/cut/coul/cut/omp, 2246  
pair\_style nm/cut/coul/long, 2246  
pair\_style nm/cut/coul/long/omp, 2246  
pair\_style nm/cut/omp, 2246  
pair\_style nm/cut/split, 2246  
pair\_style none, 2249  
pair\_style oxdna/coaxstk, 2250  
pair\_style oxdna/excv, 2250  
pair\_style oxdna/hbond, 2250  
pair\_style oxdna/stk, 2250  
pair\_style oxdna/xstk, 2250  
pair\_style oxdna2/coaxstk, 2252  
pair\_style oxdna2/dh, 2252  
pair\_style oxdna2/excv, 2252  
pair\_style oxdna2/hbond, 2252  
pair\_style oxdna2/stk, 2252  
pair\_style oxdna2/xstk, 2252  
pair\_style oxrna2/coaxstk, 2255  
pair\_style oxrna2/dh, 2255  
pair\_style oxrna2/excv, 2255  
pair\_style oxrna2/hbond, 2255  
pair\_style oxrna2/stk, 2255  
pair\_style oxrna2/xstk, 2255  
pair\_style pace, 2257  
pair\_style pace/kk, 2257  
pair\_style peri/eps, 2259  
pair\_style peri/lps, 2259  
pair\_style peri/lps/omp, 2259  
pair\_style peri/pmb, 2259  
pair\_style peri/pmb/omp, 2259  
pair\_style peri/ves, 2259  
pair\_style polymorphic, 2263  
pair\_style python, 2268  
pair\_style quip, 2272  
pair\_style rann, 2273  
pair\_style reaxff, 2279  
pair\_style reaxff/kk, 2279  
pair\_style reaxff/omp, 2279



pair\_style rebo, 1979  
 pair\_style rebo/intel, 1979  
 pair\_style rebo/omp, 1979  
 pair\_style resquared, 2285  
 pair\_style resquared/gpu, 2285  
 pair\_style resquared/omp, 2285  
 pair\_style saip/metal, 2288  
 pair\_style saip/metal/opt, 2288  
 pair\_style sdpd/taitwater/isothermal, 2290  
 pair\_style smatb, 2292  
 pair\_style smatb/single, 2292  
 pair\_style smd/hertz, 2294  
 pair\_style smd/tlsph, 2295  
 pair\_style smd/tri\_surface, 2296  
 pair\_style smd/ulsph, 2297  
 pair\_style smtbq, 2299  
 pair\_style snap, 2303  
 pair\_style snap/kk, 2303  
 pair\_style soft, 2307  
 pair\_style soft/gpu, 2307  
 pair\_style soft/omp, 2307  
 pair\_style sph/heatconduction, 2309  
 pair\_style sph/idealgas, 2310  
 pair\_style sph/lj, 2312  
 pair\_style sph/rhosum, 2313  
 pair\_style sph/taitwater, 2314  
 pair\_style sph/taitwater/morris, 2315  
 pair\_style spin/dipole/cut, 2319  
 pair\_style spin/dipole/long, 2319  
 pair\_style spin/dmi, 2321  
 pair\_style spin/exchange, 2322  
 pair\_style spin/exchange/biquadratic, 2322  
 pair\_style spin/magelec, 2325  
 pair\_style spin/neel, 2326  
 pair\_style srp, 2328  
 pair\_style srp/react, 2328  
 pair\_style sw, 2330  
 pair\_style sw/angle/table, 2335  
 pair\_style sw/gpu, 2330  
 pair\_style sw/intel, 2330  
 pair\_style sw/kk, 2330  
 pair\_style sw/mod, 2330  
 pair\_style sw/mod/omp, 2330  
 pair\_style sw/omp, 2330  
 pair\_style table, 2339  
 pair\_style table/gpu, 2339  
 pair\_style table/kk, 2339  
 pair\_style table/omp, 2339  
 pair\_style table/rx, 2343  
 pair\_style table/rx/kk, 2343  
 pair\_style tdpd, 2218  
 pair\_style tersoff, 2346  
 pair\_style tersoff/gpu, 2346  
 pair\_style tersoff/intel, 2346  
 pair\_style tersoff/kk, 2346  
 pair\_style tersoff/mod, 2351  
 pair\_style tersoff/mod/c, 2351  
 pair\_style tersoff/mod/c/omp, 2351  
 pair\_style tersoff/mod/gpu, 2351  
 pair\_style tersoff/mod/kk, 2351  
 pair\_style tersoff/mod/omp, 2351  
 pair\_style tersoff/omp, 2346  
 pair\_style tersoff/table, 2346  
 pair\_style tersoff/table/omp, 2346  
 pair\_style tersoff/zbl, 2355  
 pair\_style tersoff/zbl/gpu, 2355  
 pair\_style tersoff/zbl/kk, 2355  
 pair\_style tersoff/zbl/omp, 2355  
 pair\_style thole, 2360  
 pair\_style threebody/table, 2363  
 pair\_style tip4p/cut, 2036  
 pair\_style tip4p/cut/omp, 2036  
 pair\_style tip4p/long, 2036  
 pair\_style tip4p/long/omp, 2036  
 pair\_style tip4p/long/soft, 2097  
 pair\_style tip4p/long/soft/omp, 2097  
 pair\_style tracker, 2367  
 pair\_style tri/lj, 2369  
 pair\_style ufm, 2371  
 pair\_style ufm/gpu, 2371  
 pair\_style ufm/omp, 2371  
 pair\_style ufm/opt, 2371  
 pair\_style vashishta, 2373  
 pair\_style vashishta/gpu, 2373  
 pair\_style vashishta/kk, 2373  
 pair\_style vashishta/omp, 2373  
 pair\_style vashishta/table, 2373  
 pair\_style vashishta/table/omp, 2373  
 pair\_style wf/cut, 2377  
 pair\_style yukawa, 2379  
 pair\_style yukawa/colloid, 2380  
 pair\_style yukawa/colloid/gpu, 2380  
 pair\_style yukawa/colloid/omp, 2380  
 pair\_style yukawa/gpu, 2379  
 pair\_style yukawa/kk, 2379  
 pair\_style yukawa/omp, 2379  
 pair\_style zbl, 2383  
 pair\_style zbl/gpu, 2383  
 pair\_style zbl/kk, 2383  
 pair\_style zbl/omp, 2383  
 pair\_style zero, 2385  
 pair\_write, 975  
 partition, 977  
 plugin, 978  
 position (*lammps.Atom property*), 581  
 position (*lammps.Atom2D property*), 582  
 prd, 979  
 print, 983

processors, 985  
PyLammps (*class in lammps*), 577  
python, 989

## Q

quit, 996

## R

radius (*lammps.Atom property*), 581  
read\_data, 997  
read\_dump, 1018  
read\_restart, 1023  
region, 1026  
replicate, 1032  
rerun, 1033  
reset\_atom\_ids, 1036  
reset\_box() (*lammps.lammps method*), 560  
reset\_mol\_ids, 1038  
reset\_timestep, 1039  
restart, 1040  
run, 1043  
run() (*lammps.PyLammps method*), 578  
run\_style, 1046

## S

set, 1050  
set\_fix\_external\_callback() (*lammps.lammps method*), 569  
set\_variable() (*lammps.lammps method*), 564  
shell, 1058  
size (*lammps.NeighList property*), 584  
special\_bonds, 1060  
suffix, 1063  
system (*lammps.PyLammps property*), 579

## T

tad, 1065  
temper, 1069  
temper/grem, 1071  
temper/npt, 1073  
thermo, 1074  
thermo\_modify, 1075  
thermo\_style, 1079  
third\_order, 1085  
third\_order/kk, 1085  
timer, 1086  
timestep, 1088  
torque (*lammps.Atom property*), 581  
type (*lammps.Atom property*), 580

## U

uncompute, 1089  
undump, 1090

unfix, 1091  
units, 1092

## V

variable, 1096  
variables (*lammps.PyLammps property*), 580  
velocity, 1116  
velocity (*lammps.Atom property*), 581  
velocity (*lammps.Atom2D property*), 582  
version() (*fortran function*), 532  
version() (*lammps.lammps method*), 559  
version() (*lammps.PyLammps method*), 577  
video() (*lammps.IPyLammps method*), 583

## W

write\_coeff, 1119  
write\_data, 1120  
write\_dump, 1123  
write\_restart, 1124  
write\_script() (*lammps.PyLammps method*), 578